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Mechanism of Action of Liver Growth Induced by Peroxisome Proliferators

by

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Thesis submitted to The University of Nottingham for
the degree of Doctor of Philosophy

17th February 2011



In The Name Of God (Allah) The Most Merciful The
Most Gracious

Abstract

Humans are ubiquitously exposed to peroxisome proliferators including hypolipidemic agents, industrial solvents and natural products. Because of this and the fact that peroxisome proliferators cause non-genotoxic hepatocarcinogenesis in rodents, it is of importance to elucidate the mechanism of action of the peroxisome proliferators in order to provide an assessment of the hazard, if any, of these compounds to humans. It is also known that the peroxisome proliferators begin their actions by inducing hepatic DNA synthesis. Thus, the aim of this thesis was to find genes that could be responsible for triggering the induction of hepatic DNA synthesis caused by peroxisome proliferators, specifically ciprofibrate.

First, it was important to indicate when the induction of hepatic DNA synthesis actually happens. This was done with BrdU immunohistochemical procedures. The induction of hepatic DNA synthesis with ciprofibrate in mice was observable only after 4 days making it difficult to specify when the induction actually happened. In rats the induction of hepatic DNA synthesis was found to peak at 24 hours and this system gave the better opportunity to find the genes responsible. The difference in the timing of induced hepatic DNA synthesis between mice and rats implied that there could be a species difference in the mechanism of each species' response to PPAR. With immunohistochemistry it was noticed that there was a difference in the lobular localization of hepatic DNA synthesis in the liver tissues of rats and mice dosed with different inducers, with the rat livers exhibiting periportal distribution while hepatic DNA synthesis in the mice seemed to be distributed throughout the liver tissue.

The effects of ciprofibrate or cyproterone acetate on liver gene expression in rats were studied, using cDNA microarrays, transcriptome sequencing and quantitative real-time PCR. A 1- 5 hour treatment period was chosen to detect the immediate early gene response, while a 24 hour time point was chosen to elucidate the confounding effects from the hepatic DNA synthesis seen

during the 24 hour stimulation. The results showed that ciprofibrate altered the expression of numerous genes including previously known PPAR α agonist-responsive genes involved in processes such as PPAR signalling pathways, fatty acid metabolic pathway, cell cycle, palmitoyl-CoA hydrolase activity, lipid metabolism, inflammatory responses, and stress responses, in addition to a large number of novel candidate genes.

Three novel induced genes *G0s2*, *Ccnd1* and *Scd1*, (and two marker genes *CYP4A1* and *CYP3A1*) were confirmed with quantitative real-time PCR. The *G0s2*, *Ccnd1* and *Scd1* were found to be up-regulated at the hours 1 and 3 after dosing and not 24 hours, and the *G0s2* and *Scd1* were specific for the ciprofibrate suggesting they were involved in a distinct PPAR α pathway responsible for the hepatic DNA synthesis. The complete database of the transcriptional response provided here opens doors of opportunity for further research to identify genes responsible for the liver growth induced by peroxisome proliferators.

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Finally, I would like to thank the Libyan people for providing financial support for my study and thank you to all my family and friends for their help and support in making this PhD a pleasurable time of my life.

Dedication

This thesis is dedicated to

my Mother and Father, my Husband Wail and to my Children

Mender, Asaa and Mohammed

This thesis is dedicated to

the memory of my grandfather Ayad Amer

and to the

Martyrs of the Libyan Cause

Declaration

I declare, that this work was achieved in the duration of my PhD studies at the University of Nottingham, and is my original effort unless otherwise stated. Information from other resources has been fully recognized. No part of this thesis has been submitted for assessment leading to a degree.

Abeer Amer

17th February 2011

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Presentations and publications arising from this thesis

Platform presentations

1- **A. H. Amer**, A. M Al Kholaiifi, T. Gant and D. R. Bell. (2010). Induction of liver growth by peroxisome proliferators. At the British Toxicology Society, Edinburgh. UK.

Poster presentations

1- **A. H. Amer**, A. M. Al Kholaiifi and D. R. Bell. (2008). Effects of peroxisome proliferators on zonated DNA synthesis in rat and mouse liver. At the British Toxicology Society, Surrey.

Abstracts

1- **A. H. Amer**, A. M Al Kholaiifi, T. Gant and D. R. Bell. (2010). Mechanism of action of liver growth induced by peroxisome proliferators. *Toxicology*. 278, (3), 350-351.

2- A. H. Amer, A. M. Al Kholaiifi and D. R. Bell. (2008). Effects of peroxisome proliferators on zonated DNA synthesis in rat and mouse liver. *Toxicology*. 253, (1), 26-27.

Papers

1- A. Al Kholaiifi, **A. Amer**, B. Jeffrey, T. J. Gray, R. A. Roberts and D. R. Bell. (2008). Species-specific kinetics and zonation of hepatic DNA synthesis induced by ligands of PPAR alpha. *Toxicol Sci*. 104 (1) 74-85.

2- **A. Amer**, F. Aboshofa, A. Al Kholaiifi, T. Gant and D. Bell. (2011). Mechanism of action of liver growth induced by peroxisome proliferators. (manuscripts in progress).

List of Abbreviations

ACO	Acyl-CoA
AhR	Aryl hydrocarbon receptor
ALT	Alanine aminotransferase
ANOVA	Analysis of variance
AP-mice	Alderley park mice
β -actin	Beta actin
BrdU	5-Bromo-2-deoxyuridine
BSA	Bovine serum albumin
C/EBP α	CCAAT-enhancer-binding protein alpha
CAR	Constitutive androstane receptor
CCl ₄	Carbon tetrachloride
Cnd1	Cyclin D1 a cell cycle
cDNA	Complementary DNA
CPA	Cyproterone acetate
Ct	Cycling threshold
CV	Central vein/ perivenous
CYP3A1	Cytochrome P450, family 3 subfamily A, polypeptide 1
CYP4A1	Cytochrome P450, family 4 subfamily A, polypeptide 1
DAB	3,3' diamino-benzidine tetrahydrochloride
DECP	Diethyl pyro carbonate

DEHP	Di-(2-ethyl-hexyl) phthalate
DMSO	Dimethyl sufoxide
DPX	Distyrene, plasterine and xylene
EdU	5-ethynyl-2'-deoxyuridine
F	Forward primer
F-344/NHsd	Fisher-344/NHsd rats
FDA	Food and drug administration
FDR	False discovery rate
G0	Gap zero
G0s2	G0/G1 switch 2
GAL	GenePix Array List
GPR	GenePix results
GstYb4	Glutathione S-transferase mu 3
H	Hepatocytes
HEPES	4-(2-hydroxyethyl)-1-piperazineethanesulfonic acid
HGF	Hepatocyte growth factor
HNF4 α	Hepatic nuclear factor-4 α
IGFBP1	Insulin-like-growth-factor-binding protein-1
IP	Intra-peritoneal
IQR	Inter quartile range
KEGG	Kyoto encyclopedia of genes and genomes

LI	Labelling index
MAPK/ERK	Mitogen-activated protein kinase/extracellular signal-regulated kinase
MCP	Methylcyclopentane
MEEBO	Mouse exonic evidence based oligonucleotide
MIAME	Minimum information about a microarray experiment
NCTR	National centre for toxicological research of the U.S. A
NF-kB	Nuclear factor kappa beta
NP	Non-parenchymal cells
NTC	No template control
OD	Optical density
P	Probe
PBS	Phosphate buffer saline
PCN	Pregnenolone-16 α -carbonitrile
PCR	Polymerase chain reaction
PPAR	Peroxisome proliferator activated receptor
PPRE	Peroxisome proliferator hormone response element
PS	Portal space/ periportal
PXR	Pregane X receptor
R	Reverse primer
RGD	Rat genome database
RNAseq	Transcriptome sequencing analysis

RPKM	Reads per kilobase of exon model per million mapped reads
RT	Reverse transcriptase
RXR	Retinoid X receptor
S phase	Synthesis phase
Scd1	Sterol-coenzyme A desaturase 1
SD	Standard deviation
SDS	Sodium lauryl (dodecyl) sulfate
SSC	Saline-sodium citrate buffer
STAT 3	Signal transducer and activator of transcription 3
T3	Thyroid hormone
TBE	Tris boric acid EDTA
TBS	Tris buffer saline
TCPOBOP	1,4-Bis[2-(3,5-dichloropyridyloxy)]benzene
TGF α	Transforming growth factor alpha
TGF β	Transforming growth factor beta
TIFF	Tagged image file format
TNF α	tumour necrosis factor alpha
Tween-20	Polyoxyethylene sorbitane monolaureate
WY-14,643	Pirinixic Acid/ 4-chloro-6-[(2,3-dimethylphenyl)amino]-2-pyrimidinyl]thio]-acetic acid

Chapter 1 Introduction

Section 1.1 The Liver

Section 1.1.1 Histology and physiology of the liver

The liver is the largest organ in the body. Found on the right side of the abdomen under the lungs covered partially by the lower ribs, it consists of two main lobes, the right lobe and the left lobe with the right being the largest (Mader, 2004). The liver has a central role in metabolic homeostasis of the body, as it is responsible for the metabolism, synthesis, storage and redistribution of nutrients, carbohydrates, fats and vitamins. One of the main functions of the liver is detoxification, which removes waste and xenobiotics by metabolic conversion and biliary excretion (Kurt, 1991).

The hepatic parenchyma is organized into units called liver lobules (Michalopoulos and De-Frances, 1997), as illustrated for the pig liver in Figure 1.1. The classic liver lobules are divided with connective tissue and appear in a hexagonal shape with a central vein in the middle and portal spaces (triads) at the corners.

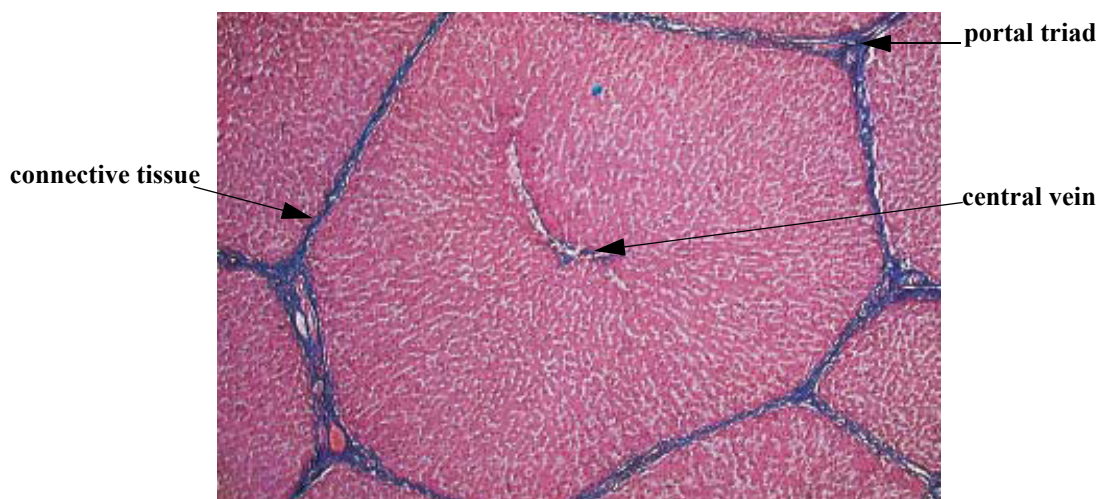


Figure 1.1 Histological view of the classic liver lobules. Histological view of a pig liver tissue, with a light microscope. Staining was with azan stain. The central vein is in the middle of the slide and the portal space (triad) are shown at the sides. The liver lobules are clearly identified with connective tissue defining each lobule (blue in colour). Magnification X40. Figure from (Kühnel, 2003).

A different structure of the liver is the portal lobule where the structure appears as a triangle with a central vein at each of the corners and a portal space in the middle. A third structure is called the liver acinus which appears as a diamond shape connecting a central vein to a portal space then a central vein and a portal space then back to the same central vein. It was reported that the liver acinus is the functional unit of the liver and is divided into three zones: Z1 periportal, Z2 intermediate and Z3 perivenous (Ross, 2003) indicated as 1, 2 and 3 in Figure 1.2.

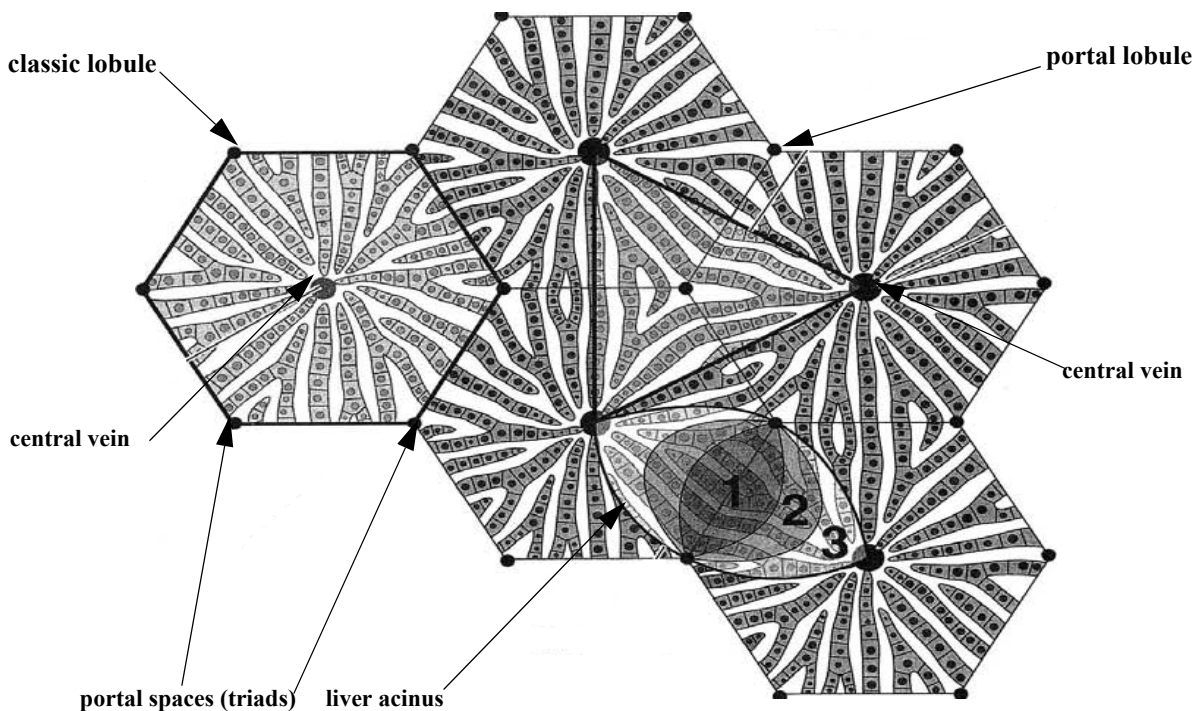


Figure 1.2 Diagram of the liver structure. Shown is the **classic liver lobule** where a central vein is in the middle and six portal triads are at the periphery. The **portal lobule** appears as a triangle with a central vein at each of the corners and a portal triad in the middle. The **liver acinus** appears as a diamond shape connected between a portal triad, central vein, portal triad and central vein this is the functional unit of the liver and is divided into three zones (the area around the central vein is zone 1 (Z1), the area near the terminal portal vein is zone 3 (Z3) and the area in between is zone 2 (Z2)). Figure from (Henrikson et al., 1997).

Liver cells closest to the terminal portal vein (periportal) receive blood richer in oxygen and nutrients than those located near the central vein (perivenous). The cells in these different zones perform different biochemical functions and respond differently to injuries, with regard to the cell organelle number and size, ploidy, pattern of enzymes activity, autonomous innervations, number of non parenchymatous cells, concentration of hormones, substrates and their metabolites, and the partial O_2 pressure (Taub, 2004). Figure 1.3 depicts the structure of a liver lobule, with hepatocytes branching from the central vein to the portal vein (triad) in rows 1-2 cells thick called plates. The portal space includes the hepatic portal vein, the hepatic artery and the bile duct.

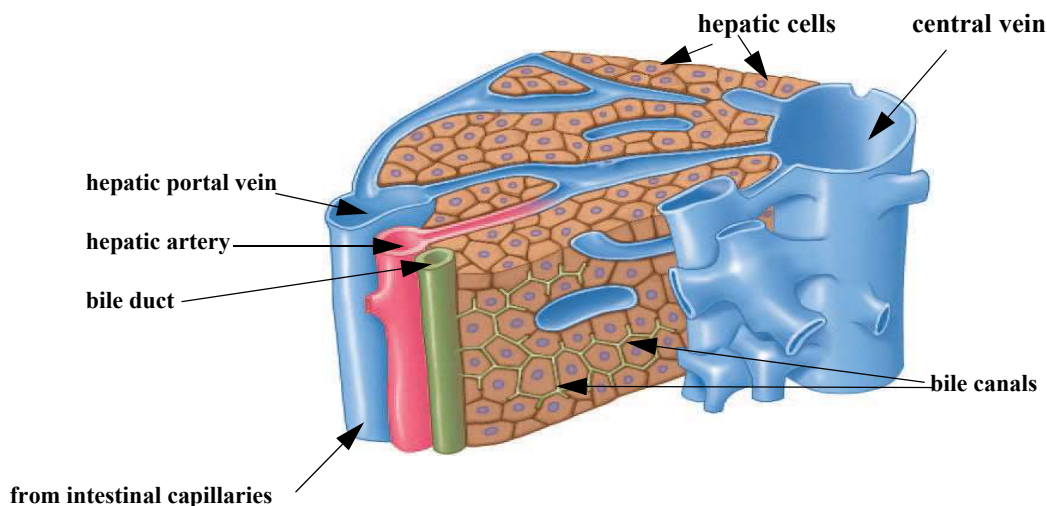


Figure 1.3 Diagrammatic structure of a liver lobule. the hepatocytes form rows of one or two cells thick and are called plates, the plates start at the central vein, also known as the inter lobular vein, and branch out in a star shaped way to the portal space. Between the cell plates are the sinusoids which contain the Kupffer cells and the epithelium cells (not shown), between each two hepatocytes starts the Bile canals which get larger and larger to make the bile ducts. A bile duct along with a portal vein and a hepatic artery get arranged in a group making the portal space. Figure from (Mader, 2004).

The main cell type found in the liver is parenchymal cells also known as the hepatocytes, which makes up most of the hepatic cells. Other cell types are the endothelial cells, Kupffer cells, lymphocytes, stellate cells and lipocytes (Dudek, 2000)(Taub, 2004) (Figure 1.4).

Endothelial cells line the sinusoids of the liver, and provide a large surface area for nutrient absorption, and Kupffer cells are situated in the sinusoids and have a phagocyte function. Also, it is thought that Kupffer cells produce cytokines (Taub, 2004).

Lymphocytes are considered part of the innate immune system that resides within the liver to help resist infection. Hepatic cells have various functions, including the storage of vitamin A and the production of the extracellular matrix and most of the factors that lead to hepatic fibrosis. All of these cell types are activated by hepatic injury (Taub, 2004).

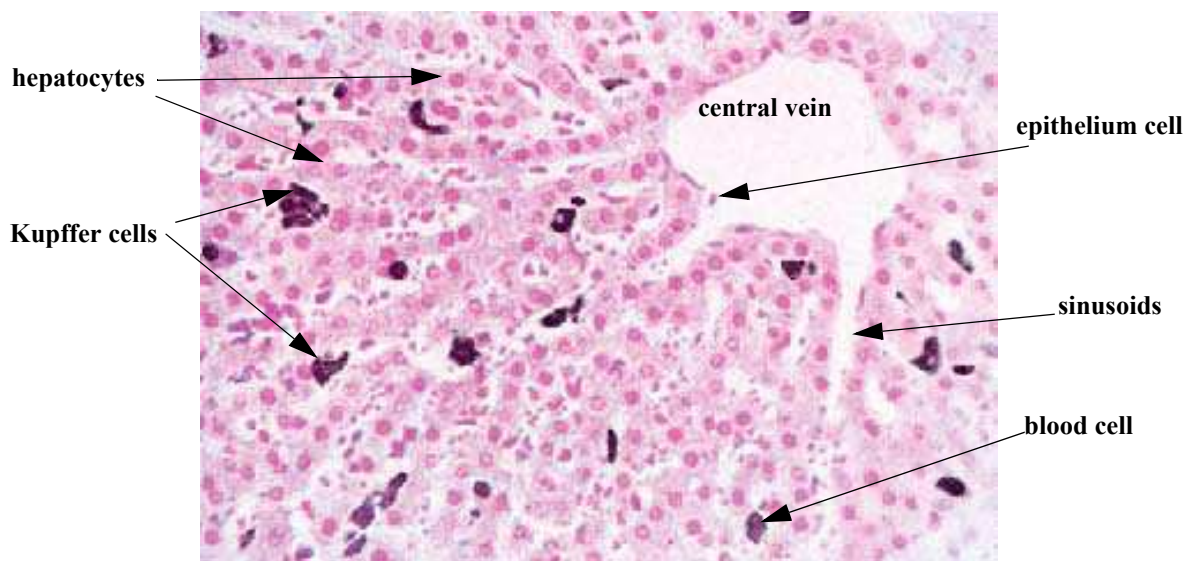


Figure 1.4 Histological section of liver cells. Histological section of liver tissue showing the hepatocytes in rows making the plates. Between the plates are the sinusoids, the endothelium cells are covering the sinusoids and some blood cells are shown in between the cells. The Kupffer cells are shown as dark stained cells. The stain is Giroud-Leblond; nuclear staining with carmine red; magnification: X300, from (Kühnel, 2003).

Section 1.1.2 Liver regeneration (growth)

Normally, mature hepatocytes live long and do not divide; they only proliferate in response to toxic injury and infection. Loss of liver function results in proliferation and restoration of functional liver tissue (Mangnall et al., 2003).

The fascinating two thirds partial hepatectomy (PH) model in rodents was first pioneered by (Higgins, 1933) as a model for regrowth. In this model, two thirds of the liver was surgically removed, and the remaining liver enlarged until the original liver mass was restored, approximately 1 week after surgery, after which the regenerative process stopped. In this experimental system, liver regeneration does not require the recruitment of liver stem cells or progenitor cells, but involves replication of the mature functioning liver cells. The regenerative process is compensatory because the size of the resulted liver is determined by the demands of the organism, and, once the original mass of the liver has been re-established, proliferation stops (Michalo-

poulos and DeFrances, 1997).

Section 1.1.3 Basic characteristics of liver regeneration

Because hepatocytes are mostly found in the G₀ phase of the cell cycle, hepatic cell division is rarely found in the normal adult liver (Michalopoulos and DeFrances, 1997). It was reported that after partial hepatectomy the liver cells start dividing, and by day three the liver mass begins to increase, while the whole mass restoration is complete in just 5-7 days (Grisham, 1962).

Thus, after partial hepatectomy in rats nearly 95% of hepatocytes rapidly re-enter the cell cycle, and hepatic DNA synthesis (cells in the S phase) increases after 12 hours and peaks around 24 hours (Mangnall et al., 2003). However, the induction of non-parenchymal DNA synthesis occurs later (36-48 hours for kupffer cells).

Subsequent to the partial hepatectomy, the levels of hepatic DNA synthesis are lower, as the liver mass requires an average of 1.6 cycles of replication in all cells to be completely restored, taking around 7 days. The expression of the gene encoding β -actin (a growth related gene) after 2 hours is high until about 36 hours, while the insulin-like-growth-factor-binding protein-1 (IGFBP1) (encoded by a cell-cycle-regulated gene) peaks after 2-3 hours. Genes encoding the α isoform of CCAAT enhancer binding protein (C/EBP α), are down-regulated within the period of maximal growth and are expressed again after the growth phase has occurred. This is clarified in Figure 1.5 (Taub, 2004).

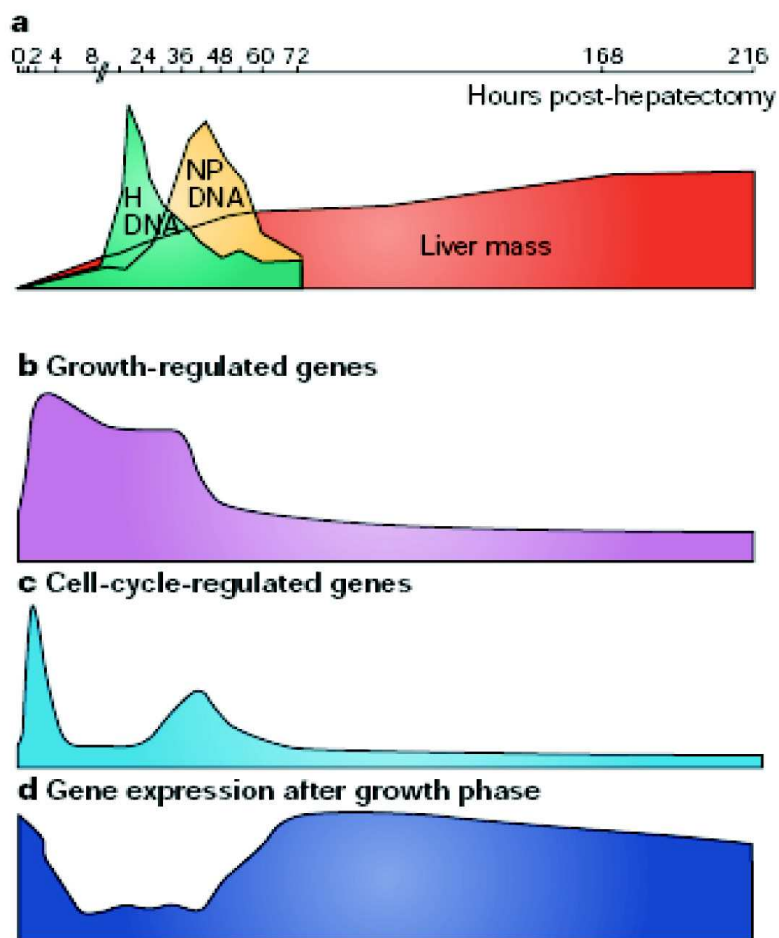


Figure 1.5 Diagrammatic illustration showing patterns of DNA synthesis and induction of gene expression during rat liver regeneration. a) 24 hours after partial hepatectomy, (H; green) DNA synthesis in hepatocytes peaks, but in the non-parenchymal cells DNA synthesis (NP; yellow) reaches its highest at 36-48 hours. Re accumulation of liver mass in red is finished in around a week. b) The pattern of induction of the genes expression for growth regulated genes, (β -actin). c) While the induction pattern of insulin like growth factor binding protein-1 (*IGFBP1*) representing the gene expression for cell-cycle-regulated genes. d) The genes encoding the α isoform of CCAAT enhancer binding protein (*C/EBP α*), are down-regulated within the period of maximal growth and are expressed again after the growth phase has occurred.(Taub, 2004).

By comparison, in mice after partial hepatectomy the DNA synthesis peak occurs later (36- 40 hours after partial hepatectomy) and varies between strains (Sigal et al., 1999).

Early studies *in vitro* with isolated hepatocytes, identified a number of potential hepatocyte growth factors such as HGF, TGF α and TGF β . It was difficult to confirm that any of these fac-

tors had an vital role in liver regeneration, but it was noted that growth-factor-and cytokine-regulated pathways are activated during liver regeneration (Taub, 2004).

Transcription factors have been acknowledged to play a major role in making hepatocytes proliferate (Fausto, 2000), apparently through the induction of the hepatic expression of numerous immediate early genes linked to the cell cycle (Searce et al., 1996). Knockout mouse studies were used to identify specific genes induced during the induction of DNA synthesis during hepatic regeneration. These studies established that cytokines might be responsible for regenerative response (Taub et al., 1999), and IL-6 could collaborate with additional factors to trigger the other up-regulated genes (Cressman et al., 1996). Other genes that are regulated during hepatic regeneration are the genes that encode the cyclin D1 (Schwabe et al., 2003)(Talarmin et al., 1999)(Diehl, et al., 1994), IGFBP1 91, STAT 3 and AP1 (Leu, et al., 2001)

One transcription factor, termed NF- κ B, has been suggested to have a role in liver regeneration (Tewari et al., 1992). A potent inducer of *NF- κ B* is tumour necrosis factor α (TNF- α) (Grilli et al., 1993)(Menegazzi et al., 1997).

The beginning of DNA synthesis is well synchronized in rat hepatocytes, starting in cells that surround the portal vein of the liver lobule and proceeding towards the central vein (Taub, 2004). Hepatic proliferation is also noted in the liver regeneration caused by apoptosis or necrosis that is induced by toxic chemicals e.g. CCL₄ (Taub, 2004) and peroxisome proliferators (Moody et al., 1991).

Section 1.2 Peroxisome proliferation

Section 1.2.1 Peroxisomes

Peroxisomes are single membrane-limited cytoplasmic organelles present in both animal and plant cells. They were first detected in 1954 and are known as “microbodies”. They were re-

ferred to as peroxisomes after it was found that they contain catalase and oxidase, which are involved in the degradation and production of hydrogen peroxide (Deduve, 1965). Other functions of the peroxisomes include β -oxidation of fatty acids and cholesterol metabolism (Holden and Tugwood, 1999).

Peroxisomes are mostly found in liver and kidney cells (Bernhard and Rouiller, 1956) (Rhodin, 1954). Leighton and his group found that peroxisomes account for nearly 2.5% of protein in rat liver (Leighton et al., 1968). The following year Weibel and his group found that the peroxisomes occupy about 1.5% of the parenchymal cell volume (Weibel et al., 1969).

Hepatocyte peroxisomes are affected in number and size by chemicals not related in structure, called the peroxisome proliferators (Reddy, 2004).

Section 1.2.2 Peroxisome proliferators (PPs)

Peroxisome proliferators are a group of structurally unrelated chemicals with different applications, such as hypolipidemic drugs, and industrial and agricultural chemicals (Moody et al., 1991).

It is reported that peroxisome proliferators induce peroxisome proliferation in livers of some rodents. Peroxisome proliferation involves changes in the ultra structure and metabolic capacity of the liver cell, such as the increase of peroxisome volume density, hepatomegaly, and an increase in β -oxidation of fatty acids and peroxisomal enzymes activities (Moody et al., 1991).

Interestingly, it has been reported that DNA synthesis is not affected by peroxisome proliferators in pure hepatocytes, rather they stimulate Kupffer cells to make mitogens which stimulate proliferation of parenchymal cells (Parzefall et al., 2001).

However, peroxisome proliferation involves several other changes that are not strictly 'peroxi-

somal', and these changes involve metabolic functions in the cell, as well as shifts the rate of entry of hepatocytes into cell proliferation (mitosis) and cell death (apoptosis) (Cattley, 2003).

Although many chemicals have been evaluated for peroxisome proliferation, several have been commonly used in mechanistic studies. Among the hypolipidemic drugs studied, clofibrate (ester), clofibric acid, and ciprofibrate are in clinical use (Cattley, 2003). Other hypolipidemic drugs include WY-14,643, BR-931, methylclofenapate, and nafenopin, compounds that were discovered in the search for more potent drug candidates, but were not entered into clinical use.

Among the non drug compounds used in research is the phthalate ester plasticizer di-(2-ethylhexyl) phthalate (DEHP), which is often used because it is cheap, available with high purity, and easy to dose (Cattley, 2003).

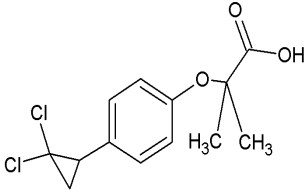
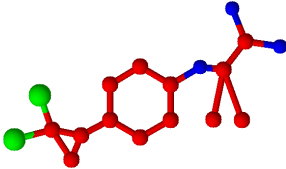
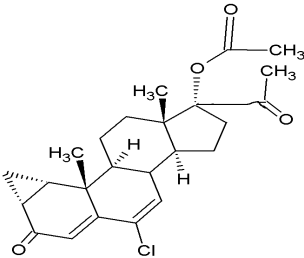
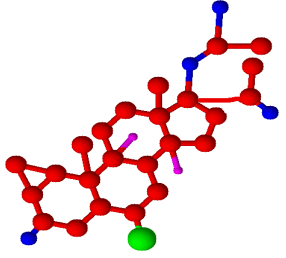
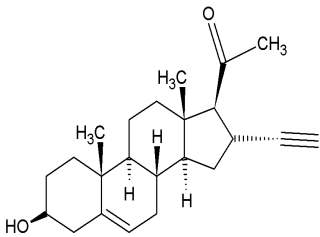
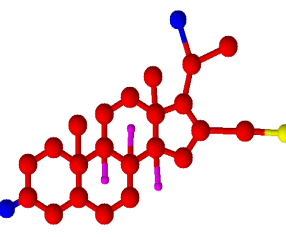
Section 1.2.3 Ciprofibrate, cyproterone acetate and pregnenolone-16 α -carbonitrile

Ciprofibrate is a strong peroxisome proliferator activated receptor-alpha (PPAR α) ligand (Mukherjee et al., 2002) from the fibrate hypolipidemic drug family, and a potent hepatocarcinogen (Meyer et al., 2003) (Yadetie et al., 2003).

The anti androgen cyproterone acetate (CPA) and pregnenolone-16 α -carbonitrile (PCN) are pregnane X receptors (PXR). Cyproterone acetate is reported to cause liver tumours and has a mutagenic effect in rats (Topinka et al., 2004a). It is also known to induce cell proliferation (Schulte-Hermann et al., 1980) and apoptosis (Kasper and Mueller, 1999) in rat liver *in vitro* and *in vivo*. Cyproterone acetate is used as a hormonal therapy to treat prostate cancer as it suppresses the action of testosterone and dihydrotestosterone on cells (Green et al., 2002). It is also reported to be used in some countries other than America and Japan as a hormone treatment for acne, with some contraceptive benefits (van Vloten et al., 2002).

Table 1.1 illustrates the chemical structure of ciprofibrate, CPA and PCN used in this study.

Table 1.1 Chemical formula and structure of ciprofibrate, cyproterone acetate and pregnenolone-16 α -carbonitrile. This table shows the systematic IUPAC (International Union Of Pure And Applied Chemistry) name, the chemical formula and 2 and 3 dimensional structure of the chemicals used to perform the induction of the hepatic DNA synthesis in this project. Chemical structures were drawn using CDS/ISIS DRAW Program.

Systematic IUPAC Name	Chemical Formula	Structure	3D Structure red=carbon blue=oxygen green=chlorine yellow=nitrogen pink=hydrogen
<p>ciprofibrate</p> <p>2-[4-(2,2-dichlorocyclopropyl) phenoxy] 2-methylpropanoic acid</p>	$C_{13}H_{14}Cl_2O_3$		
<p>cyproterone acetate (CPA)</p> <p>chloro-6-hydroxy-17 alpha methylene-1 alpha, 2 alpha pregnadiene-4,6 dione-3, 20 acetate</p>	$C_{24}H_{29}ClO_4$		
<p>pregnenolone-16α-carbonitrile (PCN)</p> <p>3S,8S,9S,10R,13S,14S,16R,17S)-17-acetyl-3-hydroxy-10,13-dimethyl-2,3,4,7,8,9,11,12,14,15,16,17-dodecahydro-1H-cyclopenta[a]phenanthrene-16-carbonitrile</p>	$C_{22}H_{31}NO_2$		

Section 1.2.4 Peroxisome proliferator activated receptor-alpha (PPAR- α)

A number of years ago PPAR α was the first of three structurally related peroxisome proliferator receptors (PPARs) to be identified of the soluble nuclear hormone receptor superfamily. Peroxisome proliferators are believed to act through this family of proteins (PPAR α) (Issemann and Green, 1990). The other two receptors were later indicated and are now known as PPAR β and PPAR γ (Dreyer et al., 1992)(Kliewer et al., 1994), but they do not mediate peroxisome proliferation (Peraza et al., 2006).

PPAR along with constitutive active androstane receptor (CAR) and aryl hydrocarbon receptor (Ahr) are known to operate as sensors of xenobiotic entrance into the cell. Mice carrying knock-outs in genes for the receptors cooperating with xenobiotic were revealed to be deficient for the promoting effects of these compounds (Yamamoto et al., 2004) (Peters et al., 1997a).

Also, PPARs are thought to be transcription factors that are activated by ligands and by interaction with elements situated 570 bp upstream of the peroxisomal enzyme acyl CoA oxidase receptive genes (Tugwood et al., 1992). Studies have demonstrated that PPAR α is an obligatory factor in peroxisome proliferation in rodent hepatocytes *in vivo* and *in vitro* (Klaunig et al., 2003).

It has been suggested that Human PPAR α have many similar functional characteristics to the rodent receptors, suggesting that the former also may be activated by peroxisome proliferators and regulate specific gene expression (Ashby et al., 1994). However, PPAR α is less abundant in human liver than in rodent liver, which has led to the suggestion that species differences result from quantitative differences in gene expression.

Section 1.2.5 Toxicological changes induced by peroxisome proliferators

Although there are no acknowledged toxicities linked with endogenous ligands of PPAR α , there is a number of findings related to toxicity and the synthetic PPAR α ligands in animal models (Peraza et al., 2006).

For example it has been reported that administration of clofibrate (a fibrate peroxisome proliferator) can cause some signs of maternal toxicity if used prior to or during pregnancy in high doses (Cibelli et al., 1988)(Stefanini et al., 1989)(Wilson et al., 1991).

Other PPAR α agonists, like phthalates and trichloroethylene in rodents have been reported to cause altered ovulation (Davis et al., 1994), reduced fertility rates (Peters and Cook, 1973)(Singh et al., 1974), teratogenesis including cardiac, skeletal, and neural tube defects (Gao et al., 2003)(Ritter et al., 1985) impaired spermatogenesis and altered development of the male reproductive tract (Corton and Lapinskas, 2005).

It is also noted that exposure to perfluorooctanoic acid causes reduced fetal weights and cardiac and skeletal malformations (Lau et al., 2004). However, the toxic effects were not proven to be linked with PPAR α pathways as clarified by Peters et al. (1997b) who administered Di(2-ethylhexyl) phthalate (DEHP) during organogenesis, in wild-type and PPAR α -null mice and this caused neural tube defects in both, indicating that PPAR α was not required to mediate the effect.

On the other hand, fibrate therapy is linked with cholelithiasis (gallstones), as it has been recorded in humans treated with fenofibrate, clofibrate or bezafibrate which were associated with an increased incidence of cholelithiasis (Caroli-Bosc et al., 2001)(Raedsch et al., 1995) and also myopathy and rarely rhabdomyolysis (Bridgman et al., 1972) (Alsheikh-Ali et al., 2004).

But it is well-known that PPAR α ligands cause hepatocellular carcinomas in rodents, and they have also been connected to other malignancies, as they cause pancreatic acinar cell tumours and Leydig cell tumours (Reddy et al., 1980). Furthermore, pancreatic acinar cell tumours and Leydig cell tumours have been reported to occur only in rats, but not in mice (Klaunig et al., 2003). It was also reported that clofibrate and Wy-14,643 treatments increase the growth of human breast cancer cell lines (Suchanek et al., 2002).

Section 1.3 Mechanism of action of carcinogenesis

Among the first stages of carcinogenesis provoked by chemicals are initiation and promotion, when both initiation and promotion occur this is known as “complete carcinogenesis” and the peroxisome proliferators would cause both initiation and promotion activities (Kobliakov, 2010). The initiation stage is provoked by genetic changes in the genome caused by genotoxic metabolites of carcinogens, whilst the promotion stage is initiated by a non genotoxic mechanism and does not effect the DNA structure (Gujaeva et al., 1998).

Promotion requires particular scenarios, including increased cell proliferation, decreased intercellular interactions, and inhibition of apoptosis (Kobliakov, 2010). This promotion stage, caused by non genotoxic compounds is not well studied and it is unknown if the pathways are similar to the initiation stage or not (Kobliakov, 2010).

Additionally, it is stated by Wu that the promoter induces the production of reactive oxygen species in the cells and this affects intercellular interactions, stimulates the cell migration, and inhibits apoptosis (Wu et al., 2006). However, the pathways responsible for these changes have yet to be resolved.

Section 1.3.1 Peroxisome proliferators induce liver cancer in rodents

Peroxisome proliferators are known for their ability to cause hepatocellular carcinoma in laboratory rodents (Reddy et al., 1980). The effect of this is recorded by (Cattley et al., 1991) who showed that in rodents, after just one year of treatment with Wy-14,643 (a potent peroxisome proliferator) there was 100% incidence of multifocal liver cancer. Gonzalez and his group established that the PPAR α mediated pathways that control lipid metabolism and the cell proliferation pathways are independent of each other. This was achieved with PPAR α humanized transgenic mice, and this suggests that these mice would not be susceptible to peroxisome proliferator induced hepatocarcinogenesis (Cheung et al., 2004).

Other studies showed that just one week of exposure to Wy-14,643 causes hepatomegaly and hepatocytes entering S-phase. Also, this exposure resulted in hepatocellular neoplasms, including some carcinomas after 11 months exposure in wild type mice and not in PPAR α null mice. This makes it clear that PPAR α is essential for the increase in hepatic replicative DNA synthesis in response to peroxisome proliferators.

Although it is known that PPAR α cause cancer in long term exposures in rodents, there are significant differences in species response, and PPAR α is not recorded to induce peroxisome proliferation and hepatocarcinogenesis in humans (Yadete et al., 2003).

Section 1.3.2 Peroxisome proliferators are non-genotoxic carcinogens

Chemical carcinogens are commonly classified on the basis of their mechanism of action; genotoxic (mutagenic) and non genotoxic (non mutagenic) (Weisburger and Williams, 2000). An important aspect of peroxisome proliferators is that they are non-genotoxic carcinogens (Ashby et al., 1994)(Peters et al., 1997a).

It has been proposed that during carcinogenesis peroxisome proliferators promote liver tumours by increasing the production of H_2O_2 leading to the production of peroxisomal fatty acyl CoA oxidase (Yeldandi et al., 2000). Moreover, through carcinogenesis and tumour growth intercellular interactions are affected and this is important for the normal regulation of organs. The morphology of the cells are also changed, and the epithelial mesenchymal transition is detected (Radisky et al., 2005).

It is understood that peroxisome proliferators reduce the expression of glutathione peroxidase and increase the oxidative stress in hepatocytes caused by the large increase in the H_2O_2 levels and the gathering of lipofuscin in the liver cells (Yeldandi et al., 2000)(Kobliakov, 2010). It is also reported that peroxisome proliferators begin hepatocarcinogenesis in rodents by increasing cell proliferation (Reddy et al., 1980) (Becuwe and Dauca, 2005). However, the mechanism that PPAR α ligands cause cancer is still vague (Ashby et al., 1994)(Klaunig, 2003)(Peters et al., 2005).

Luci (Luci et al., 2007) argued that the hepatocarcinogenesis is a result of oxidative stress and an impaired balance between apoptosis and cell proliferation in the liver, while Shah and Morimura (Shah et al., 2007a) and (Morimura et al., 2006) found that PPAR α humanized mice appear to be resistant to the induction of peroxisome proliferators and less sensitive to the development of liver tumours.

Section 1.4 Induction of hepatic DNA synthesis by peroxisome proliferators

It is clear that DNA synthesis plays a significant role in carcinogenesis and that peroxisome proliferators provoke hepatic DNA synthesis that is associated with carcinogenesis (Marsman et al., 1988). It is therefore desirable to understand the mechanism of the induction of DNA synthesis by PPAR α ligands, particularly since these ligands increase the normal liver size, in contrast to the regenerative growth pathways induced by partial hepatectomy (Mangnall et al., 2003).

Section 1.4.1 Acute and chronic effects of peroxisome proliferators on hepatic DNA synthesis

Menegazzi's group showed that rats treated with a peroxisome proliferator known as nafenopin or with CPA (a PXR) would cause hepatic DNA synthesis that starts 18 to 24 hours after treatment. They also reported that the levels of hepatic TNF- mRNA, NF-B and AP-1 were not changed indicating a difference from the results obtained after partial hepatectomy, although the hepatic cells entered the S phase of the cell cycle at similar times (Menegazzi et al., 1997).

In a recent study, Al-kholaifi showed that the effect of ciprofibrate on hepatic DNA synthesis in male F-344/NHsd rats peaked between 24 and 48 hours after treatment, while 129S4/SvJae and 129S4/SvJae PPAR α -null mice treated with ciprofibrate showed hepatic cells entering the S phase only after 3-4 days (Al Kholaifi, 2008)(Al Kholaifi et al., 2008).

It is known from former studies that the replicating hepatic nuclei are not distributed randomly within the rat liver, but are found predominantly in the periportal region (Barrass et al., 1993). Barrass's results were achieved in male Sprague-Dawley rats treated with methylcyclopentane (MCP) and clofibric acid for 1-26 weeks. This was confirmed by Al kholaifi with Fisher rats, while most mouse species' livers didn't show any preference in the zonation of hepatic DNA

synthesis (Al Kholaiifi et al., 2008). Other zonation studies on mouse liver using a different approach/ method on young non treated B6C3F1 and C57BL mice showed that the distribution is mostly in the mid-zone Z2 (Amacher et al., 1998).

The difference in the kinetics of induction of hepatic DNA synthesis between mouse and rat after exposure to PPAR α ligands suggests that there might be a difference in the mechanism of hepatic DNA synthesis of the two species, and how each of the species' livers respond to PPAR α .

Frick and Blaauboer showed in separate studies that humans do not display the same range of hepatocarcinogen peroxisome proliferator-induced responses seen in rats and mice (Frick et al., 1987)(Blaauboer et al., 1990). Also it has been demonstrated that non-human primates and guinea pigs are unaffected by peroxisome proliferators (Bell et al., 1998). So, establishing an explanation for, and understanding the risks to man of exposure to these chemicals has been a major focus for research (Lake et al., 2000).

Section 1.5 Gene expression in response to peroxisome proliferators

PPAR α forms a heterodimer with the retinoid x receptor α (RXR) -after being provoked by the peroxisome proliferators- which then binds to precise regions on the DNA (called peroxisome proliferator hormone response elements; PPREs) of the targeted genes (Tugwood et al., 1992) as shown in Figure 1.6.

The PPAR α -RXR α heterodimers attach to DNA sequences with repeats of the sequence AGGTCA (AGGTCA X AGGTCA) separated by one nucleotide (Jpenberg et al., 1997). This heterodimer would be able to stimulate transcription of a large number of genes, including those involved in lipid metabolism such as cytochrome P450 4A and acyl CoA oxidase.

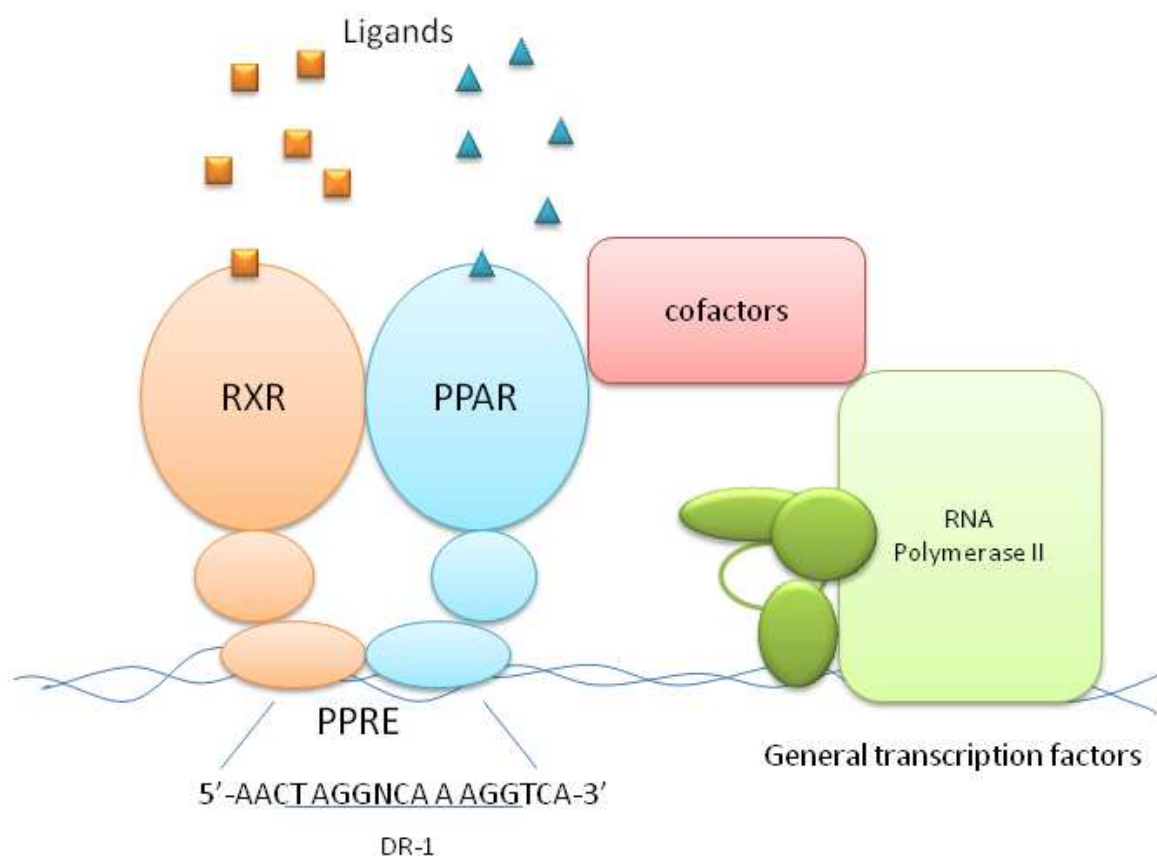


Figure 1.6 Schematic diagram of the PPAR mechanism of action. The blue triangles are peroxisome proliferator that bind to the PPAR, leading to heterodimer with retinoid x receptor (RXR) which has been activated by its ligands (orange squares). The PPAR/RXR heterodimer binds within the DNA binding C domain to peroxisome proliferator hormone response elements (PPRE) through a DNA sequence with repeats of the sequence AGGTCA (AGGTCA X AGGTCA) separated by one nucleotide (this would be located at the promoter of the target gene). The diagram is a modified version from (Keller et al., 2000).

Several studies showed that peroxisome proliferators are involved in the regulation of several genes that are involved in hepatic proliferation, metabolism, immune modulation, mitochondrial and peroxisomal fatty acid β -oxidation (Desvergne and Wahli, 1999) (Latruffe and Vamecq, 1997). Other studies reported that ciprofibrate is involved in modulating the expression of genes implicated in cell proliferation such as cyclins (Rininger et al., 1996)(Peters et al., 1998).

Yadatie examined the effects of 50 mg kg⁻¹ body weight of ciprofibrate per day for 60 days on gene expression in the liver of rats by using a cDNA microarray analysis. From this it was found

that a number of up-regulated genes were involved in lipid and sugar metabolism while other genes were implicated in growth and stress responses (Yadatie et al., 2003).

Peroxisome proliferators are reported to induce the transcription of the *CYP4A1* gene (Hardwick et al., 1987), and the induction of *CYP4A1* is linked with the transcription of genes encoding the 3-oxoacyl CoA thiolase and the bifunctional enzyme acyl-CoA (*ACO*) (Furuta et al., 1982)(Reddy et al., 1986).

Bell also characterized the induction of *CYP4A1* and acyl-CoA *in vitro* and found them highly induced after dosing rat hepatocytes with clofibric acid for 4 days, or with methylclofenapate, nafenopin, clofibric acid or mono (ethylhexyl) phthalate for 1 h and 8 h. This suggested that the different time course of induction of *CYP4A1* and acyl-CoA oxidase is not related to the type of peroxisome proliferator but to the process of peroxisome proliferation (Bell and Elcombe, 1991a).

In vivo studies in rats dosed intraperitoneally with methylclofenapate also showed significant induction of *CYP4A1* after 6 h, 8 h, 24 h and 30 h, while the acyl-CoA showed no increase at 8 h but increased significantly after 24 h (Bell et al., 1991). As the induction of the *CYP4A1* gene is the first sign of the peroxisomal reaction, it is possibly the most significant enzymic indicator/marker of peroxisome proliferation (Orton and Parker, 1982)(Hardwick et al., 1987).

In other reports the effects of peroxisome proliferators on immediate early gene expression have been related to the extracellular signal-regulated kinases and phosphorylation of the upstream regulator MAPK/ERK kinase signal transduction pathway (Rokos and Ledwith, 1997).

As PPARs can induce the regulation of a number of genes, PPARs can also down-regulate gene expression as they are necessary for the suppression of apolipoprotein CIII mRNA expression,

but the mechanism is not clearly understood (Peters et al., 1997a). CYP2C11 expression is clearly down-regulated by peroxisome proliferators as reported in (Corton et al., 1998).

Gonzalez and his group demonstrated that hepatic microRNA expression was largely activated by PPAR α . They studied a microRNA involved in cell growth, *let-7C*, and found it was suppressed following 4-h treatment, 2-week or 11-month constant treatment with the potent PPAR α agonist Wy-14,643 in wild-type mice. *let-7C* was shown to target *c-myc*. The induction of *c-myc* through *let-7C* consequently amplified expression of the oncogenic mir-17-92 cluster; this did not arise in PPAR α null mice (Shah et al., 2007a).

(Suzuki et al., 2010) examined the effect of Wy-14,643 on DNA damage in rat livers treated orally for 14 days and found that a number of DNA repair genes were induced such as *Gadd45*, *Apex 1*, *Xrcc5* and *Mlh1*. These results implied that hepatic DNA was damaged by Wy-14,643 but might be repaired via the activation of these DNA-repair genes.

The development of microarray tools has significantly changed gene expression analysis research. However, the reliability and reproducibility of RNA microarray data can be challenged. Therefore, confirmation of results with quantitative real-time PCR or a similar technique is essential (Baker et al., 2004). Baker found using cDNA microarray analysis and confirmation with real-time PCR that in clofibrate treated rats, hepatic expression of acyl CoA, topoisomerase II- α and CYP4A1 were highly induced. Another tool used recently to study global gene expression is the RNAseq analysis technique. Also known as illumina sequencing, this technology is still very expensive and needs experience in bioinformatics, but it is fast and the illumina sequencing data is highly reproducible, with reasonably low technical variation in enabling identification of differentially expressed genes (Marioni et al., 2008).

Section 1.6 The aim of the Thesis

The overall aim of this thesis is to shed some light on the mechanism of action of hepatic growth induced by peroxisome proliferators activating receptor ligand (PPAR α), specifically ciprofibrate. This work is based on the assumption that increased DNA synthesis is causally related to the development of liver cancer.

To investigate how peroxisome proliferators induce hepatocyte proliferation and which genes they induce to cause cancer, the specific aims were;

- 1- To indicate when the induction of hepatic DNA synthesis provoked by ciprofibrate happens in mice and rats, by immunohistochemical technique.
- 2- To investigate where in the liver tissue the induction of hepatic DNA synthesis occurs in mice and rats dosed with ciprofibrate.
- 3- To characterize the genes responsible for triggering the induction of hepatic DNA synthesis caused by ciprofibrate in comparison with cyproterone acetate (CPA) by studying the immediate early induced genes. This was done with cDNA microarray, Transcriptome sequencing analysis and Quantitative Real Time PCR.

Chapter 2 Materials and Methods

Section 2.1 Materials

Section 2.1.1 Animals

Male F-344/NHsd (Fisher) rats (14-15 weeks, 260±20g) were bought from Harlan Laboratories, Inc. UK. DBA/2JCr1 (22±2g) and C57BL/6Cr1 (20±2g) mice were used at age 9-10 weeks, and were purchased from Charles River Laboratories, Inc. UK.

Animals were matched for sex, strain, supplier and age, and were randomized on arrival on the basis of body weight in to the appropriate number of groups.

Animals were maintained under specific pathogen-free conditions, in plastic cages and were kept at 24±4°C. The humidity was 70±5%, with a 12 h day/night light cycle, with food and water available ad libitum throughout the experimental period.

The animals were humanely killed with a single overdose of Dolethal (pentobarbital 200 mg ml⁻¹, purchased from Vetoquinol Company). The animals' experiments were performed in accordance with the Home Office guidance (Scientific Procedures) Act 1986.

Section 2.1.2 Chemicals

The peroxisome proliferator used was ciprofibrate as a powder and was a generous gift from Dr. T. J. B Gray, Sanofi- Aventis (Alnwick, UK).

The pregnane X receptor agonists used were cyproterone acetate CPA and pregnenolone-16 α -carbonitrile PCN, and were bought from Sigma-Aldrich® as powders.

Section 2.1.3 Immunohistochemistry materials

All chemicals used were of the highest quality available. 5-bromo-2'-deoxyuridine (BrdU), 3,3'-diaminobenzidine tetrahydrochloride (DAB), bovine serum albumin (BSA), polyoxyethylene sorbitane monolaureate (Tween-20) and 10% (w/v) formalin were purchased from Sigma-Aldrich®.

A primary anti-BrdU mouse monoclonal antibody was purchased from GE healthcare (UK), and a secondary antibody (blotting grade affinity-purified goat anti-mouse IgG (H+L) horseradish peroxidase conjugate) was purchased from Bio-Rad.

Ultrapure tris base was obtained from Melford Laboratories Ltd. DPX (Distyrene, plasticiser and xylene), hydrochloric acid and 30% (w/v) hydrogen peroxide were obtained from Fisher Scientific, ammonium hydroxide from Aldrich, and cobalt chloride from AnalaR. Amersham cell proliferation kits were purchased from GE healthcare (UK).

Poly-L-lysine-coated slides (Polysine™) were obtained from Fisher, pure paraffin wax (melting point 56°C) from RA lamb, Peel-A-Way® disposable histology molds from Polysciences Inc, and TAAB embedding stubs (25mm) from TAAB laboratory and microscopy.

Glacial acetic acid, xylene, ethanol, methanol, sodium phosphate and haematoxylin stain were purchased from Sigma-Aldrich®. Pure water was produced in this laboratory at a quality of < 0.2 µS.

Section 2.1.4 Microarray materials

Section 2.1.4.1 RNA isolation materials

TRI reagent® solution (a mixture of guanidine thiocyanate and phenol in a mono-phase solution) and 1-bromo-3-chloro-propane, Isopropanol and ethanol were bought from Sigma-Aldrich®. SDS Sodium lauryl (dodecyl) sulfate ($C_{12}H_{25}SO_4Na$) and ethidium bromide were bought from Fisher Scientific.

Section 2.1.4.2 Synthesis and labelling of cDNA

cDNA synthesis and labelling was with an amino allyl cDNA labelling kit from Ambion the RNA Company. Reagents used and included in the kit for cDNA labelling reactions are, 10X RT buffer, Oligo(dT) Primers [Oligo(dT)18, 50 mM] random decamers (50 mM), RNase inhibitor (10 units/ml), dNTP Mix (no dTTP) (10 mM each: dATP, dCTP, dGTP, dTTP)+ AA dUTP Mix, 3 mM dTTP, 3 mM 5-(3 aminoallyl)-2'-dUTP.

Also M-MLV reverse transcriptase (200 units/ml), 3 M sodium acetate (pH 5.5), glycogen (5 mg ml⁻¹), 1 M sodium hydroxide, DMSO (100% dimethyl sulfoxide), coupling buffer, 1 M HEPES (pH 7), 4 M hydroxylamine, water for 75% ethanol, 100% ethanol and nuclease-free Water.

The fluorescent dyes used for the labelling were Alexa Fluor® 555 and Alexa Fluor® 647 reactive dyes from Invitrogen.

Section 2.1.4.3 Microarray slides and hybridization materials

Whole genome mouse Mouse Exonic Evidence Based Oligonucleotide (MEEBO) array slides were used. The arrays were printed over 2 slides (A+B), and were accompanied with a specific

GenePix Array List (GAL) file.

The array slides were Gentix Aldehyde Plus arraying slides and were a kind gift from Dr T. Gant's lab, Systems Toxicology Group, Leicester University.

For the microarray hybridization of the cDNA, the tRNA used was from Invitrogen and 2X-enhanced cDNA hybridization buffer from Genisphere.

Section 2.1.4.4 Microarray scanner

An Axon 4200 scanner and GenePixPro(6) program was used to scan and analyse the slides. Both were situated at Dr T. Gant's laboratory, Systems Toxicology Group, Leicester University.

Section 2.1.5 Transcriptome sequencing materials

RNA sequencing was done with the AB SOLiD 3 platform, using short reads, at Dr Aziz Aboobaker's lab, Next Generation Sequencing Facility, Queens Medical Centre (QMC), University of Nottingham. Associated equipment for the transcriptome analysis was a COVARIS S2 sonicator, a DIGILAB hydroshear nanodrop 2000 and 3000 and a Qubit nucleic acid quantification agilent bioanalyser.

Section 2.1.6 Quantitative real- time PCR materials

Section 2.1.6.1 Synthesis of cDNA

Total hepatic RNA was reverse transcribed to first strand cDNA using a High Capacity RNA-to cDNA kit, and TaqMan® Gene Expression Master Mix which were from Applied Biosystems.

Section 2.1.6.2 Primers and probes

Real- time RT-PCR primers and probes for rat genes were designed using PRIMER 3 (Skaletsky, 2000) at <http://frodo.wi.mit.edu/primer3/>.

The primers and probes were checked for hairpins and dimers with AutoDimer v1 software and the primers were checked for specificity by Blast searching against the rat genome using the standard rodent database, at <http://www.ebi.ac.uk/Tools/blastall/nucleotide.html>.

The probes and primers were synthesized by Eurofins, MWG Operon. The sequence of the primers and probes and the dyes and quenchers that they are attached to are listed in Table 2.1.

Table 2.1 RT-PCR oligonucleotides primers and probes for rat genes. Primers and probes are designated by letters indicating the forward primer F, the reverse primer R or the probe P. Sequences are given from 5'-->3'. The reporter dye is at the 5' end of the oligonucleotide. At0647N was used as an alternative to Cy5 where indicated.

Rat gene	Oligonucleotide sequence	Genebank accession no.	Reporter dye
<i>AhR</i> F R P	GCA GCT TAT TCT GGG CTA CA CAT GCC ACT TTC TCC AGT CTT A TAT CAG TTT ATC CAC GCC GCT GAC ATG	Af082124	HEX- BHQ1
<i>β-actin</i> F R P	CTG ACA GGA TGC AGA AGG AG GAT AGA GCC ACC AAT CCA CA CAA GAT CAT TGC TCC TCC TGA GCG	V01217	Cy5-BHQ2
<i>Ccnd1</i> F R P	GCG TAC CCT GAC ACC AAT CT GGC TCC AGA GAC AAG AAA CG CTG GAT GCT AGA GGT CTG CGA	NM_171992	HEX-BHQ1
<i>CYP3A1</i> F R P	AGT GGG GAT TAT GGG GAA AG CAG GTT TGC CTT TCT CTT GC TAG AGC CTT GCT GTC ACC CA	NM_013105	FAM-BHQ1
<i>CYP4A1</i> F R P	TCA TGA AGT GTG CCT TCA GC TGT GTG ATC ATG GGC AAG TT ATC CAG GCC ATT GGG AAC TT	NM_175837	FAM-BHQ1
<i>G0s2</i> F R P	GGT GTG GTG CTC GGT CTA GT ACA AAG TCG CCT CCT GTG TC CAG GCC CTG ATA GCA GAA GG	NM_0010096 32	At0647N- BHQ2
<i>Scd1</i> F R P	TCC TGC TCA TGT GCT TCA TC GGA TGT TCT CCC GAG ATT GA TAC TGC TGG GGC GAA ACT TT	NM_139192	HEX-BHQ1

Section 2.1.7 Software used in data analysis

The NORT.T normalizing program was obtained from the Systems Toxicology Group, Leicester University (Dr Shu-Dong Zhang and Dr Timothy W. Gant) and was used for the initial normalizing of the scanning results and the microarray comparison experiments. The program can be downloaded from http://143.210.176.81/SystemsToxicology/Microarray_Softwares/Softwares_NorTT.aspx

The microarray experimental information [(in accordance with Minimum Information About a Microarray Experiment (MIAME))] was stored in ArrayTrack, a software system for managing and interpreting microarray gene expression data developed by the Center for Toxicoinformatics at the National Centre for Toxicological Research (NCTR) of the U.S. Food and Drug Administration (FDA). The database generated from this thesis is accessible under the University of Nottingham, file ABEER, from http://edkb.fda.gov/webstart/extdb_arraytrack/3.4/. R software version 2.9.0 (R foundation for statistical computing) was used to analyse the final results for the microarrays.

The data from the RNA sequencing facilities (University of Nottingham Next Generation Sequencing Facility) were analysed with CLC Genomics Workbench 3.7 software. The original data can be found at http://spldeepseq.nottingham.ac.uk/~aziz/wtp_rat/.

The rat genome used was from Ensembl at http://www.ensembl.org/Rattus_norvegicus/Info/Index, and the translation to gene names was done with rat genome database RGD tools at <http://rgd.mcw.edu/rgdweb/search/genes.html?100>. Microarray heatmaps were established with High-Throughput GoMiner (Zeeberg, 2005). This software uses just + or - so it does not show the differences in between but it gives the main pathways.

Section 2.2 Methods

Section 2.2.1 Dosing of animals

The chemicals were administrated to the animals in different dosing schedules depending on the study, ensuring that the least suffering and stress was applied to the animals at all times.

The dosing schedules were:

- 1- Acute dosing schedule.
- 2- Chronic dosing schedule.
- 3- Gavage dosing schedule.

Section 2.2.1.1 Acute dosing schedule

In the acute study 100 mg kg⁻¹ 5-bromo-2'-deoxyuridine (BrdU) was given to the animals by intra peritoneal (IP) injection, in a volume of 5 ml kg⁻¹, two hours prior to killing the animals.

The BrdU was dissolved in phosphate buffer saline (PBS; 15mM NaH₂PO₄, 150mM NaCl, pH adjusted to 7.4 with NaOH) then sterilized with a micro-filter (pore size of 0.2 μm).

All the animals were humanely killed with a single intraperitoneally overdose of Dolethal and death was confirmed by cervical dislocation of the neck. Figure 2.1 represents a cartoon of the experimental schedule of the acute dosing protocol.

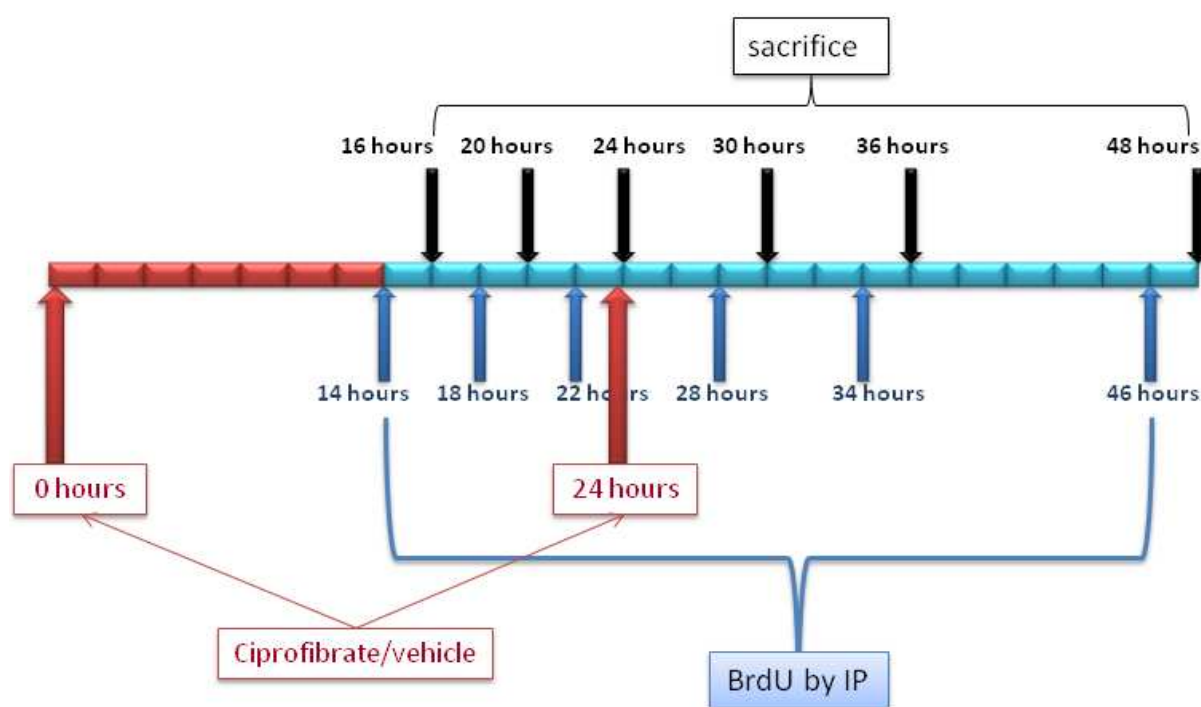


Figure 2.1 Schematic representation of acute and gavage dosing protocols. Cartoon describes the acute dosing schedule for studying the time course of induction of hepatic DNA synthesis. The blue arrows represent the number of hours when the acute IP dosing of the BrdU (100mg kg⁻¹ body weight) accrued. The black arrows show the time of scarifying the animals 2 hours after the BrdU. The red arrows indicate the time of gavage dosing of the drug or vehicle at 0 or 24 hours. Each sequence of the time course line specifies a 2 hour time limit, the red is the time before the BrdU started.

Section 2.2.1.2 Chronic dosing schedule

The dosing of the BrdU to the animals in the chronic experiments was carried out by acclimatizing them to 10% (v/v) Sainsbury's orange juice as their sole source of fluid for 7 days then on orange juice supplemented with 0.08% (w/v) BrdU. All drinking bottles containing BrdU were covered with aluminum foil to protect them from light, and were changed on a daily basis.

All the animals were humanely killed with a single intraperitoneally overdose of Dolethal and death was confirmed by cervical dislocation of the neck. In Figure 2.2 and Figure 2.3 a cartoon is illustrated showing the experimental schedule of the chronic dosing protocol for the testing of the effects of BrdU (Figure 2.2) and for the experimental design of the chronic dosing schedule.

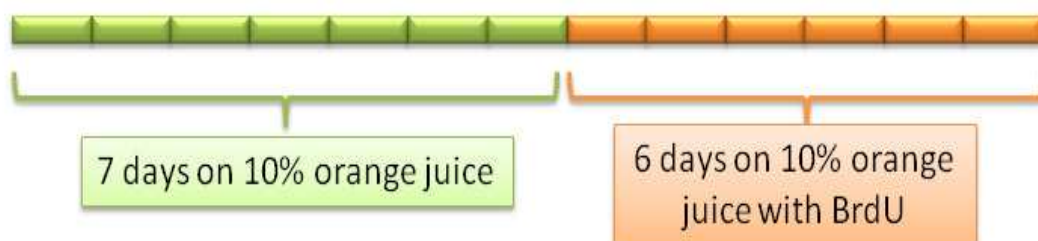


Figure 2.2 Schematic representation of chronic dosing protocol alone. Cartoon describes the chronic dosing schedule for studying the BrdU effects in mice. The green sequences show the number of days the animals were given 10% orange juice as sole source of drinking fluid. The orange sequences represent the number of days the animals had BrdU with the orange juice as their sole source of drinking fluid. Each sequence of the time course line specifies a 1 day time limit.

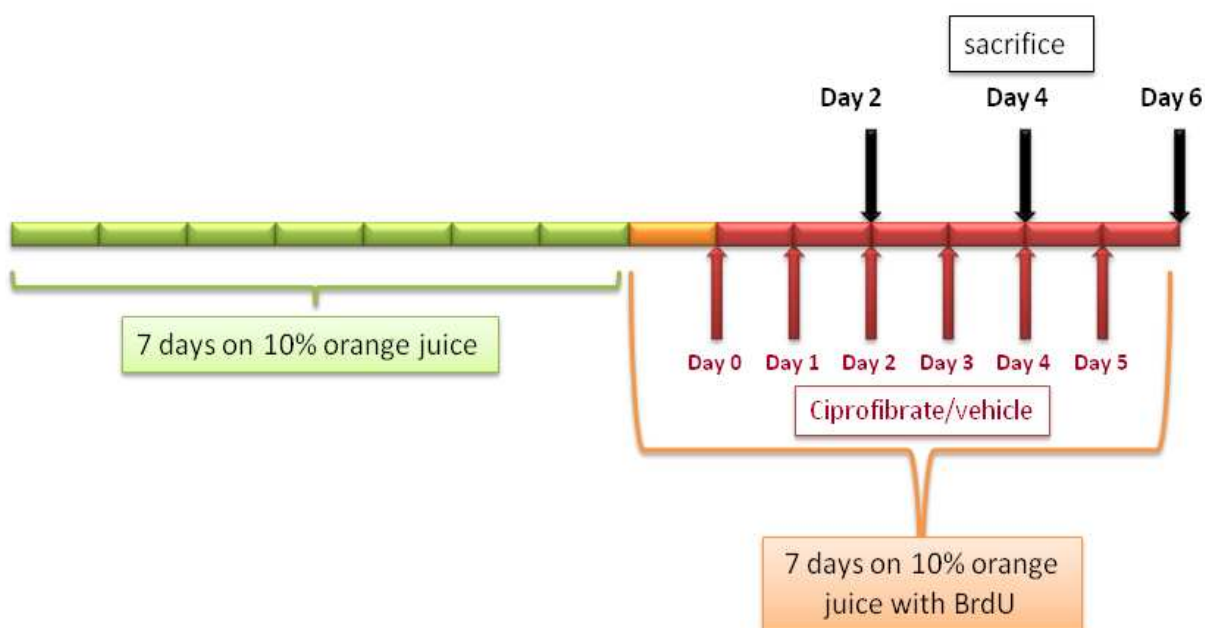


Figure 2.3 Schematic representation of chronic and gavage dosing protocols. Cartoon describes the chronic dosing schedule for studying the time course of induction of hepatic DNA synthesis. The green sequences show the number of days the animals were given 10% orange juice as their sole source of drinking fluid. The orange sequence represent the number of days the animals had BrdU with the orange juice as their sole source of drinking fluid. The red sequences show the number of days the animals had started the ciprofibrate with the BrdU in the orange juice as their sole source of drinking fluids. The red arrows indicate the time of gavage dosing of the drug or vehicle and the black arrows show the time of scarifying the animals. Each sequence of the time course line specifies a 1 day time limit.

Section 2.2.1.3 Gavage dosing schedule

The ciprofibrate, cyproterone acetate (CPA), pregnenolone-16 α -carbonitrile (PCN) in the time course was 50mg kg⁻¹ ciprofibrate for 1, 3, 5, 16, 20, 24, 30, and 48 hours in the rat study and 100mg kg⁻¹ for 2, 4 and 6 days in the mouse study. For the CPA and PCN the dose was 100mg kg⁻¹ for 24 hours. The doses for the dose response was 50, 100, 200, and 300mg kg⁻¹ after 24 hours.

These were administrated to the animals in corn oil as a vehicle by a gavage injection that enables the drug to be administrated straight to the stomach.

Section 2.2.2 Animal observations and tissue collection

The animals were observed 2-3 times daily for signs of distress or discomfort. Daily measurements of body weight were recorded at the same time (10-11am) each day, to monitor the well-being and health of the animals.

Immediately after killing the animals, blood samples were collected in plain tubes by cardiac puncture. After the blood had clotted (2-4 hours) it was centrifuged at 14000 Xg and the serum frozen for the clinical chemistry procedures.

The liver weights of the animals were recorded and the percentage of the body weight to liver weight calculated. Aliquots of fresh liver and a part of the intestine (as positive control) were fixed for immunohistochemical analysis (formalin fixation), or snap frozen in liquid nitrogen for storage at -80°C prior to biochemical analysis.

Section 2.2.3 Immunohistochemistry technique for BrdU paraffin sections

This method was used to determine the percentage of BrdU-labelled cells during induction of DNA synthesis in the animals' liver as a result of the ciprofibrate's or cyproterone acetate's action. The two-step indirect method was originally optimized by (Al Kholaiifi, 2008).

In this method the thymidine analog BrdU was dosed to the animals orally or injected.

Section 2.2.3.1 Tissue processing

Tissues were freshly collected from the animals. The livers and a section of the intestine were stored in a fixative 10% (v/v) formalin until used. The tissues were left in the fixative not more than 3 days. Fixed tissues were dehydrated and infiltrated with paraffin wax by processing in a Shandon Citadel 2000 Automated Tissue Processor (protocol shown in Table 2.2).

Table 2.2 Protocol for tissue processing.

Solution	Time
70% ethanol	6.5 hours
80% ethanol	1 hour
90% ethanol	1 hour
95% ethanol	1 hour
100% ethanol	1 hour
100% ethanol	1 hour
100% ethanol	1 hour
xylene	1 hour
xylene	1 hour
xylene	1 hour
paraffin wax	1 hour
paraffin wax	1 hour

After the tissue processing, the tissues were manually embedded in paraffin wax, and left on ice for at least 48 hours before sectioning with a microtome set to 4.5 μ m.

The sections were floated on a preheated water bath to 37°C before mounting on Polylysine™ adhesive coated glass slides. The slides were then left overnight on a hot plate at 37°C to dry.

Section 2.2.3.2 BrdU immunohistochemical staining

Immunohistochemical staining was performed by heating the sections for 2-3 minutes or until the wax was melted, then placing them in xylene for 5 min. The sections were rehydrated in a decreasing series of ethanol concentrations (100%, 70%, 50%) and then washed in water. The slides were then treated with fresh 3% (v/v) hydrogen peroxide in methanol to quench the endogenous peroxidase, and then washed twice with PBS.

To retrieve the BrdU antigen a heat induced epitope retrieval technique was used, where the slides were plunged in boiling 10mM citric acid followed by 10 min. at 60°C, using a 700 Watt-microwave (heating induces epitope retrieval by breaking the DNA, and unmasking the BrdU). Slides were then washed with PBS.

The slides were incubated horizontally, and in a warm, moist chamber with a 0.5% (w/v) bovine serum albumin (BSA) blocking solution to block non-specific binding of cytosolic BrdU. Then the slides were incubated for 45 min with Amersham mouse monoclonal anti-BrdU antibody [1:1000 dilution in tris buffer saline (TBS)], then washed with distilled water.

The secondary antibody, blotting grade goat anti-rabbit IgG horseradish peroxidase conjugate, was added for 30 min. The slides were then rinsed in PBS.

The slides were returned to a vertical orientation for incubation in the staining solution DAB reagent, before being washed with water.

The slides were then subjected to a standard histological staining procedure as shown in Table 2.3, where the slides were dipped in haematoxylin solution, washed, dipped in acid alcohol, then rinsed in running tap water, then in ammonia in water, and washed again in tap water. Then for the final dehydration of the slides, they were placed in an increasing series of ethanol (50%,

70%, 95%, 100%).

Finally the slides were treated with 100% xylene, and then mounted with DPX and covered with a cover slip each to protect the tissue from shrinking.

The slides were ready to be examined, preferably after 12-24 hours so the slides were completely dried.

Table 2.3 Protocol for BrdU immunohistochemical staining.

Application	Preparation of Application	Time
Heat	heat the slides until the wax melts	2-3 minutes
Xylene	Xylene	5 minutes
Ethanol	100%-70%- 50%	8- 4- 3 minutes respectively
H ₂ O	H ₂ O	2 minutes
H ₂ O ₂	3% H ₂ O ₂ (180ml methanol+20ml hydrogen peroxide)	15 minutes
Phosphate buffer saline PBS	0.1M is 11.5g Na ₂ HPO ₄ di-sodium hydrogen orthophosphate 2.96g NaH ₂ PO ₄ .2H ₂ O sodium phosphate and 5.84g sodium chloride NaCl in 1litre distilled water	5 minutes twice (X2)
Citric acid	10mM 2.1g citric acid in 1L dH ₂ O, adjust pH at 6.0 using 1N sodium hydroxide NaOH.	Boiled citric acid 10 minutes at 40°C
PBS	Phosphate buffer saline (as before)	5 minutes
Bovine serum albumin	BSA 5% (blocking solution) is 5g Bovine serum albumin, 0.5ml Tween-20 in 100ml PBS.	15 minutes
Primary antibodies	(anti-BrdU mouse monoclonal antibody from Amersham). Freshly prepared 1:1000 dilution buffer is 3g BSA+100µl Tween-20 in 100ml TBS (Tris buffer saline) pH 7.8.	45 minutes
H ₂ O	H ₂ O	5 minutes

Table 2.3 Protocol for BrdU immunohistochemical staining.

Secondary antibodies	(peroxidase anti-mouse IgG from Bio-rad) freshly prepare 1:100/slide. Dilution buffer is 0.087g monosodium phosphate monohydrate $\text{NaH}_2\text{PO}_4 \cdot \text{H}_2\text{O}$, 0.194g disodium phosphate (anhydrous) Na_2HPO_4 , 1.75g sodium chloride NaCl and 2g BSA in 200ml distilled H_2O	30 minutes
PBS	Phosphate buffer saline (as before)	9 minutes
Staining solution DAB	(development reagent is 200 μl hydrogen peroxide+ A + 4ml of B) A development reagent is 176ml TBS 10mM Ph 7.6 (1.21g tris base+5.64g NaCl , adjust PH to 7.8 using HCl in 1 liter distal water) with 20ml intensifier 0.25% cobaltous chloride. B is the DAB 25mg/ml stock solution, and is 0.5g DAB powder in 20ml phosphate buffer.	8 minutes
H_2O	Distal H_2O	3 minutes
Haematoxylin solution.	(Harris) Haematoxylin 100ml+glacial acetic acid 4ml, filter before use.	30 seconds
H_2O	Running tap water	3 minutes
Acid alcohol	(for differentiation) 700 ml commercial grade ethanol with 300ml distilled water and 10ml concentrated hydrochloric acid.	5 seconds
H_2O	Rinse in running tap water	1 minute
Ammonia	0.2% Ammonia in water	1 second
H_2O	Tap water	3 minutes
Ethanol	Ethanol 50%-70%- 95%-100%	1 minute each
Xylene	Xylene (X2)	1 minute each
DPX	Mount with DPX (Distyrene, plasterine, and xylene) and cover with cover slide.	~1 minute each slide

Section 2.2.4 Examination of the slides

The immunohistochemically stained slides were examined in one of two ways, depending on the study: 1- Labelling index studies

2- Zonal distribution studies

Section 2.2.4.1 Labelling index studies

To determine the hepatic DNA labelling index, one slide (with a slice of liver and gut) from each animal was used and 2000 nuclei were examined from random fields of view of each slide (the mean was calculated as the number of animals). Nuclei labelled with BrdU give a dark black/brown colour (Figure 2.4) indicating that the cell has undergone DNA division. These cells were also counted as well as non-labelled nuclei. The percentage of the labelled nuclei was then calculated. A CX21 Olympus microscope was used with a graticule eye piece. The magnification used was X400.

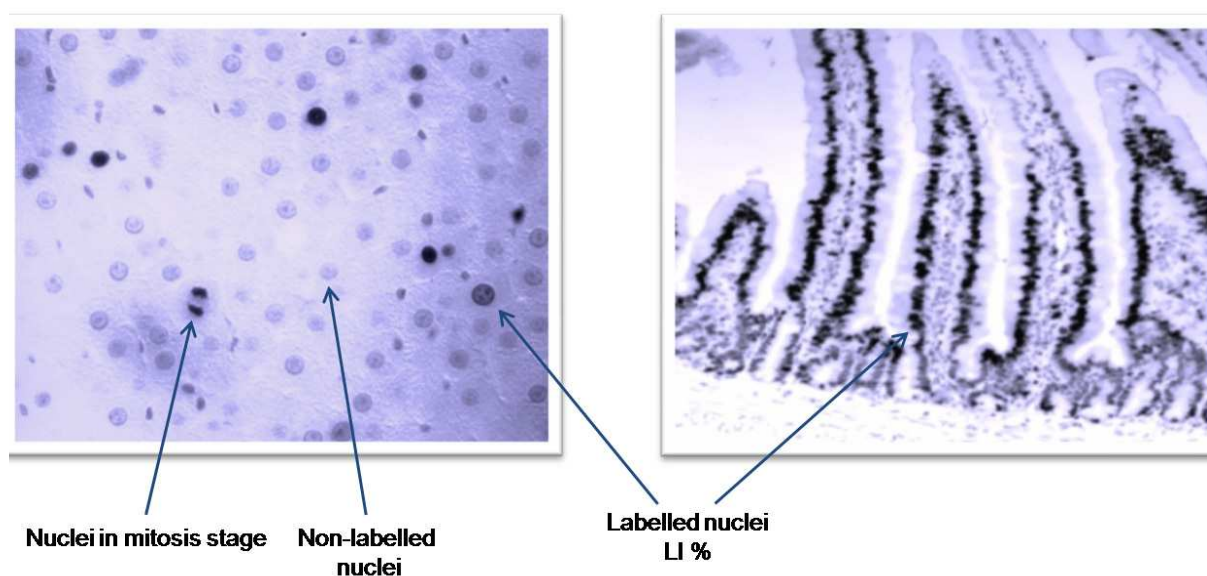


Figure 2.4 BrdU labelled hepatocytes. On the left a section of the liver with non-labelled and labelled hepatocytes. On the right a section of the intestine used as a control and shows the labelled epithelium cells in a dark colour and the non-labelled cells are pale at the top. The magnification of the liver is X400, and of the intestine is X100

Section 2.2.4.2 Zonal distribution studies

To determine the zonal distribution of the BrdU-labelled hepatocytes; the number of labelled and non-labelled nuclei per field was counted.

A field is defined from (Barrass et al., 1993) as a radius of five to seven cells around either the portal space, which is the periportal region, or the central vein which is the perivenous region (Figure 2.5).

Ten random areas of both periportal and perivenous regions were counted (five each). Small central veins or portal spaces of similar sizes were selected for the analysis of the perivenous or periportal. One slide per animal was used. The total number of labelled nuclei in five fields was recorded for each of the periportal and perivenous zones.

An Olympus CX21 reflected light upright microscope was used at X400, with a graticule eye piece.

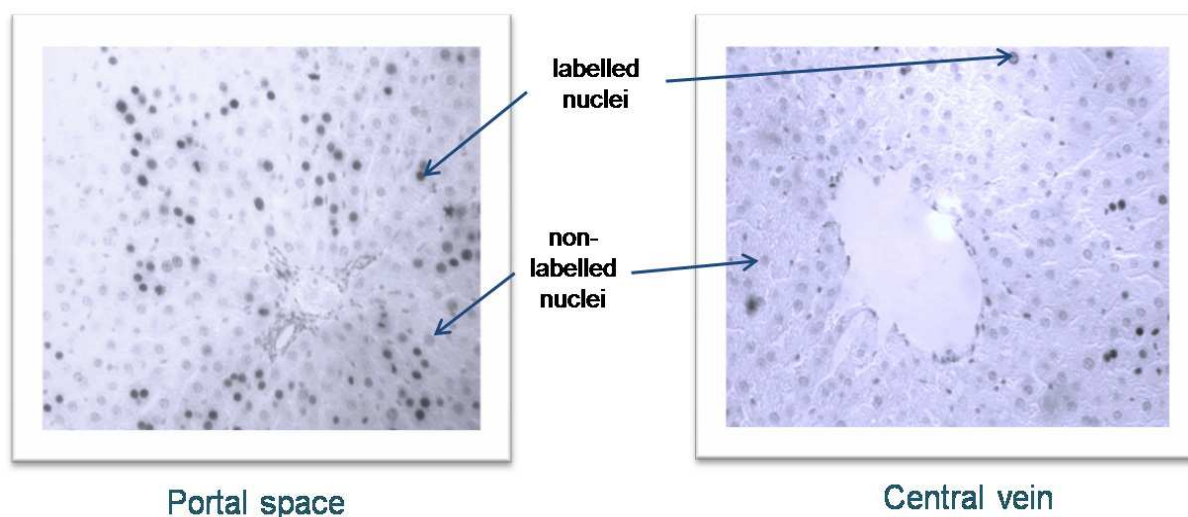


Figure 2.5 The periportal and perivenous zones. On the left is a portal space containing the portal vein, the hepatic artery and the bile duct and the surrounding hepatocytes which are the periportal zone. On the right is the central vein with the surrounding hepatocytes which count as the perivenous zone. The magnification is X100.

Section 2.2.5 Microarray analysis

Section 2.2.5.1 RNA isolations

Total RNA was isolated using TRI reagent® solution from frozen liver cells with the following procedure:

TRI reagent® solution (1 ml) was added to 80-90 mg of frozen liver tissue pieces in a 1.5 ml Eppendorf tube, and homogenized with a mini homogenizer for 2-3 min. The mixture was left to sit at room temperature for 5 min, before 200 µl 1-bromo-3-chloro-propane (Sigma) was added and the mixture shaken vigorously for 15 seconds.

Then the samples were vortexed on full speed for 5-10 seconds, and incubated for 2-3 min at room temperature. The samples were centrifuged at 14000 Xg for 14 min, at 4°C.

The upper layer was carefully transferred to a new RNase-free 1.5ml Eppendorf tube and 600µl isopropanol was added. After mixing, the tube was incubated at room temperature for 10 min.

The samples were centrifuged at 14000 Xg for 10 min to pellet the RNA. All subsequent procedures were performed at 4°C (on ice). The supernatant was removed carefully, then the pellet was washed twice with 0.5 ml ice cold 75% ethanol, centrifuging between each wash step.

The pellet was resuspended in ~50-100µl diethyl pyro carbonate (DECP) water and stored at -80°C until use.

Note; DECP water was made by mixing 1ml DECP (diethyl pyro carbonate) and 9 ml pure ethanol (absolute) in 1 litre distilled water, and then autoclaved.

Filter pipette tips and clean gloves were used for all the procedures. The gloves were changed on a regular basis.

Section 2.2.5.2 Synthesis and labelling of cDNA from RNA

This was done with a modified technique, using an Amino allyl cDNA labelling Kit Ambion® the RNA company.

To reverse transcribe the RNA, 1µl RT primer was added to around 20µg total RNA and denatured at 75°C for 9 min. Added to this was 2µl 10X RT buffer, 1µl RNase inhibitor, 1µl dNTP mix (dATP, dCTP, dGTP), 1µl dTTP+AA dUTP mix and 2µl M-MLV reverse transcriptase.

The mixture was incubated at 42°C for 1 h and 45 min, then 4µl 1M NaOH was added before incubation at 65°C for 15 min to remove the template RNA by alkaline hydrolysis. The reaction was then neutralized with 10µl 1M HEPES buffer.

The cDNA was recovered by ethanol precipitation, by incubating with 3.4µl 3M sodium acetate and 100µl 100% ethanol at -80°C for 1-2 h.

The cDNA was pelleted by centrifugation at 14000 Xg at 4°C for 15 min, the supernatant was carefully aspirated and discarded, and then the cDNA pellet was washed by adding ~0.5 ml of 75% (v/v) ice cold ethanol and vortexed briefly, before centrifuged for ~5 min at 14000 Xg at 4°C, and carefully removed and the supernatant discarded.

To remove the last traces of ethanol the tube was re-centrifuged for ~5 min at 14000 Xg at 4°C, and a small sized pipet was used to remove the residual fluid.

The cDNA was then resuspended in 4.5µl coupling buffer and 2.5µl nuclease-free water, mixed thoroughly and gently vortexed, and then the tube was centrifuged briefly.

To couple dye to the amino modified cDNA, 3µl fluorescent dyes Alexa Fluor® 555 or Alexa Fluor® 647 were dissolved in 3µl 100% DMSO, then the dyes were added to the cDNA and

incubated in the dark at room temperature for 1 hour.

6 μ l of 4M hydroxylamine was added to the mixture, before incubation for 15 min at room temperature in the dark to terminate the coupling reaction.

The labelled cDNA was purified using rehydrated NucAway spin columns after hydrating the columns with 650 μ l nuclease free water for 1-2 h just before use. The cDNA was transferred on to the matrix of the NucAway spin columns and centrifuged at 750 Xg for 2 min (according to the manufacturer's instructions).

The dyed -labelled cDNA was run through the column to a 1.5 ml Ependorff tube positioned beneath the column and the free dye was retained in the spin column matrix.

The dye-labelled cDNA was prepared for concentration by ethanol precipitation by adding 9 μ l 2M sodium acetate and 250 μ l 100% ethanol and storage at -80°C overnight.

The next day the labelled cDNA was centrifuged for 15 min at 14000 Xg at 4°C then the supernatant was carefully removed, and the dye-labelled cDNA pellet was washed by adding ~0.5 ml of 75% (v/v) ice cold ethanol and vortexed briefly.

The tube was then centrifuged for ~5 min at 14000 Xg at 4°C, and carefully removed and the supernatant discarded. To remove the last traces of ethanol the tube was re-centrifuged for ~5 min at 14000 Xg at 4°C, and then with a small sized pipet the residual fluid was removed.

The cDNA was a small pellet, 1-2 mm in diameter, which was visibly red for the Alexa Fluor® 555 or blue for the Alexa Fluor® 647.

The dye-labelled cDNA was suspended in 25-40 μ l RNA-free water and the success of the labelling was assessed with a nanodrop machine to measure the optical density (OD) of the cDNA

and the labelled dyes.

Note that all the labelled fluorescent steps were done in the dark.

Section 2.2.5.3 Hybridization of the cDNA

The dye-labelled cDNA was mixed with 2 μ l tRNA (4ml ml⁻¹) and 40-80 μ l hybridization buffer and centrifuged briefly to collect the mix, then denatured for 5 min at 100°C and incubated for 1 h at 42°C.

The MEEBO array slides and the cover slips were prepared for use by washing the oligo arrays twice in 0.2% (w/v) sodium dodecyl sulfate (SDS) for 2 min, then twice in ddH₂O for 2 min each, and then dried by centrifugation at 250 Xg for 4 min.

The cover slips were washed in 1% (w/v) SDS for 30 min and then underwent 5 X 5 min washes with dH₂O, before being spun dry/wiped dry and stored in a dust free environment.

The cDNA was spread on the MEEBO array slides by placing the slide over a template slide (to indicate where the array is located on the slide) and placing the coverslip over the array before carefully and slowly pipetting the labelled-cDNA between the slide and the coverslip at either end. The sample draws itself underneath the coverslip by capillary movement covering the entire array area. This was done in the dark.

The slides were then kept in the dark and placed on a slow moving shaker for a couple of minutes to assist an even spread of the cDNA on the array slide.

Nucleic acid hybridization was then performed by incubating the slides overnight at 42°C in a Gentix hybridization chamber. To ensure that the slides were kept humid, 5ml water was added in the chamber.

The slides were washed after hybridization, for 5 min with 1X SSC buffer and 0.03% (w/v) SDS after removing the coverslips, and then were washed in 0.2X SSC for 3 min, and in 0.05X SSC for 3 min.

The slides were then dried by centrifuging them at 250 Xg for 4 min.

Section 2.2.5.4 Scanning of the array slides

The slides were scanned with a GenePix 4200A Axon scanner and a Genepix professional (6) and a Genepix professional (6) software program, with hardware settings of 635 (standard red) and 532 (standard green) for the wavelengths and laser power of 70% and filters of Cy5 and Cy3 for the high and low wavelengths respectively.

The slides were scanned at two wavelengths (532 nm for the Alexa Fluor® 555 dye and 635 nm for the Alexa Fluor® 647 dye).

The ratio was always on 635/532 with the GenePix array list (GAL) file. The images were saved as 3 multi-images, Tagged Image File Format TIFF (compressed).

Section 2.2.5.5 Fitting features of the scanned array slides

The images were opened for analysis with the GenePix Pro (6) program at a ratio formulation of 635/532. The blocks from the GenePix Array List (GAL) files were subsequently aligned on the images from the scanner. The relevant GAL file (A or B) was superimposed on the slide and each of the blocks and features aligned automatically. The features were then checked manually for empty or irrelevant features.

After fitting all the blocks and the features for each of the A and B slides, and the results from the analysis were saved as a GenePix results (GPR) file which could be opened in Excel.

Section 2.2.5.6 Normalization of the GPR files

The files were normalised with NORTT, which is a normalise program and Student's t- test gene expression data program 1.1.0, from the MRC Toxicology Unit.

The normalization was an intensity-dependent normalization with settings to treat the background fluorescence as zero and to pre-condense data by gene identity. The labelling method was chosen as Cy3 for the control (555) and Cy5 for the treated (674). Data that is flagged by GenePix as having failed to record a fluorescence in both channels was ignored. A normalised data file *.NOR was generated and saved. Data which had a single channel only, failed the threshold test, or failed to pass the circularity test in GenePix 5.0 was written to a *.NEG file.

Section 2.2.5.7 Analysis of the microarray data

For the comparison and the quality control of the microarray data, microarray statistical analysis was done with Microsoft Excel 2007, where the power of the normalised log₂ ratio of median was used.

To compare between the microarray data with linear regression, a VLOOKUP (VLOOKUP searches for a value in the first column of a table array and returns a value in the same row from another column in the table array) was used to identify the value of one array data corresponding to the value in the second array of data. The linear regression line was fitted with Graphpad prism 5. The microarray experimental data was stored and normalised in the Array Track and then the R software was used to analyse and identify the genes of interest.

High-Throughput GoMiner program (Zeeberg, 2005) was used to interpret the the microarray data.

Section 2.2.6 Transcriptome sequencing

Section 2.2.6.1 Sequencing

RNA sequencing was performed on RNA liver samples from F344 rats treated with ciprofibrate/corn oil for 3 hours. The University of Nottingham Next Generation AB SOLID 3 platform Sequencer was used, where it was broken up to ~20 Kbp with a sonicator, then each sequence was ligated with a labeled sequence of nucleotides, this generate a monoclonal sequencing unit (which is the sequence attached to beans on a 1:1 basis).

The monoclonal sequence units were then placed on slides, and then the sequencer read the information from the slides as short reads (which are readings of short parts of the whole DNA). These short reads were assembled with Ensembl software.

Section 2.2.6.2 Expression analysis

After generating the data it was analysed with CLC Genomics Workbench 3.7 software. To analyse differential expression, it was of importance to tell the workbench program how the samples were related. This was done by setting up an experiment to define the relationship between the samples, which was defined as a two group experiment (control and treated) n=4. The samples were specified as unpaired multi-group comparison with the number of groups as 2 groups.

Normalization with CLC Genomics Workbench program was essential in order to ensure that the samples were comparable and assumptions on the data for analysis were met, and to remove the bias effects of the sample preparation and array processing. This was done with a quantile normalization method before the statistical analysis was carried out.

Quality control was performed on the normalised data with CLC Genomics Workbench pro-

gram to enable visual inspection of the distribution and variability of the data, and to allow identification of any unwanted systematic differences between the samples.

The tools used were box plots/analyzing distributions, Hierarchical Clustering of samples and Principal Component Analysis. After the above steps the samples were ready for statistical analysis. Following statistical analysis the retrieval of the genomic information was processed using rat genome data RGD base tools, by entering the Ensembl number of the genes manually to recover the gene name and description.

Section 2.2.7 Quantitative real-time reverse transcriptase polymerase chain reaction (qRT-PCR)

Section 2.2.7.1 High Capacity RNA-to-cDNA

cDNAs were made from each of the 4 biological replicates obtained for each of the 3 treatments at the 4 timepoints. This was done using the high capacity RNA-to-cDNA kit (Applied Biosystems) according to manufacturers' instructions. As negative controls, the reaction was set up without a RT enzyme mix (-RT) to ensure no contamination from the buffer, and the whole mix without the RNA (-RNA) was used to control for genomic DNA contamination. The samples were prepared by using up to 2 µg of total RNA per 20 µl reaction as in Table 2.4.

Table 2.4 Preparation of the RT reaction mix.

Component	Volume/reaction (μ l)	Volume/reaction (μ l)
	+RT	-RT
2X RT buffer	10.0	10.0
20X RT enzyme mix	1.0	-
nuclease-free water	quantity sufficient to 20 μ l	quantity sufficient to 20 μ l
RNA sample	up to 2 μ l	up to 2 μ l
total per reaction	20	20

The tubes were mixed well and briefly centrifuged to spin down the contents and to eliminate air bubbles. Everything was on ice until used.

The samples were run with the program shown in Table 2.5 to initiate the reverse transcription reaction.

Table 2.5 Optimal conditions for use with high capacity RNA-to-cDNA kit.

	Temperature $^{\circ}$ C	Time (min)
Step 1	37	60
Step 2	95	5
Step 3	4	infinity

The cDNA was stored at -20° C until used.

Section 2.2.7.2 Standard curves for qRT-PCR products

Standard curves for each of the genes used (*CYP4A1*, *CYP3A1 G0s2*, *Ccnd1*, *AhR*, β -*actin* and *Scd1*) were created by using 5-fold serial dilutions of a control cDNA to demonstrate the qRT-PCR efficiency. This was done in triplicate.

The standard curves for each of the genes were generated with MX4000 software using a least mean squares curve fitting logarithm. The regression value r^2 was generated for each gene as ($Y = m \log X + b$) where m is the slope of the line.

The average amplification efficiency was determined as it is directly related to the slope of the curve throughout the cycling reaction. The amplification efficiency is $10^{-1/\text{slope}}$. This corresponds to the number of template molecules that are duplicated every cycle.

Section 2.2.7.3 Quantitative real-time PCR using TaqMan® gene expression

This was done with TaqMan® gene expression master mix real-time PCR, which supplies a fluorescence reading of messenger RNA mRNA expression throughout each cycle of the PCR procedure, which in turn uses a quantitative study of mRNA expression based on PCR cycling threshold (Ct) values using MX4000 software. The primers and probes were amplified, with one or more endogenous control gene in the same reaction. The endogenous control was chosen on the fact that the gene induction was not changed in the control and samples treated with ciprofibrate or CPA.

The qRT-PCR was used to study the changes in gene expression of *CYP4A1*, *CYP3A1 G0s2*, *Ccnd1*, and *Scd1* by using *AhR* and β -*actin* as normalization genes. *CYP4A1* and *CYP3A1* were used as positive controls for the ciprofibrate and CPA respectively. No template control (NTC), -RT and -RNA were run in parallel as negative controls, and each sample was done in two rep-

licates.

The best annealing temperature, the amounts of Forward, Reverse, primers and cDNA were all optimized with stepwise procedures (data not shown). The real-time PCR reaction was determined as in (Table 2.6) and amplified as in (Table 2.7) with a TaqMan thermal cycler MX400.

Table 2.6 Real-time PCR reaction. The amount of master mix is 18 μ l and then DECP water was added to the final amount of 45 μ l. This was ready to be spread in two 20 μ l well on a 96 well plate. The concentrations of the primers were 10 pmol/ μ l for the primers and 5 pmol/ μ l for the probes.

Gene	Forward primer (μ l)	Reverse primer (μ l)	Probe (μ l)
<i>AhR</i>	1.5	1.5	1.5
<i>β-actin</i>	1.5	1.5	1.5
<i>Ccnd1</i>	1.5	2.25	2.25
<i>CYP3A1</i>	2.25	2.25	2.25
<i>CYP4A1</i>	2.25	2.25	2.25
<i>G0s2</i>	2.25	2.25	2.25
<i>Scd1</i>	2.25	2.25	2.25

Table 2.7 Real-time PCR cycle program. The conditions of the real-time PCR reaction, the whole reaction was 90 min.

Number Of Cycles	Temperature ($^{\circ}$ C)	Time
1	95	10 min
40	95	20 sec
40	60	1 min

Section 2.2.7.4 Statistical analysis of qRT-PCR

Biogazelle qBasePluse software was used to normalise and analyse the gene expression with the relative quantity method. The magnitudes of RNA number of the genes targeted were determined from the experiments Ct values. The copy number of the samples was normalised to that of the endogenous controls *AhR* and *β-actin*.

Section 2.2.8 Quantity and quality measurement of RNA and cDNA

To quantitate the RNA, absorbance at 260nm was determined with a CECIL CE9500 Spectrophotometer, while a Nanodrop machine was used to measure the quantity of the cDNA and the amount of fluorescent labelling of the 555 and 647 dyes.

The concentrations of nucleic acid in the samples were determined according to 1 OD (optimal density) at 260nm = 20-33 ng/μl of single stranded DNA (for cDNA), and 1 OD at 260nm = 40 ng/μl of RNA.

Agarose gels [2% (1.0g/50ml)] were used to check the quality of the RNA. They were prepared by adding 1g agarose powder to 50 ml 1X tris boric acid EDTA (TBE) (10X TBE was prepared by adding 108g Tris base, 55g Boric acid, 9.3g Na₂EDTA to 1 litre diluted water) with 0.5ml 10 X SDS. Gels were post-stained with ethidium bromide.

Section 2.2.9 Measurement of serum Alanine Aminotransferase ALT

Measurement of serum Alanine Aminotransferase (ALT) activity was carried out using Vitros ALT slides (Ortho-Clinical Diagnostics). The serum ALT test was performed as a service, in the Clinical Chemistry Section, Pathology Department in the Queens Medical Centre (QMC), Nottingham.

Section 2.2.10 Statistical analysis

All data is represented as mean \pm standard deviation. Statistical significance was tested by Unpaired Student's t- test to compare between two data sets with two tailed distributions and a two sample equal variance. Dunnett's Multiple Comparison Test with a one way analysis of variance (ANOVA) was used for multiple comparisons. This was done with GraphPad prism 5.0 software (Inc, SanDiego, CA).

Statistical analysis for the RNA sequencing data was done with Student's t- test. Also the false discovery rate FDR was controlled (the false discovery rate is the proportion of false positives among the declared positive). The method used in CLC Genomics Workbench for controlling the FDR is that of Benjamini (Benjamini and Hochberg, 1995).

The inspection of the result of the statistical analysis was done with CLC Genomics Workbench software.

Chapter 3 Results

Section 3.1 Induction of hepatic DNA synthesis

Mouse liver was initially chosen, as it might provide a better model than rat liver, because of the availability of knockout mice. However, previous studies on strains of mice dosed with peroxisome proliferators (Styles et al., 1990) showed that the induction of hepatic DNA synthesis in mice starts at 24 hours, while others (Al Kholaiifi, 2008) and (Al Kholaiifi et al., 2008) found that it did not start until after 3-4 days after dosing. It was therefore necessary to investigate when exactly the hepatic induction happens in different strains of mice.

Section 3.1.1 Studies in mice

Section 3.1.1.1 Optimization of immunohistochemical technique

The aim was to optimize the protocol for immunohistochemistry, specifically the secondary antibody concentrations and the dewaxing time for the histological process.

A set of liver sections were stained by immunohistochemical protocol as described in Section 2.2.3.2 using the Amersham mouse anti-BrdU labelling system to detect BrdU incorporation into DNA. Two antibodies were used, the primary antibody was anti-BrdU mouse monoclonal antibody (1:1000) and different concentrations of secondary antibody (blotting grade goat anti-rabbit LgG horseradish peroxidase conjugate) from Bio-rad company. The concentrations were 1:50, 1:100 and 1:500. Preliminary experiments revealed that the Bio-rad antibody gave a stronger signal than Sigma and Amersham antibodies (data not shown).

As shown in Figure 3.1 the immunoreactivity observed in slide A (1:50) is very strong but the noise is also very high. The other hand slide C (1:500) has very low noise but BrdU is not de-

tected in the hepatocytes. Slide B (1:100) shows a good signal-to-noise ratio, and this concentration of secondary antibody was used for immunohistochemical staining for the rest of the experiments.

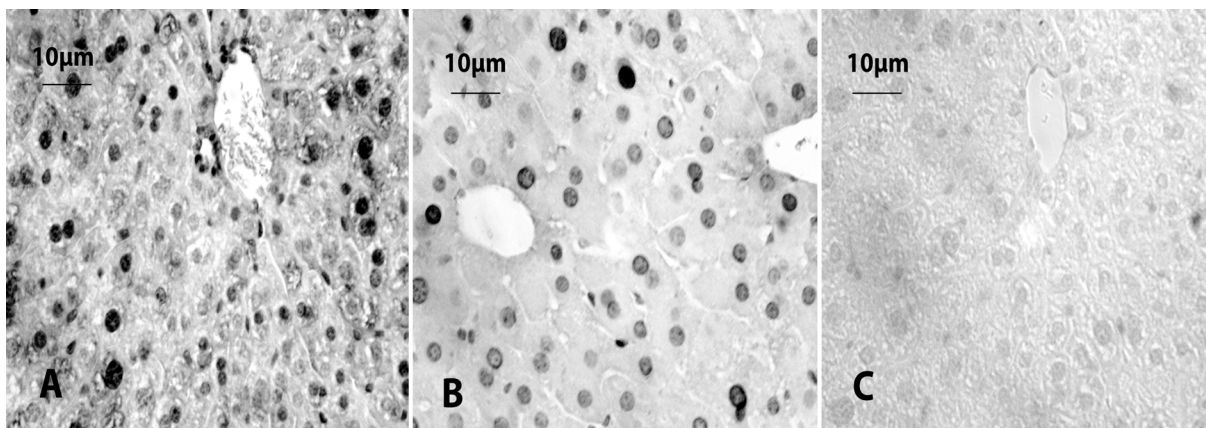


Figure 3.1 Optimization of secondary antibody concentration for immunohistochemistry. The figure shows slides stained with different concentrations of secondary antibody on 4.5µm thick liver sections from male 129S4/SvJae mice 9-10 weeks old. The animals were treated with 0.05 mg/kg BrdU with orange juice as their sole source of fluid, then gavaged with 100 mg/kg ciprofibrate for 4 days. The livers were then processed as in Section 2.2.3.1. The slides were stained according to immunohistochemical protocol using Amersham cell proliferation kit with different concentrations of secondary antibody from Bio-Rad. In slide A the concentration of the secondary antibody: buffer is 1:50, in slide B its 1:100 and in slide C 1:500. The slides are Harris haematoxylin counterstained. The photos were taken with a gray scale camera. The sections were examined under a light microscope at 400x. The scale bar=10µm.

The next part was to optimize the dewaxing time. The existing protocol required leaving the slides in xylene pots for 2 X15 minutes.

After preparing the slides (Section 2.2.3.1) they were left at 40°C for 2 minutes (or until the wax started to melt) then in xylene for 5 minutes. Liver and gut sections stained with this protocol had a high signal-to-noise ratio when compared with the 30 minute dewaxing time (Figure 3.2). Gut sections were used for all the experiments as a positive control (Figure 3.2 E and F). The intestinal epithelium consistently undergoes cell renewal, so it will show if the BrdU is taken up by the animal or not, and the labelled nuclei have a high signal to noise ratio especially in the intestinal crypts where the proliferation and differentiation occurs (Wille et al., 2004).

Similar optimization was undertaken to reduce the time needed for dehydration, hydration, and the time for primary antibody immunodetection of the incorporated BrdU (data not shown).

The overall time required for the immunohistochemical staining protocol was reduced from ~6 hours to ~3.5 hours. Thus, these data show that reducing the time for the immunohistochemical staining protocol does not necessarily decrease the quality of the slides. This revised procedure was used for the immunohistochemical staining for the rest of the experiments.

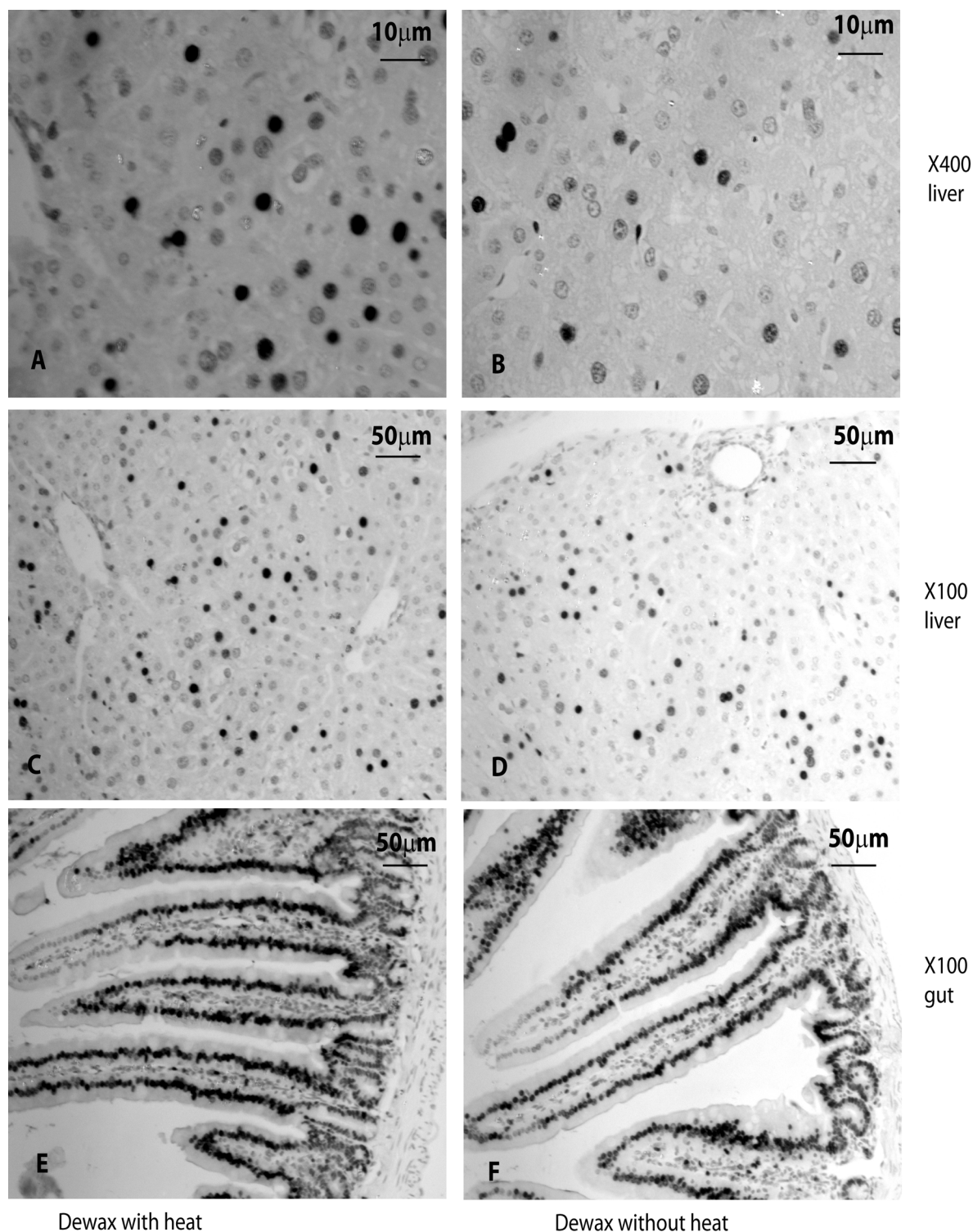


Figure 3.2 Optimizing of the dewaxing procedure. Liver and gut sections from male 129S4/SvJae mice. The animals were treated with BrdU for 5 days and gavaged with ciprofibrate for 4 days as shown in Figure 3.1. The slides were then stained with immunohistochemical protocol (Section 2.2.3.2). A,C (liver sections) and E (gut section) were dewaxed by putting the slides at 40°C for 2 minutes, then (while still hot) left in xylene for 5 minutes. B,D (liver sections) and F (gut section) were dewaxed by leaving in xylene for 2x15 minutes. Slides were counterstained with Harris haematoxylin. Sections were examined under a light microscope at 400X. Photos were taken with a gray scale camera. For A and B the scale bar=10µm for C,D,E and F the scale bar=50µm.

Section 3.1.1.2 Effect of ciprofibrate on hepatic DNA synthesis in mice

Previous work (Al Kholaiifi, 2008) in 129S4/SvJae mice showed that induction of hepatic DNA synthesis starts 3-4 days after dosing with peroxisome proliferators (methylclofenapate MCP), and this differs from work done by (Styles et al., 1990) on C57BL/6J mice who found that DNA synthesis starts as early as 24 hours after dosing with MCP.

This experiment was designed to study the time course of hepatic response to ciprofibrate in different strains of mice, to see whether the time course is distinct in different strains. Two inbred strains of mice were chosen C57BL/6JCrI which are resistant to liver carcinogenesis and DBA/2JCrI which are relatively susceptible to liver carcinogenesis (Diwan et al., 1986).

Section 3.1.1.2.1 Effect of BrdU in DBA/2JCrI mice

Before studying the effects of ciprofibrate on liver growth in DBA/2JCrI it was of importance to exclude any toxic effect from the BrdU -which is used as a DNA label- by studying the effect of BrdU on body weight (the effect on body weight acts as an indication for toxicity).

Male DBA/2JCrI mice were acclimatised for 7 days with 10% orange juice as their sole source of fluid, then dosed with 0.05% or 0.08% 5-Bromo-2'-deoxyuridine (BrdU) in 10% orange juice for 6 more days. The animals body weight was measured daily. The results established that, there was no significant difference between the weight of animals when beginning the experiment and at time of death (Student's un-paired t- test) (Figure 3.3).

From this result we can draw a conclusion that there was no gross toxic effect from the BrdU.

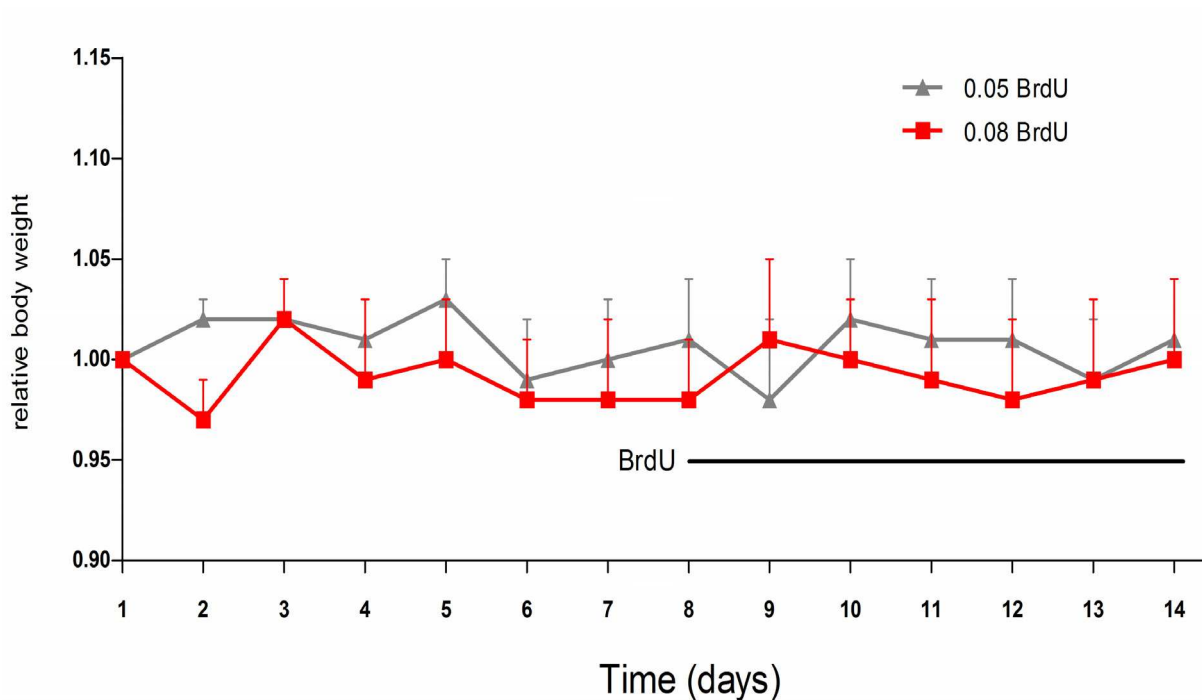


Figure 3.3 Effects of BrdU on body weight in DBA/2JCr1 mice. Groups of (matching weight) male DBA/2JCr1 mice aged 9-10 weeks old were acclimatised to 10% orange juice as their sole source of fluid, then were dosed with 0.08% (indicated by red squares) or 0.05% (indicated by gray triangles) BrdU in the orange juice for six more days (the beginning of the BrdU dosing is indicated by the horizontal black line). Mice body weights were measured on a daily basis for each individual mouse and relative body weight was calculated relative to the body weight for each individual mouse on day 0. BrdU administration started on day 8, see Section 2.2.1.2. Data are the mean \pm SD of 6 animals. With un-paired Student's t- test there was no significant difference between the weight of animals when starting BrdU and at the time of death. $P < 0.05$.

Section 3.1.1.2.2 Effect of ciprofibrate in DBA/2JCrI mice

The aim of this experiment was to test the effect of ciprofibrate on the induction of hepatic DNA synthesis in DBA/2JCrI mice.

Male DBA/2JCrI mice were acclimatized to 10% orange juice for 7 days then on day 8 they were given 0.08% BrdU with orange juice as their sole source of fluid. After one day of exposure to BrdU, 100mg kg⁻¹ day⁻¹ ciprofibrate or corn oil (which is used as a vehicle) was administered to the animals by gavage daily (see Section 2.2.1). Figure 3.4 shows the body weight relative to the body weight of each individual mouse on day 0.

By day 3 it became apparent that one of the mice -in a control group- was preventing the rest of the group from eating by aggressive behavior, so the average body weight of the group came down on that day. The mouse was isolated, and the body weight of the group rose.

The results showed that the animals' body weight was significantly larger at the end of the experiment compared with the beginning of the experiment, (un-paired Student's t- test, P<0.05).

This shows that the BrdU and the ciprofibrate have no toxic effect on the body weight of the DBA/2JCrI mice, and the growth of the animals was normal in comparison with the controls.

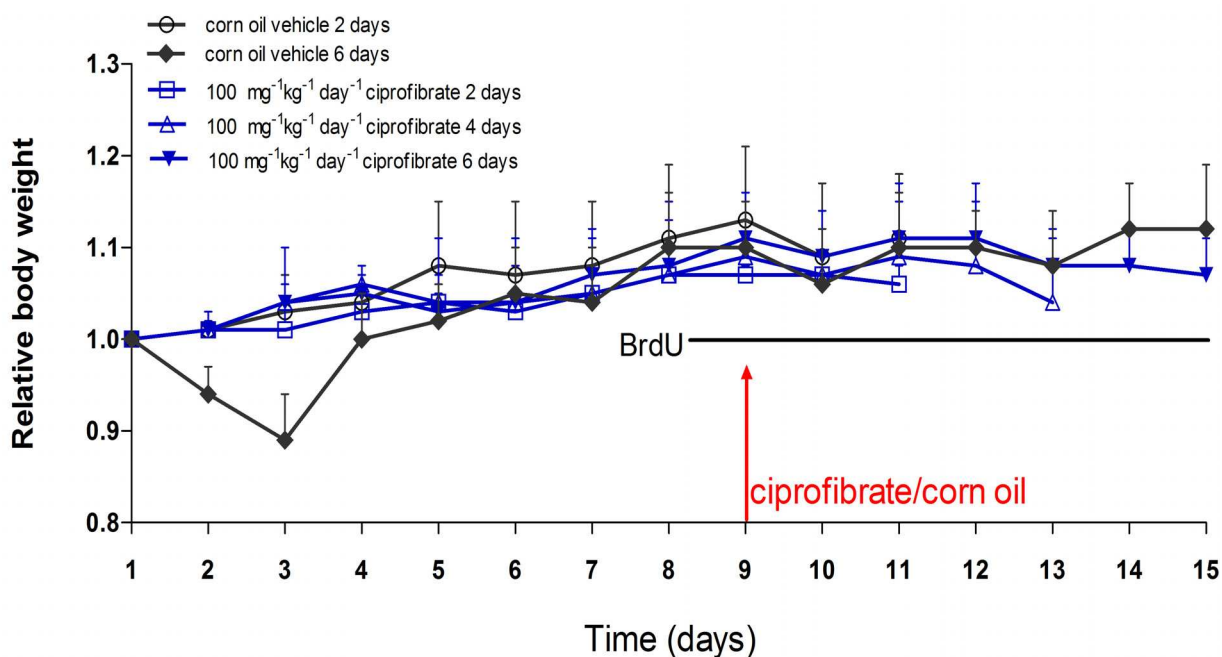


Figure 3.4 Effect of ciprofibrate on body weight in DBA/2JCrI mice. Groups of (matching weight) male DBA/2JCrI mice aged 9-10 weeks old were acclimatised to 10% orange juice then given 0.08% BrdU with orange juice as their sole source of fluid. After one day of exposure to BrdU the animals were gavaged with corn oil (vehicle) or 100mg kg⁻¹ day⁻¹ ciprofibrate (Section 2.2.1), and then killed after 2, 4 and 6 days of exposure to the ciprofibrate/corn oil. Mice body weights were measured on a daily basis and body weight was calculated relative to the body weight for each individual mouse on day 0. The BrdU administration started on day 8 and the red arrow indicates the beginning of administration of ciprofibrate/corn oil. Data are the mean \pm SD of 6 animals. Significant difference between the weight of animals when starting the experiment and at the time of death was done with unpaired t- test ($P < 0.05$).

The effect of ciprofibrate on liver growth of DBA/2JCrI mice is presented as relative liver weight and hepatic DNA synthesis (Figure 3.5).

Relative liver weight was significantly increased from 5.4 \pm 0.6% in the pooled control to 7.8 \pm 0.3% and 9.2 \pm 0.4% in the 4 and 6 days treated groups respectively, but there was no significant difference between the pooled control and the treated group for 2 days (6.2 \pm 0.4%) with Dunnett's multiple comparison test ($P < 0.05$) (Figure 3.5A).

The induction of hepatic DNA synthesis (the percentage of the BrdU-labelled hepatocytes to the total number of hepatocytes) was assessed as described in Section 2.2.4.1. The mice treated with

ciprofibrate for 2 days had no significant difference from the pooled control with Dunnett's multiple comparison test. On the other hand, animals dosed for 4 and 6 days with ciprofibrate had a significantly larger hepatic labelling index indicating more hepatic DNA synthesis occurs than the pooled control, as shown in Figure 3.5B.

These data show that ciprofibrate increases liver weight and hepatic DNA labelling index after 4 days of dosing with ciprofibrate, but does not significantly affect the liver weight or hepatic DNA synthesis after two days' dosing. This result is substantively the same as previous findings in 129S4/SvJae mice (Al Kholaiifi et al., 2008). In conclusion the ciprofibrate has no effect on body weight in DBA/2JCr1 mice treated with ciprofibrate up to 6 days. In liver growth studies, ciprofibrate did not show any significant increase on liver weight or on the induction of hepatic DNA synthesis after 2 days exposure to ciprofibrate but the statistically significant effect started after 4 days exposure.

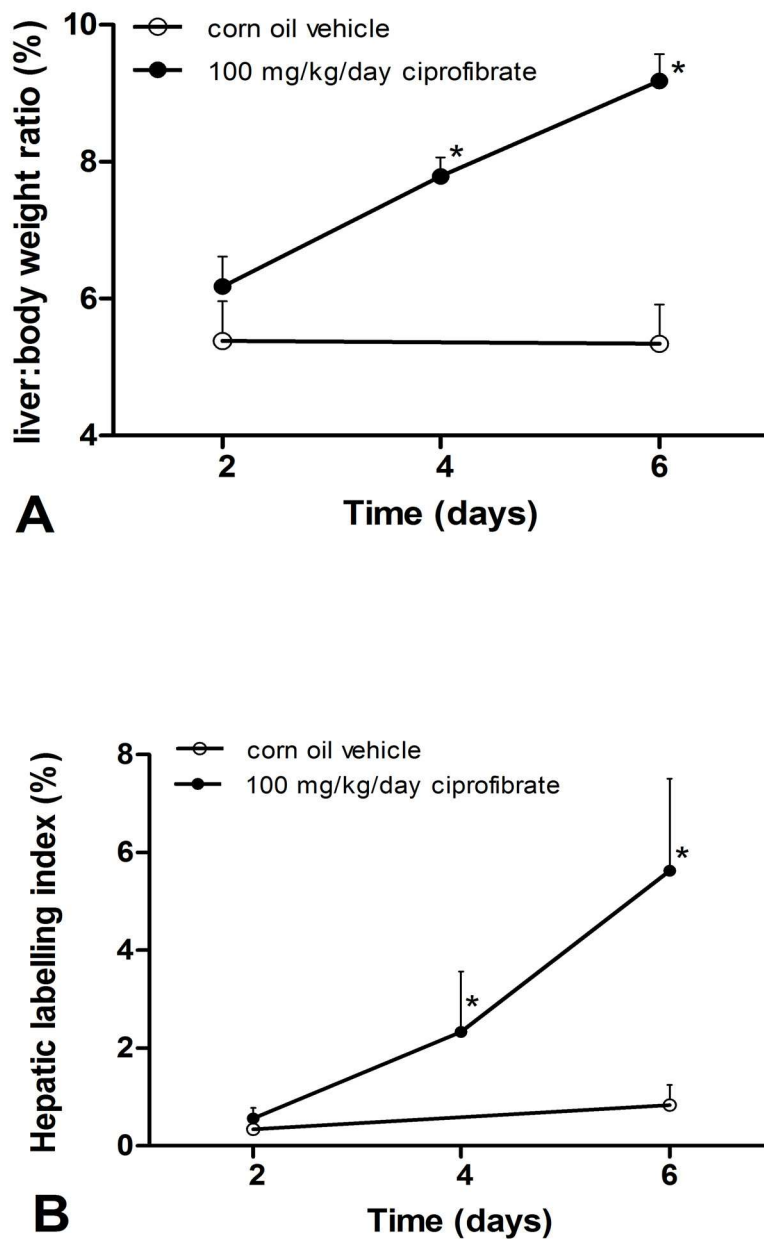


Figure 3.5 Effects of ciprofibrate on liver growth in DBA/2JCr1 mice. Groups of six DBA/2JCr1 male mice aged 9-10 weeks were dosed with 100mg kg⁻¹ day⁻¹ ciprofibrate or corn oil vehicle as described in Figure 3.4, and killed on days 2, 4 and 6. The left panel **A** shows liver to body weight ratio, the right panel **B** shows hepatic labelling index (the percentage of the labelled hepatocytes to the total number of hepatocytes). Comparison with Dunnett's multiple comparison test is against the pooled control group and statistically significant difference is indicated by an asterisk ($p < 0.05$).

Section 3.1.1.3 Effect of ciprofibrate in C57BL/6JCrI mice

Styles (Styles et al., 1990) has previously shown that methylclofenapate induced DNA synthesis at 24 hours after administration in the C57BL/J mouse strain. This experiment was designed to test the effect of ciprofibrate on the induction of hepatic DNA synthesis in C57BL/6JCrI mice by using immunohistochemical detection of BrdU.

Male C57BL/6JCrI mice were acclimatised to 10% orange juice for 7 days as their sole source of fluid, and then on day 8 they were given 0.08% BrdU with orange juice. After one day of exposure to BrdU the mice were gavaged daily with ciprofibrate or corn oil/vehicle (as a control) (Section 2.2.1). Figure 3.6 shows the time course effects of 0.08% BrdU and 100mg kg⁻¹ day⁻¹ ciprofibrate in male C57BL/6JCrI mice on body weight relative to body weight of each individual mouse on day 0. There was no significant difference between the animals body weight when starting the experiment and at time of death for all treated and control groups. Statistics was done with un-paired Student's t- test (P<0.05).

From these results it can be indicated that there was no toxic effect from BrdU or the ciprofibrate on C57BL/6JCrI male mice as measured by body weight loss.

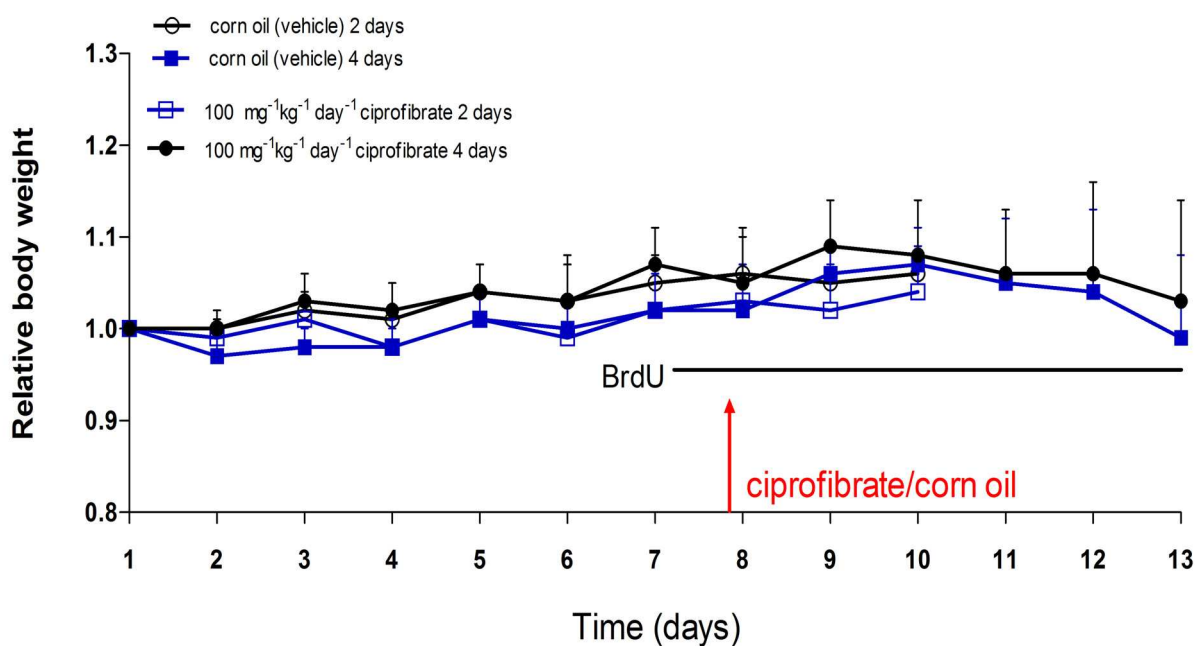


Figure 3.6 Effects of ciprofibrate on body weight in C57BL/6JCrI mice. Male C57BL/6JCrI mice aged 9-10 weeks were dosed with BrdU in the drinking orange juice and gavaged with corn oil (vehicle) or 100mg/kg/day ciprofibrate, then killed after 2 and 4 days (Section 2.2.1). Mice body weights were measured on a daily basis and relative body weight was calculated relative to the body weight for each individual mouse on day 0. The BrdU administration started on day 7 and the red arrow indicates the beginning of ciprofibrate/corn oil administration. Data are the mean \pm SD from 6 animals. There was no significant difference between the weight of the animals at the beginning of the experiment and at the time of death with students t- test ($P < 0.05$).

There was significant increases in the relative liver weights, from $4.8 \pm 0.7\%$ in the pooled control to $6.1 \pm 1.2\%$ and $8.3 \pm 0.4\%$ for the groups treated for 2 and 4 days respectively (Figure 3.7A). There was no significant difference in the hepatic labelling index between the pooled control and the group treated for 2 days with ciprofibrate, while the labelling index was significantly higher after 4 days of dosing with ciprofibrate, as shown in Figure 3.7B (statistics were done with Dunnett's multiple comparison test, $P < 0.05$).

From this we can establish that ciprofibrate has no toxic effect on body weight of C57BL/6JCrI male mice, while it increases liver weight significantly over control after 2 days' treatment. The hepatic labelling index did not show any difference after 2 days' exposure to ciprofibrate but was significantly different after 4 days' exposure.

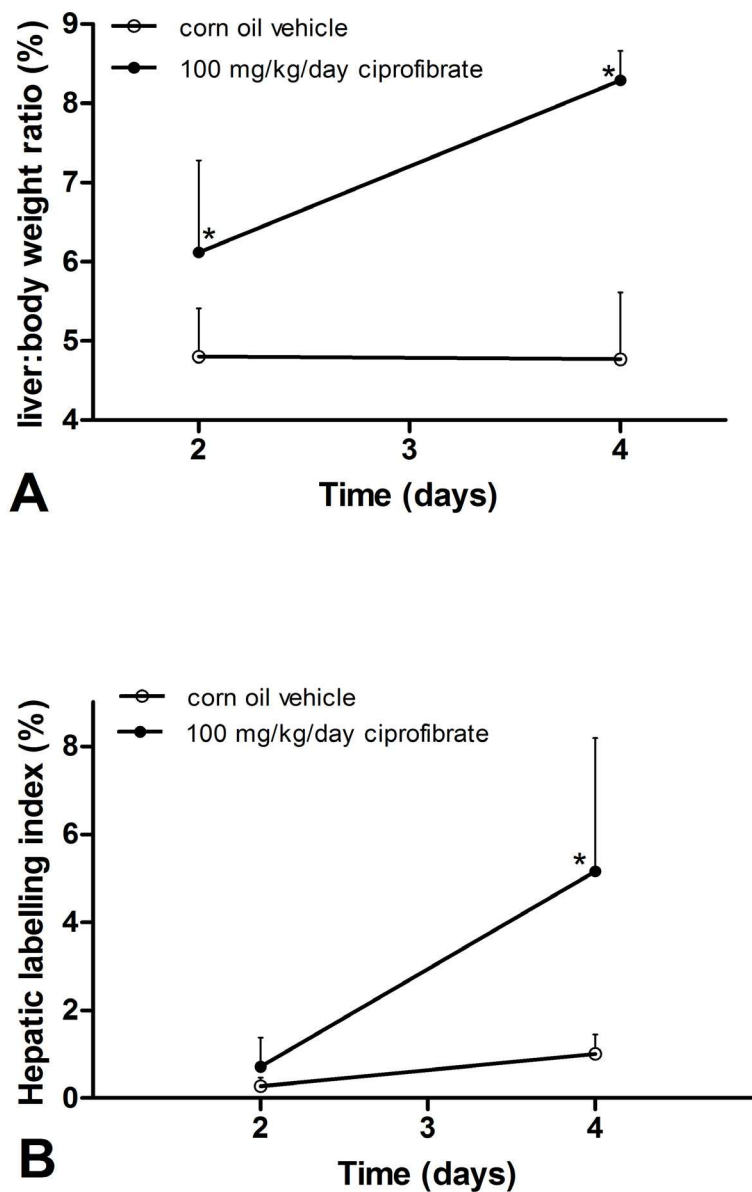


Figure 3.7 Effects of ciprofibrate on liver growth in C57BL/6Jcrl mice. Male C57BL/6Jcrl mice aged 9-10 weeks were dosed as described earlier in Figure 3.6 with 100mg/kg/day ciprofibrate or corn oil, then killed on days 2 and 4. In **A** Liver to body weight ratio was determined as described in Section 2.2.2. **B** shows hepatic labelling index (the percentage of the labelled hepatocytes to the total number of hepatocytes). Data shown are mean \pm SD (n=6). Statistically significant difference from the control group is indicated by an asterisk, with Dunnett's multiple comparison test. (P<0.05).

Section 3.1.2 Studies in F-344/NHsd rats

The demonstration that the induction of hepatic DNA synthesis with ciprofibrate in mice starts after 3-4 days makes it quite hard to determine when the induction is triggered (there is a window of ~2-3 days where the signal could have happened). Therefore, rat liver was used to determine if species difference would affect the timing of hepatic DNA synthesis.

Section 3.1.2.1 Effect of ciprofibrate on hepatic DNA synthesis in F-344/NHsd rats

Section 3.1.2.1.1 Time course of ciprofibrate effect in F-344/NHsd rats

To identify the genes responsible for the hepatic DNA synthesis, it was necessary to identify when the induction of hepatic DNA synthesis peaks in F-344/NHsd rats dosed with ciprofibrate.

Groups of male F-344/NHsd rats were dosed with 50mg kg⁻¹ day⁻¹ ciprofibrate or corn oil (vehicle) as a control, then killed after 16, 20, 24, 30, 36 and 48 hours. Two hours before killing they were dosed intraperitoneally (IP) with 100 mg kg⁻¹ BrdU. This experiment was done on two different occasions. On the first the time points were 24, 30, 36 and 48 hours, and on the second the time points were 16, 20, 24 and 30 hours. both with controls at the latest and earliest time points. The animals were from the same source, and were the same strain and age (Section 2.2.1).

Liver weight to body weight ratio is shown in Figure 3.8. Analysis with Dunnett's multiple comparison test showed a significant difference between the liver weight to body weight ratio for the pooled control 3.6±0.4% and treated groups at the time points 24 hours (4.0±0.2%), 30 hours (4.2±0.3%), 36 hours (4.1±0.3%), and 48 hours (4.8±0.2%) while for the groups killed at the time points 16 and 20 hours there was no significant difference (P< 0.05).

As a further test of whether the statistical significance found at the time points 24, 30, 36 and

48 hours is spurious the treated and control groups were compared against each other at each time point. It was found that the groups killed at the time points 24 and 30 hours had no significant difference when compared with the control groups at the same time point (24 hours treated against 24 hours control, 30 hours treated against 30 hours control). The group treated for 36 hours (there was no control at that time point) was compared against the control groups killed after 30 and 48 hours and there was no significant difference.

This indicates that the difference at the time points 24, 30 and 36 hours are not biologically significant and the significance that appeared when comparing the treated with the pooled control was spurious. On the other hand, the group killed after 48 hours had a significant difference from the control at the same time point (Student's t- test, $P < 0.05$).

From this result it is established that the liver weight to body weight ratio of F-344NH/sd rats treated with 50 mg kg^{-1} ciprofibrate is biologically and significantly high at 48 hours, while there is no biological difference for the other time points.

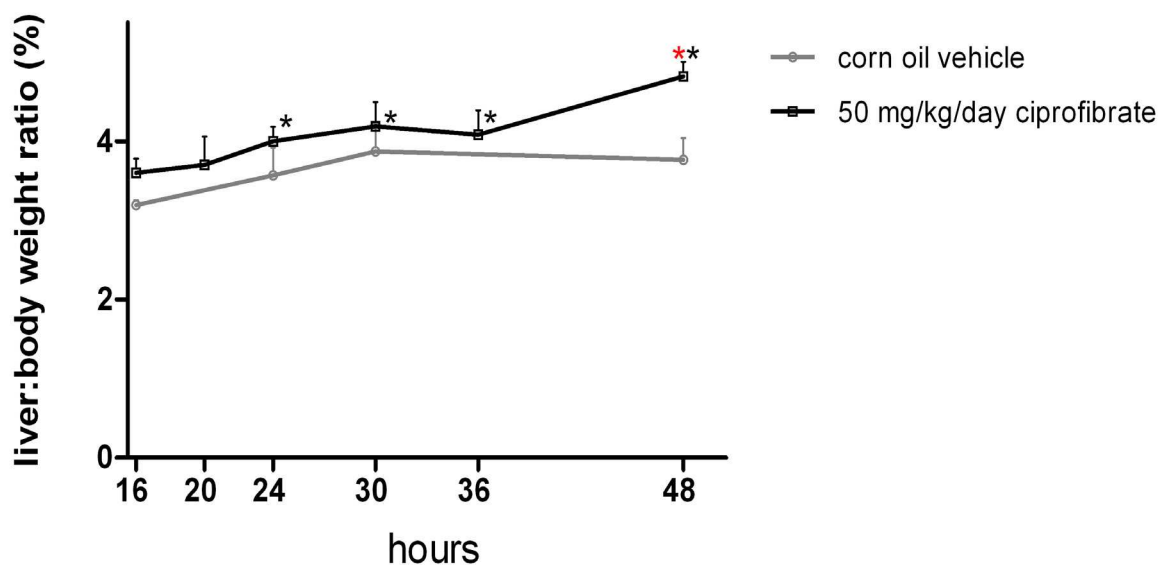


Figure 3.8 Time course of effects of ciprofibrate on liver weight in F-344/NHsd rats. Groups of male F-344/NHsd rats aged 14-15 weeks were dosed with 50mg/kg/day ciprofibrate or corn oil (vehicle), then killed at the time points indicated, BrdU labelling was intraperitoneally (IP) two hours before killing (Section 2.2). Each point represents the mean \pm SD of 6 animals, and is representative of results obtained from two experiments. Statistically significant difference from the pooled control groups is indicated by a black asterisk with Dunett's multiple comparison test ($P < 0.05$). Statistically significant difference from individual controls is indicated by a red asterisk with Student's t- test ($P < 0.05$).

Figure 3.9 shows the time course of the hepatic labelling index in male F-344/NHsd rats. The results showed a significant difference in the hepatic DNA synthesis between the pooled control groups ($0.3\pm 0.4\%$) and the group dosed with ciprofibrate at time 0 hour and killed after 24 hours ($3.8\pm 1.8\%$), 30 hours ($2.1\pm 1.1\%$) and 48 hours ($4.3\pm 1.4\%$), while the groups dosed for 16 hours ($0.2\pm 0.1\%$), 20 hours ($0.3\pm 0.2\%$) and the 30 hours ($2.1\pm 1.1\%$) had no significant difference from the pooled control group and fell to background levels. This gave two peaks; the first at 24 hours and the second at 48 hours.

From these results it can be established that for F-344NHsd rats treated with $50 \text{ mg kg}^{-1} \text{ day}^{-1}$ ciprofibrate the labelling index starts low then goes up at 24 hours and comes back down to background level, then up again at 48 hours. This could be as a response to the second dose that

is given after the first 24 hours and this would suggest the pattern that a peak develops after 24 hours from dosing. However, this needs to be investigated further.

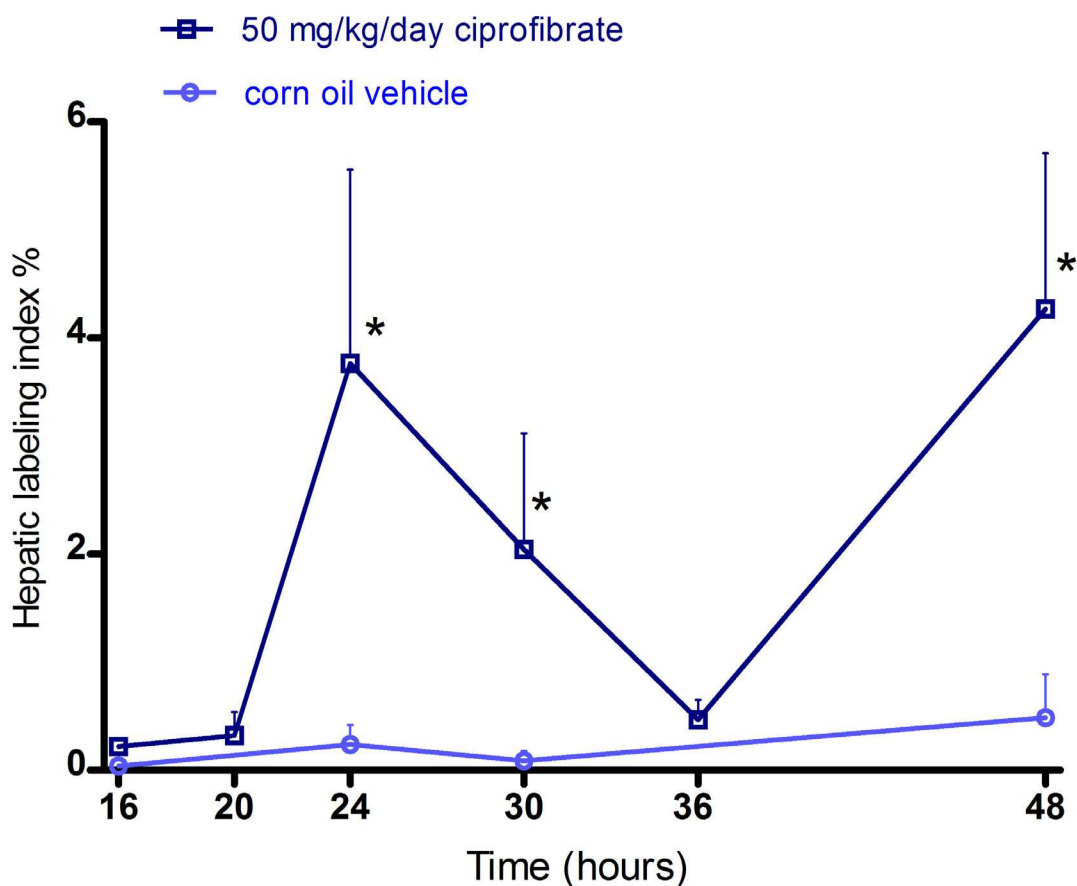


Figure 3.9 Time course of effects of ciprofibrate on hepatic DNA synthesis in F-344/NHsd rats. Groups of male rats were dosed with $50\text{mg kg}^{-1}\text{day}^{-1}$ ciprofibrate or corn oil (vehicle) as described in Figure 3.8. Immunohistochemical protocol and counting as in Section 2.2.3.2 and Section 2.2.4.1. All data are expressed as mean \pm SD of 6 animals, and are representative of results obtained from two independent experiments (in one experiment the time points were at 24, 30, 36 and 48 hour and in the other experiment the time points were 16, 20, 24 and 30 hours). Statistically significant difference from the pooled control group is indicated by an asterisk ($P < 0.05$) (Statistic done with Dunett's multiple comparison test).

Section 3.1.2.1.2 Dose response for ciprofibrate in rats

The objective of this experiment was to determine the optimal dose of ciprofibrate in F-344/NHsd rats that give the highest hepatic replicative DNA synthesis.

Al Kholaiifi (Al Kholaiifi, 2008) demonstrated that the induction of DNA synthesis in hepatocytes in F-344NHsd rats starts after 1 day of dosing with ciprofibrate, using a chronic BrdU labelling system where the hepatic DNA synthesis was acclimated in the liver. Also, Section 3.1.2.1.1 showed that the hepatic replicative DNA synthesis in F-344/NHsd rats is at the highest after 24 hours. Therefore the hepatic labelling index was determined after 24 hours.

Groups of six animals were gavaged with different concentrations of ciprofibrate dissolved in corn oil [0, 50, 100, 200, 300 mg kg⁻¹ body weight] and killed after 24 hours. Two hours before killing, the animals were given 100 mg kg⁻¹ BrdU intraperitoneally (IP) (Section 2.2). The liver weight ratios increased significantly at the doses 100 and 200 mg kg⁻¹ ciprofibrate to 4.1±0.3% and 4.6±0.5% in comparison with the control group 3.5±0.2%. The liver weight was not increased at the dose 300 mg kg⁻¹ to (3.9±0.08%) as illustrated in Figure 3.10A.

A similar pattern was found in the labelling index data. At 24 hours after dosing the labelling index for rats dosed with 50, 100 and 200 mg/kg body weight ciprofibrate were 6.3±2.6%, 6.0±3.0% and 5.9±1.0% respectively, in comparison to the control (vehicle) that was 0.2±0.2% and all three doses were significantly higher than the control with Dunnett's multiple comparison test. As for the group dosed with 300 mg kg⁻¹ ciprofibrate the induction went down to 0.76±0.64% (Figure 3.10B). This reduced response of the 300 mg kg⁻¹ implies that it is a result of a toxic reaction and is suppressing the liver growth.

These results establish that the doses 50, 100 and 200 mg kg⁻¹ body weight give a similar label-

ling index. The concentration 50mg kg^{-1} was selected for the assay as a optimal dose of ciprofibrate for F-344NHsd rats, because it was the smallest amount of compound and it gave a reasonably good labelling index.

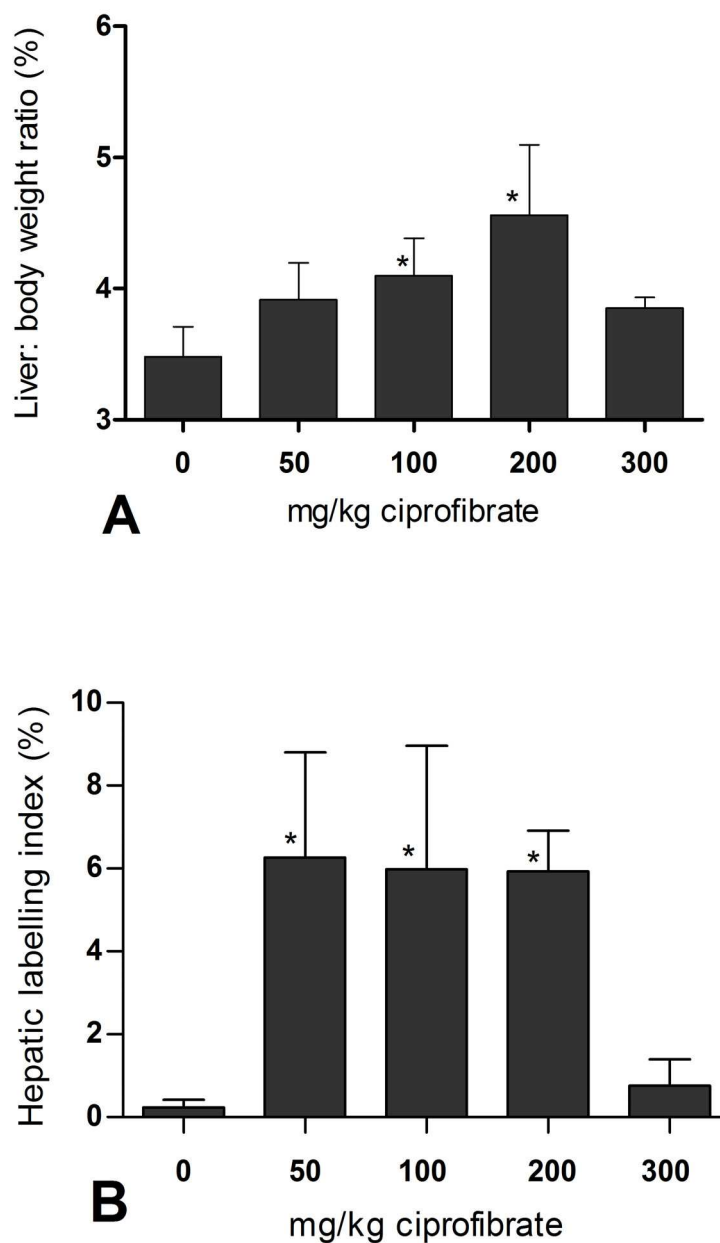


Figure 3.10 Dose response of ciprofibrate on liver growth in F-344/NHsd rats. Groups of (matching weight) male F344/NHsd rats aged 14-15 weeks were dosed at time 0 hour, with the indicated dose of vehicle or ciprofibrate, then at 22 hours they were dosed intraperitoneally with BrdU and killed at 24 hours (Section 2.2.1). **A** shows liver-to-body weight ratios. **B** shows labelling index (determined by immunocytochemical localization of BrdU see Section 2.2.4.1). All data are expressed as mean \pm SD of 6 animals, Statistically significant difference from the control group is indicated by an asterisk with Dunnett's multiple comparison test ($P < 0.05$).

Serum from the F-344HN/sd rats was assayed for alanine aminotransferase (ALT), as a test for hepatic damage (see Section 2.2.5.1). ALT level in the F-344HN/sd rat groups treated with ciprofibrate appeared higher at 100 and 200 mg kg⁻¹ than in the control group, but this was not a significant difference with Dunnett's multiple comparison test ($P < 0.05$) (Figure 3.11).

In conclusion the doses 50, 100, 200 and 300 mg kg⁻¹ ciprofibrate have no significant toxic effect on the liver tissue.

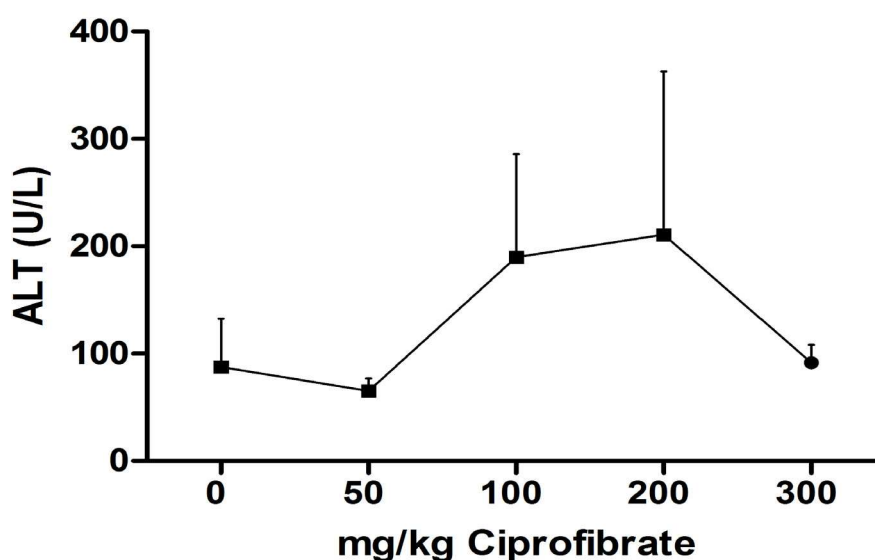


Figure 3.11 Effect of ciprofibrate on serum aminotransferase (ALT). Groups of (matching weight) male F-344NHsd rats aged 14-15 weeks were dosed with the indicated dose of vehicle or ciprofibrate, and killed after 24 hours as described earlier in Figure 3.10. Serum aminotransferase (ALT) activity was determined using Vitros ALT slides (Ortho-Clinical Diagnostics). The procedure was performed in the clinical chemistry department, Queens Medical Center (QMC) at Nottingham (see Section 2.2.5.1). Data are the mean \pm SD of 6 animals. Statistics done with Dunnett's multiple comparison test ($P < 0.05$).

Section 3.1.2.2 Effect of pregnane X receptor (PXR) ligands on hepatic DNA synthesis in F-344NHsd rats

It is known from the literature that the pregnane X receptor (PXR) ligands strongly enhance hepatic DNA synthesis of male and female Wistar rats (Schulte-Hermann et al., 1980). This experiment was designed to test the effects of the pregnane X receptor (PXR) ligands on liver growth and hepatic DNA synthesis of F-344/NHsd rats. The PXR ligands are used as a positive control, with a different mechanism for induction of hepatic DNA synthesis.

Section 3.1.2.2.1 PXR ligands in male and female rats: Pilot study

The aim of this experiment was to choose between cyproterone acetate CPA, and pregnenolone-16 α -carbonitrile PCN, based on their effects on hepatic DNA synthesis in male or female rats.

Two male and female F-344/NHsd rats were dosed with 100mg kg⁻¹ (the doses were chosen from the literature (Topinka et al., 2004b) and (Guzelian et al., 2006)) CPA or PCN or corn oil (vehicle), then killed after 24 hours. Two hours before killing they were dosed intraperitoneally with 100 mg kg⁻¹ BrdU. Liver weight to body weight ratio is shown in Figure 3.12A.

There was an increase in liver to body weight ratio for the male group treated with CPA 4.8% and 4.5% in comparison with the control at 3.9% and 3.7%, the liver weight ratio in male rats treated with PCN is 3.9% and 4.0% and for the females treated with CPA it was 3.9% and 3.6%. The labelling index for the male group treated with CPA was 6.7% and 7.6% and for the group treated with PCN was 6.6% and 6.3% in comparison with the control group 0.5% and 0.3% but the female group was extremely high at 49.2% and 53.3% (Figure 3.12B).

From these results it was decided to use CPA on males as the CPA gave the highest liver to body weight ratio and a high labelling index for the males (in comparison with male F-344/NHsd rats

treated with ciprofibrate the value was $3.8 \pm 1.8\%$ as shown in Figure 3.9), and the CPA was easier to dissolve in the corn oil than the PCN, which could only be used as a suspension. Although the female group gave a very high labelling index it was not a properly controlled experiment since there was no female control group, and so there were two variables (dose and sex) when compared with male controls. It would be interesting to investigate the effects of CPA on liver growth in female rats.

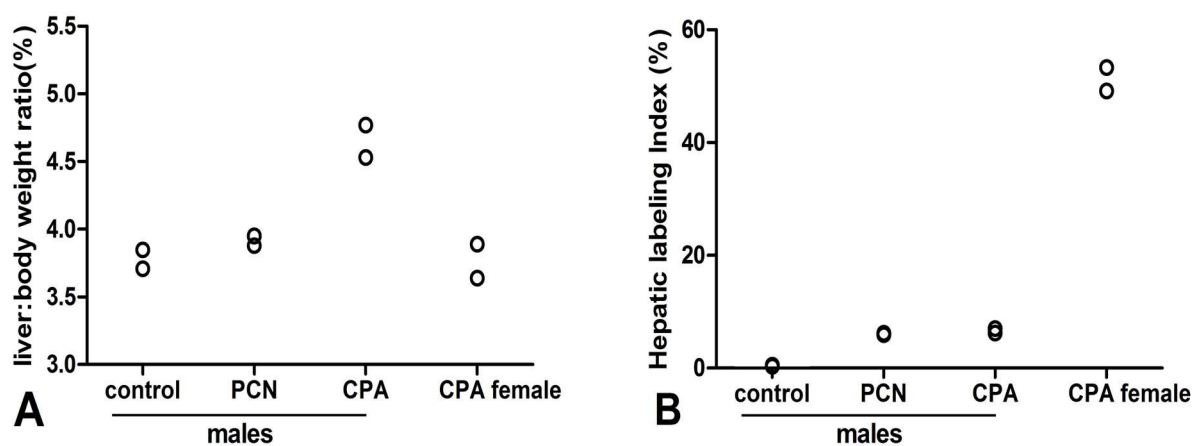


Figure 3.12 Effects of PXR ligands in male and female rats: pilot study . Three groups of male F-344/NHsd rats aged 14-15 weeks were treated with 100mg kg^{-1} PCN, CPA or corn oil and a group of females were treated with 100 mg kg^{-1} CPA. They were then killed after 24 hours. Two hours before killing they were dosed intraperitoneally with BrdU (Section 2.2). A shows liver weight to body weight ratio. B shows the hepatic DNA synthesis (as determined by immunocytochemical localization of BrdU) ($n=2$).

Section 3.1.2.2.2 Effects of CPA in male F-344/NHsd rats

The aim of this experiment was to investigate more rigorously the effects of CPA on liver growth and DNA synthesis in male F-344NHsd rats.

Rats were dosed with 100 mg kg^{-1} CPA or corn oil (vehicle), then killed after 24 hours. Two hours before killing, they were dosed intraperitoneally with BrdU. The liver was then processed and stained by immunohistochemical protocol (Section 2.2.3). Liver weight to body weight ratio is shown in Figure 3.13A.

Analysis with Student's t- test showed no significant difference between liver weight to body weight ratio for the control ($3.8 \pm 0.5\%$) and treated group ($4.2 \pm 0.3\%$). There was a significant difference in the labelling index between the control group ($0.1 \pm 0.1\%$) and the group dosed with CPA ($7.8 \pm 2.9\%$) Figure 3.13B.

This result revealed that CPA in male F-344NHsd rats had no significant effect on liver weight to body weight ratio after 24 hours, but has a significant effect on the labelling index.

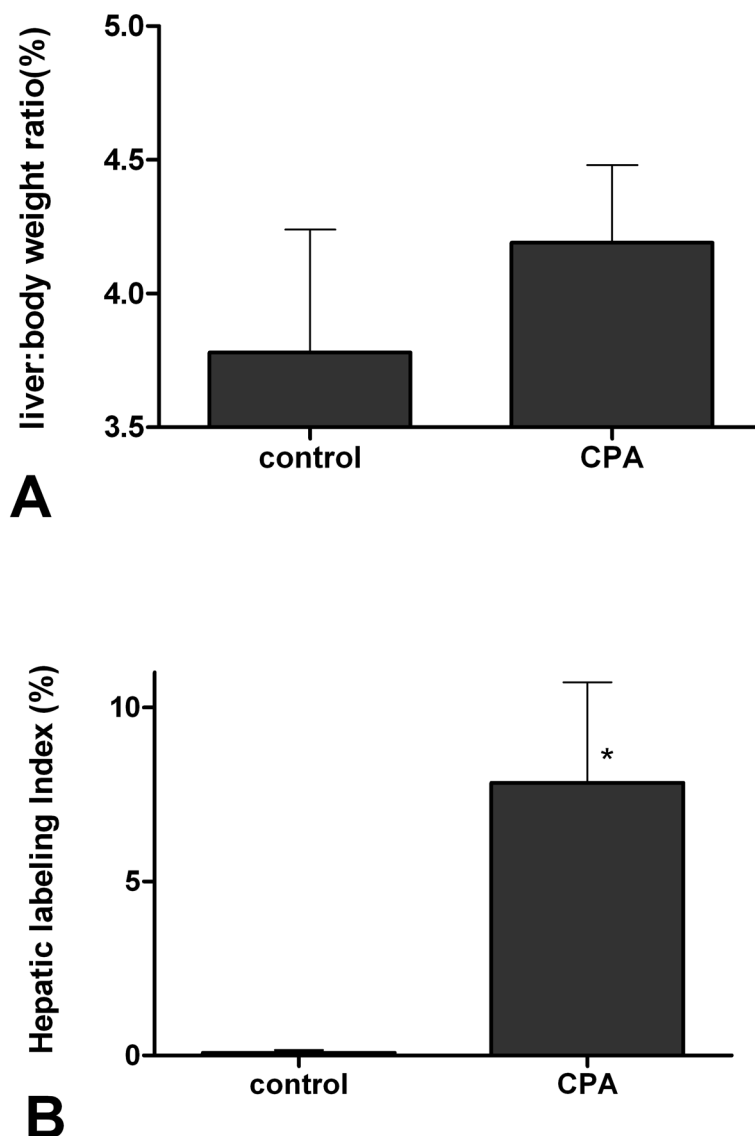


Figure 3.13 Effects of CPA on liver growth in F-344NHsd rats. Male F-344NHsd rats aged 14-15 weeks, were dosed with 100 mg kg⁻¹ CPA or corn oil (vehicle), and killed after 24 hours (Section 2.2.1). A shows liver weight to body weight ratio. B shows the immunocytochemical localization of BrdU (as in Section 2.2.4.1) for the control and CPA treated group. All data are expressed as mean±SD of 6 animals, Statistically significant difference from the control group is indicated by an asterisk, Statistics done was with Student's t- test (P<0.05).

Section 3.2 Zonal distribution of DNA synthesis in liver

The experiments below were designed to establish if the replicating hepatocytes are found mostly in the periportal or centrilobular regions for F-344NHsd rats and different strains of mice dosed with different treatments. Table 3.1 shows the strains of mouse or rat, which had been studied dosed with different treatments. It should be noted that some of the slides used were taken from (Al Kholaiifi, 2008). Al Kholaiifi dosed the animals and stained the slides with immunohistochemical technique. I then read the slides and determined the zonation of the BrdU-labelled hepatic nuclei in the periportal and perivenous zones as described in Section 2.2.4.2:

Table 3.1 Zonal distribution study.

No	Receptor	Treatment	Species/strain	Reference
1	PPAR α	ciprofibrate	F-344NHsd rats	Section 3.1.2.1
2	PXR	CPA	F-344NHsd rats	Section 3.1.2.2.2
3	CAR	TCPOBOP	129S4/SvJae mice	(Al Kholaiifi, 2008)
4	PPAR α	ciprofibrate	129S4/SvJae mice	(Al Kholaiifi, 2008)
5	PPAR α	ciprofibrate	129S4/SvJae PPAR α -null mice	(Al Kholaiifi, 2008)
6	PPAR α	ciprofibrate	C57BL/6JCrI mice	Section 3.1.1.3
7	PPAR α	ciprofibrate	DBA/2JCrI mice	Section 3.1.1.2.2
8	PPAR α	MCP	129S4/SvJae mice	(Al Kholaiifi, 2008)
9	PPAR α	MCP	AP mice	(Al Kholaiifi, 2008)

Section 3.2.1 Zonation studies in F-344/NHsd rats

Section 3.2.1.1 Induction of hepatic DNA synthesis by PPAR α ligands (ciprofibrate)

This experiment was designed to determine the lobular distribution of hepatocellular proliferation in liver of F-344/NHsd rats treated with ciprofibrate. Figure 3.14 shows immunohistochemical localization of hepatic DNA synthesis in F-344/NHsd rats treated with 50mg kg⁻¹ ciprofibrate for 4 days (Section 2.2.4.1).

Figure 3.14A and B show central veins (CV) on the right hand side. Hepatocytes closest to the (CV) are not BrdU-stained. In Figure 3.14C a portal space (PS) is on the right hand side and the stained hepatic nuclei are mostly gathered around the (PS). Closer examination of immunohistochemical-stained slides on higher power (X400) show a clear distribution of labelled nuclei Figure 3.15. The hepatocytes around the central vein show little labelling as shown in Figure 3.15A and B. While several labelled cells gather around the portal space (Figure 3.15C and D) upon visual inspection.

From this it established that the zonal distribution of hepatic replication in male F-344/NHsd rats dosed with ciprofibrate is not randomly distributed and was more situated at the periportal zone than at the central veins, and this is consistent with the results established from (Barrass et al., 1993) on Sprague-Dawley rats treated with MCP and clofibric acid.

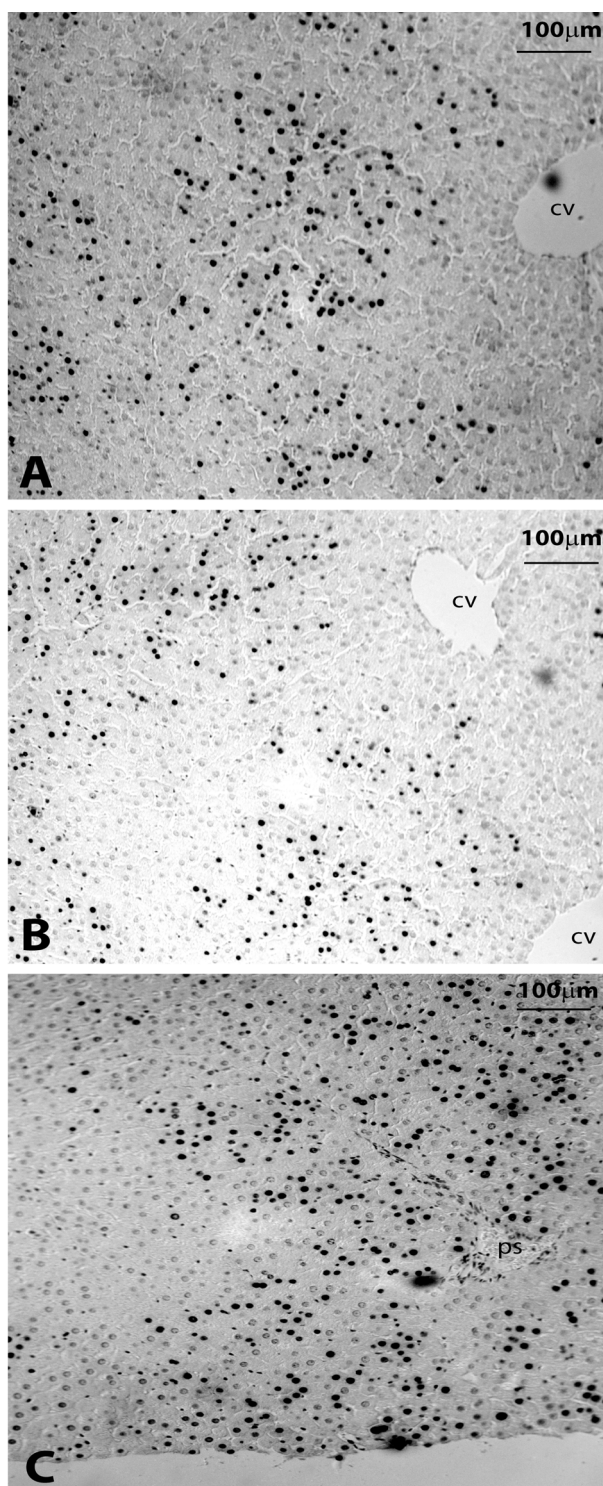


Figure 3.14 Zonal distribution of hepatic replication in F-344/NHsd rats. Immunohistochemical sections of livers from male Fisher 344/NHsd rats aged 14-15 weeks and treated with BrdU in orange juice as described in Section 2.2.1.2. Then gavaged with $50 \text{ mg kg}^{-1} \text{ day}^{-1}$ ciprofibrate for 4 days. A and B show central veins (CV); hepatocytes in the area closest to the CV are not BrdU stained. C shows a portal space (PS) on the right hand side, and the hepatocytes are mostly stained in the area around the PS. Sections were examined under a light microscope at X40. Photos were taken with a gray scale camera. The scale bar=100µm. Slides were from (Al Kholaiifi, 2008) (personal communication).

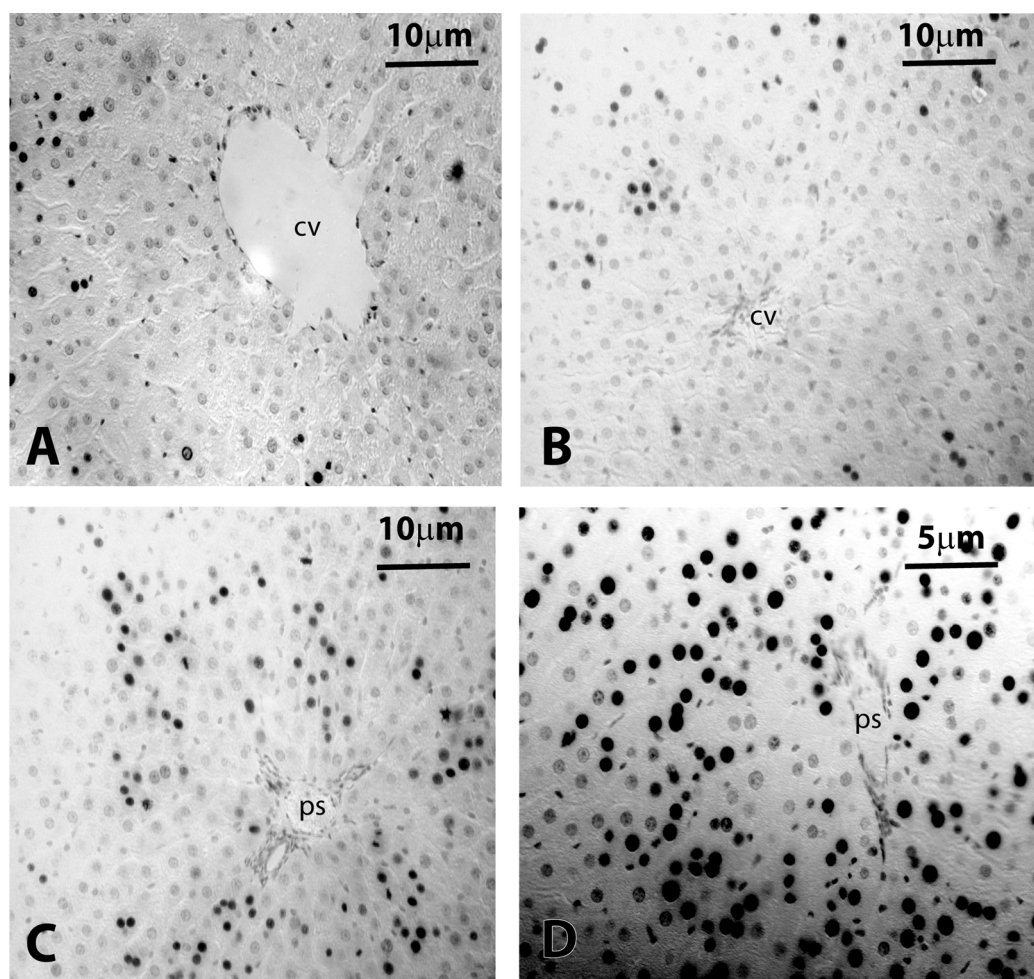


Figure 3.15 Central vein and portal space of liver sections in F-344/NHsd rats. Liver sections in male Fisher 344/NHsd rats aged 14-15 weeks and treated with BrdU. Then gavaged with 50 mg kg⁻¹ day⁻¹ ciprofibrate for 4 days Section 2.2.1. A and B panels show central veins CV and the area around it. C and D show portal spaces PS and the area around it. Sections were examined under a light microscope at X400. Photos were taken with a gray scale camera. In A, B and C the scale bar=10µm. In D the scale bar=5µm. Slides were a kind gift from (Al Kholaiifi et al., 2008) (personal communication).

The distribution of labelled hepatocytes was determined in periportal and centrilobular zones, as described in Section 2.2.4.2. There was a significant difference between the incidence of labelled hepatocytes between centrilobular and periportal regions in the control (Figure 3.16A) and treated animals according to Student's t- test ($p < 0.05$) (Figure 3.16B), with the number of labelled cells being larger in the periportal region by 4, 12 and 20 fold in the rats treated with ciprofibrate for 1, 2 and 4 days.

These results show the hepatic zonal distribution of F-344/NHsd rats liver is periportal rather than centilobular, and that treatment with ciprofibrate emphasises this result.

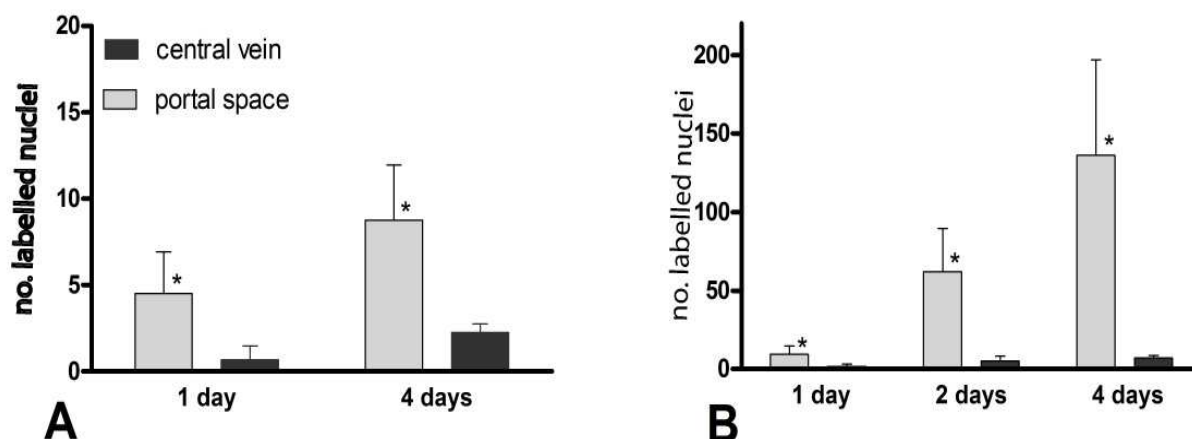


Figure 3.16 Hepatic zonal distribution of cell proliferation in rats treated with ciprofibrate.

Male F-344/NHsd rats aged 14-15 weeks and treated with BrdU (Section 2.2.1). Visualization of BrdU-stained hepatocyte nuclei was undertaken using the method of (Barrass et al., 1993) described in Section 2.2.4.2. The number of cells labelled in periportal (PS- gray bars) or centrilobular (CV- black bars) is shown in **A** for vehicle control rats orally treated with corn oil for 1 and 4 days (the number of animals is 6 & 4 respectively) and in **B** animals treated orally with 50 mg kg⁻¹ day⁻¹ ciprofibrate for 1, 2 and 4 days (the number of animals is 6, 5 and 5 respectively). The bars show the mean value, and the error bars depict one standard deviation. Statistically significant difference from the control group is indicated by an asterisk. Statistics done was with Student's t- test (P<0.05).

Section 3.2.1.2 Induction of DNA synthesis by the PXR ligand cyproterone acetate (CPA)

The aim of this experiment was to determine lobular distribution of hepatocellular proliferation in liver of F-344/NHsd rats treated with cyproterone acetate (CPA).

Male F-344/NHsd rats were treated with 100 mg kg⁻¹ CPA as described in Section 3.1.2.2.2. Immunohistochemical staining was applied on the liver tissue and the distribution of labelled hepatocytes was determined in periportal and centrilobular zones, as described in Section 2.2.4.2.

There was a significant difference in the number of labelled hepatocytes between centrilobular and periportal regions as in Figure 3.17 with the number of labelled cells being larger in the periportal region, comparable to the results that were achieved from the previous experiments on F-344/NHsd rats treated with corn oil or ciprofibrate (the statistic used was Student's t- test, $P < 0.05$, the number of animals is 6).

From these results we can establish that the hepatic zonal distribution of F-344/NHsd rats liver has a periportal distribution when treated with CPA, and this is consistent with the result found by (Barrass et al., 1993) and the results found earlier for ciprofibrate (Section 3.2.1.1).

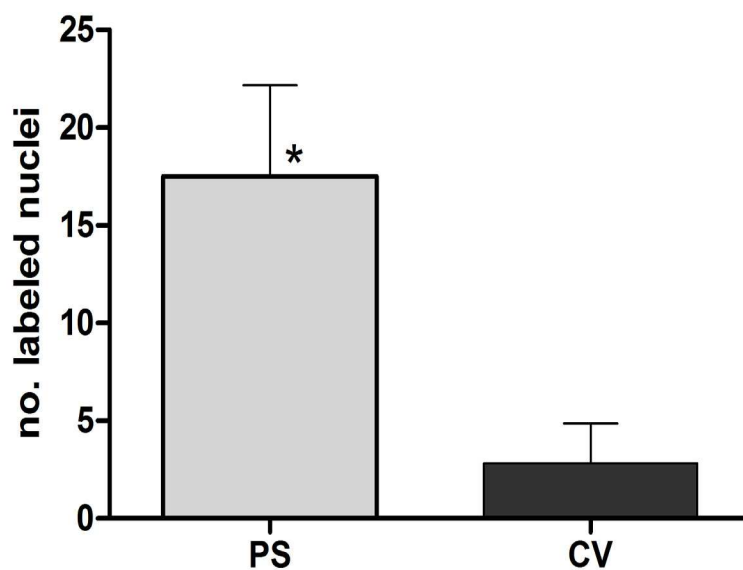


Figure 3.17 Hepatic zonal distribution of cell proliferation in rats treated with CPA. Male F-344NHsd rats aged 14-15 weeks old were dosed with 100 mg kg^{-1} CPA and killed after 24 hours. Two hours before killing they were dosed intraperitoneally with BrdU. Immunohistochemical protocol and visualization of BrdU-stained hepatocyte nuclei was undertaken as described in Section 2.2.4.2. The figure shows the number of cells labelled in the periportal region (PS- gray bar) or in the centrilobular region (CV- black bar) (the number of animals is 6). Statistically significant difference is indicated by an asterisk with Student's t- test between the PS and the CV ($P < 0.05$).

Section 3.2.2 Zonation studies in mouse

These experiments were designed to establish if the replicating hepatocytes are found mostly in the periportal or centrilobular regions for different strains of mice dosed with different hepatic inducing chemicals.

Section 3.2.2.1 Induction of hepatic DNA synthesis by the constitutive androstane receptor (CAR) agonist TCPOBOP in 129S4/SvJae mice

This experiment was designed to determine the lobular distribution of hepatocellular proliferation in male and female 129S4/SvJae wild type mice, treated with a CAR agonist TCPOBOP, as a hepatocyte mitogen, to determine if this caused differential lobular zonation of hepatocyte labelling. 129S4/SvJae mice were dosed with BrdU added to the orange juice as their sole source of fluid. After at least one day on BrdU, the animals were dosed by gavage with 3 mg kg⁻¹ TCPOBOP and killed after 2 days. Immunohistochemical staining was done on the liver tissue as in Section 2.2.3.2. Visualization of BrdU-stained hepatocyte nuclei was undertaken, essentially as described in Section 2.2.4.1.

Figure 3.18 shows immunohistochemical localization of hepatic DNA synthesis in liver sections. Figure 3.18A and B show central veins on the right hand side and portal spaces on the left. Figure 3.18C shows a CV in the middle top of the liver section. All the slides when examined on low power did not show any zonation for the BrdU-stained hepatocytes and the labelled hepatocytes were evenly distributed in the liver lobule. Examination of mice slides on higher power (X400) corroborated a homogeneous distribution of labelled nuclei. The BrdU-stained hepatocytes around the central veins and the portal spaces do not show any preferential lobular distribution as shown in Figure 3.19A, B, C and D. From this it established that the zonal distribution of hepatic replication in male 129S4/SvJae mice has no discernible difference between the periportal and the perivenous zones.

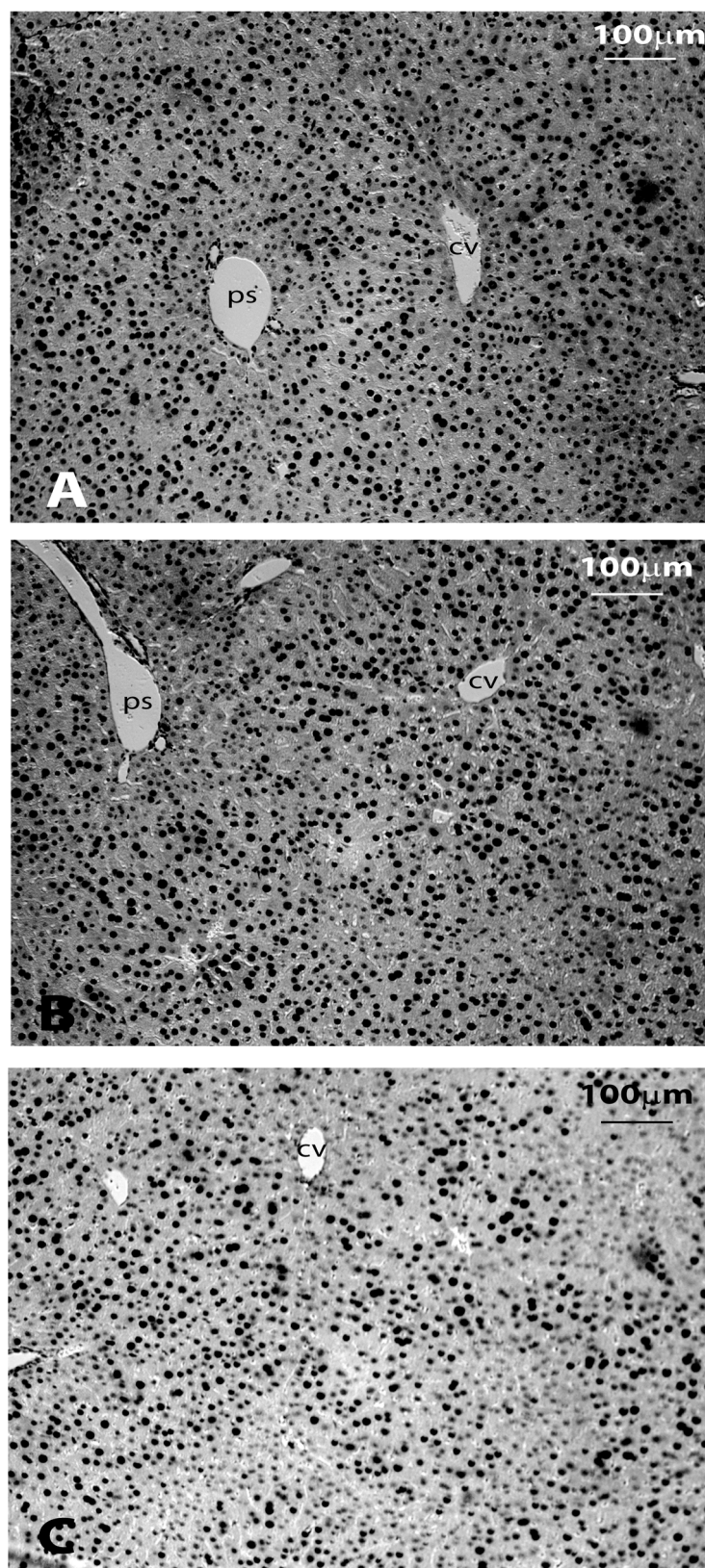


Figure 3.18 Zonal distribution of hepatic replication in 129S4/SvJae mice. Male 129S4/SvJae mice aged 9-10 weeks were dosed with BrdU then gavaged with 3 mg kg^{-1} TCPOBOP and killed after 2 days. Immunohistochemical protocol and visualization of BrdU-stained hepatocyte nuclei was undertaken on liver sections as described in Section 2.2.3.2 and Section 2.2.4.2. A- C sections were examined under a light microscope at 40X. Photos were taken with a gray scale camera. The scale bar=100µm. Slides were a gift from (Al Kholaiifi et al., 2008).

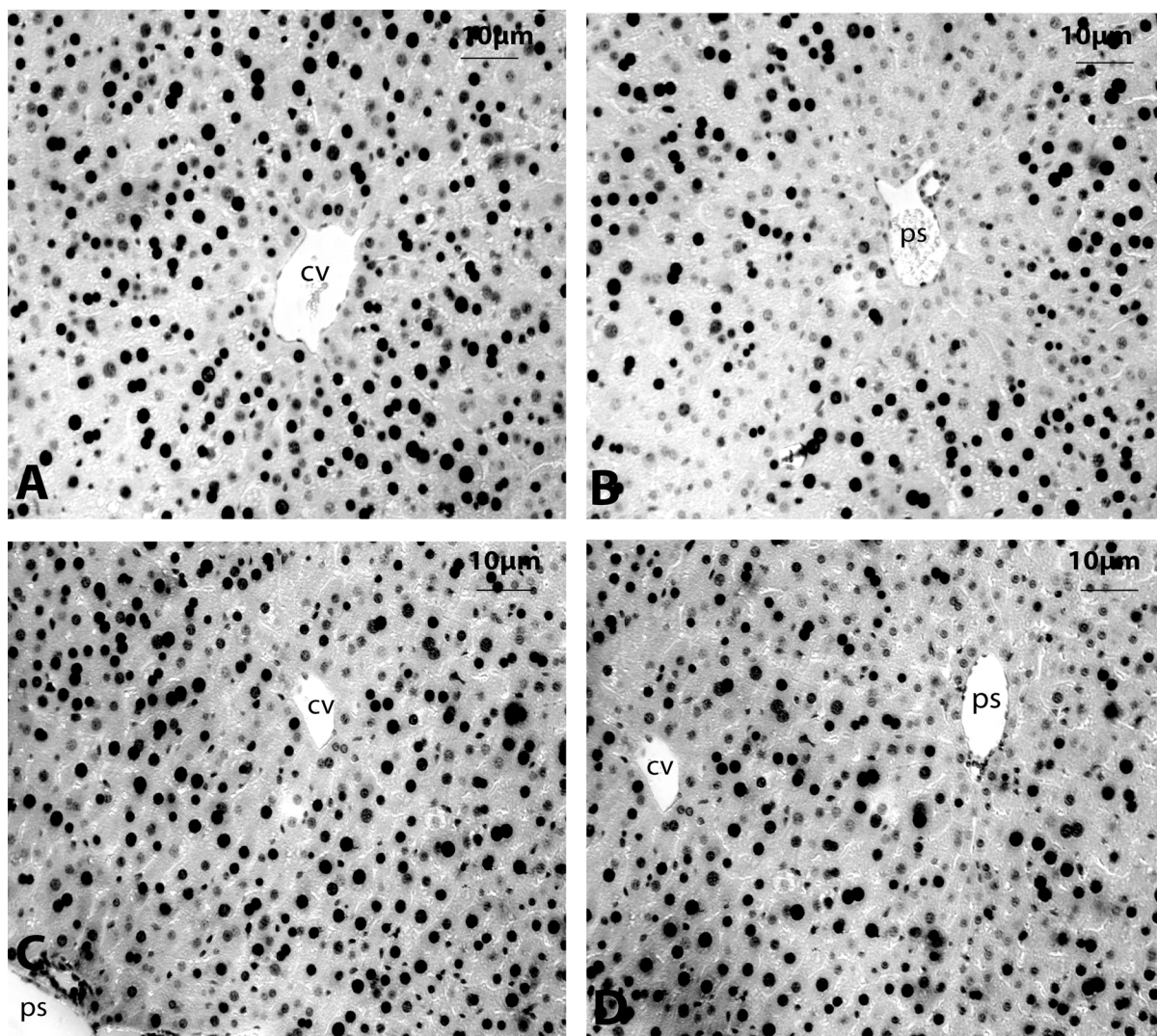


Figure 3.19 Central veins and portal spaces of liver sections in 129S4/SvJae mouse. Immunohistochemical staining of liver from 129S4/SvJae male mice, the animals were dosed with BrdU and TCPOBOP as in Figure 3.18, slides were a kind gift from (Al Kholaiifi et al., 2008). Visualization of BrdU-stained hepatocyte nuclei was undertaken as in Section 2.2.4.2. The scale bar represents 10µm. Sections A-D were examined under a light microscope at 400X. Photos were taken with a gray scale camera.

There was no significant difference in the hepatic zonation between the periportal and the perivenous regions according to Student's t- test ($P>0.05$) for male groups (control and treated) and the control female group, as shown in Figure 3.20. However, for the female group treated with TCPOBOP there was a significant zonation ($P=0.005$).

In this case, the labelled cells were preferentially perivenous (CV), although the magnitude of the effect was relatively small, with the number of labelled cells in the perivenous region being less than two times greater than those in the periportal region. The total labelling index was strongly increased by TCPOBOP.

From these results it was found there was no significant difference in the number of labelled hepatocytes between the periportal and the perivenous zones in male and female 129S4/SvJae control mice and the male mice treated with TCPOBOP, but there was a significant effect for treated female mice where the labelled cells were preferentially perivenous.

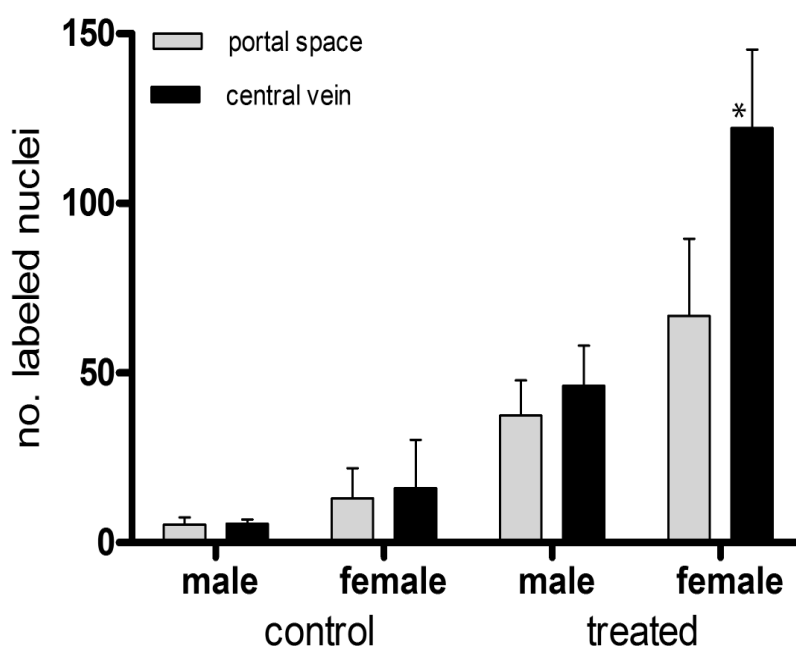


Figure 3.20 Zonal distribution of hepatic DNA synthesis in 129S4/SvJae mice treated with TCPOBOP. Male and female 129S4/SvJae mice were treated with BrdU then gavaged with a single dose of 3 mg kg^{-1} TCPOBOP and killed after 2 days. Livers were subjected to the immunohistochemical protocol as described in (Section 2.2.3.2). The number of cells labelled in periportal (PS- gray bars) or centrilobular (CV- black bars) is shown for the control ($n=4$) and treated ($n=5$) group for males and females. The bars show the mean value, and the error bars depict one standard deviation. Statistically significant difference between the CV and the PS is indicated by an asterisk. Statistic was done with Student's *t*-test ($P < 0.05$).

Section 3.2.2.2 Induction of hepatic DNA synthesis by PPAR α ligands (ciprofibrate)

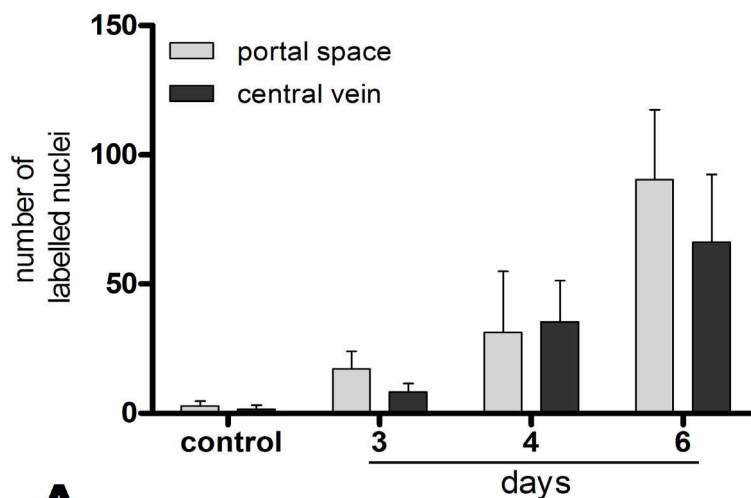
Section 3.2.2.2.1 Zonation in 129S4/SvJae mice treated with ciprofibrate.

This experiment was designed to determine the lobular distribution of hepatocellular proliferation in the liver of 129S4/SvJae mice treated with ciprofibrate.

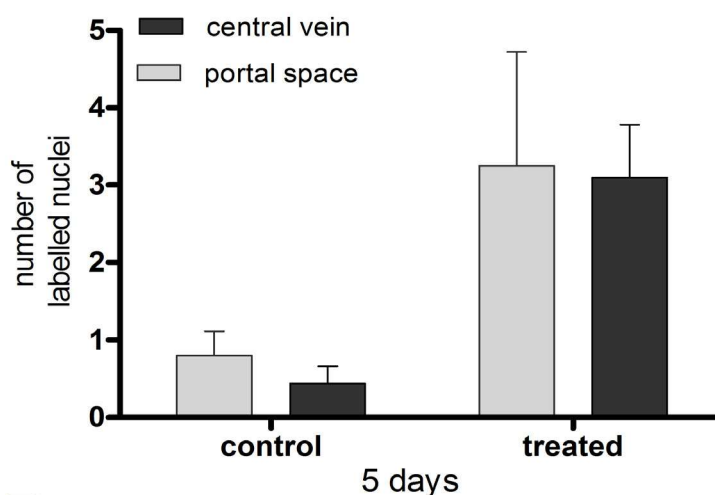
129S4/SvJae male mice were stained for BrdU labelling and treated with 100mg kg⁻¹ day⁻¹ ciprofibrate or corn oil as shown in Section 2.2.1.2 and Section 2.2.1.3. The number of labelled cells in the periportal or centrilobular areas was determined as described earlier in Section 2.2.4.2. Dosing with corn oil was for two days and ciprofibrate was 3, 4 and 6 days Figure 3.21A. There was no significant difference in the number of labelled hepatocytes between the portal space zone and the central vein zone for all the groups. There was also no significant difference in a separate experiment, where mice were administered 300 mg kg⁻¹ ciprofibrate for 5 days Figure 3.21B.

In different experiments of animals treated with corn oil for 4 and 6 days there was no significant difference between the periportal region and the centrilobular. When all the control data (2, 4, 5, and 6 days) were pooled together there still was no significant difference in lobular distribution of labelled hepatocytes (data not shown). Statistics done with Student's t- test (P<0.05).

These results show that hepatic zonal distribution of DNA synthesis in 129S4/SvJae mice treated with corn oil (vehicle)/ ciprofibrate is consistent throughout the liver zones and does not accumulate in any of the periportal or perivenous zones, and there is no effect of time (2, 4, 5 and 6 days) or different doses of ciprofibrate 100, 300 mg kg⁻¹ day⁻¹ on the distribution.



A 100mg/kg ciprofibrate



B 300mg/kg ciprofibrate

Figure 3.21 Lobular distribution of hepatic DNA synthesis in ciprofibrate-treated 129S4/SvJae mice. Male 129S4/SvJae mice aged 9-10 weeks were treated with BrdU and the liver processed and visualization of BrdU-stained hepatocyte nuclei was undertaken as in Section 2.2.4.2. The number of cells labelled in periportal (PS- gray bars) or centrilobular (CV- black bars) is shown, in **A** for animals treated orally with corn oil for the control group and with 100 mg kg⁻¹ Ciprofibrate for 3, 4 and 6 days (the mean is for 5 animals). **B** shows animals treated orally with corn oil (control) or 300 mg kg⁻¹ Ciprofibrate for 5 days (the mean is for 4 animals). There was no significant difference between the number of labelled hepatocytes in portal space and central vein areas for all groups. The bars show the mean value, and the error bars depict one standard deviation. Statistics done was with Student's t- test ($P < 0.05$).

**Section 3.2.2.2.2 Zonation in 129S4/SvJae PPAR α -null (129S4/SvJae-Ppara^{tm1Gonz/}
tm1Gonz) mice treated with ciprofibrate.**

This experiment was designed to determine the lobular distribution of hepatocellular proliferation in liver of male 129S4/SvJae nullizygous for PPAR (the tm1Gonz allele) mice which were treated with 100 mg kg⁻¹ day⁻¹ ciprofibrate.

Male 129S4/SvJae PPAR α -null mice were stained for BrdU labelling and orally dosed with 100 mg kg⁻¹ day⁻¹ ciprofibrate or corn oil (vehicle) for two and four days, as shown in Section 2.2.1.2 and Section 2.2.1.3. The number of labelled cells in the periportal or centrilobular areas were determined as described in Section 2.2.4.2. Figure 3.22 shows that there was no significant difference in the number of labelled hepatocytes between portal space and central vein zones for all the control and treated groups, although the mean value is higher in the PS for the treated groups and the control for 4 days. Statistics were done with Student's t- test P<0.05.

These results indicate that the hepatic zonal distribution of male 129S4/SvJae PPAR α -null mice treated with corn oil (vehicle) or 100 mg kg⁻¹ day⁻¹ ciprofibrate is consistent through out the liver lobule zones and does not significantly accumulate in any of the periportal or perivenous regions.

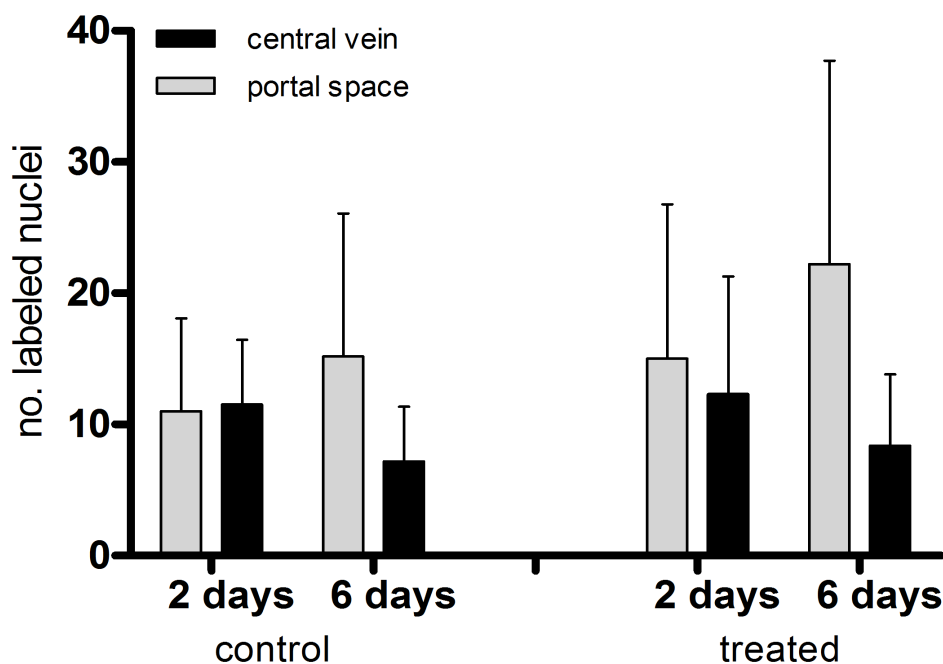


Figure 3.22 Lobular localization of DNA labelling in 129S4/SvJae ($Ppara^{tm1Gonz/tm1Gonz}$) male mice. Male 129S4/SvJae ($Ppara^{tm1Gonz/tm1Gonz}$) mice aged 9-10 weeks were treated with BrdU (Section 2.2.1.2). The number of cells labelled in periportal (PS- gray bars) or centrilobular (CV- black bars) is shown for control or animals treated orally with 100 mg kg^{-1} ciprofibrate for 2 and 6 days (for the control the mean is for 2 and 5 animals), for the treated the number of animals is 3 and 5 for days 2 and 6 respectively). The bars show the mean value, and the error bars depict one standard deviation. There was no difference between portal space and central vein for all the groups with Student's t- test ($P < 0.05$).

Section 3.2.2.2.3 Zonation in C57BL/6JCrI mice treated with ciprofibrate.

The above results did not show a significant preferential periportal distribution of hepatocyte labelling in 129S4/SvJae mice. However, it is possible that there could be zonal distribution of labelling in a different mouse strain, so this experiment was designed to determine the lobular distribution of hepatocellular proliferation in the liver of male C57BL/6JCrI mice treated with ciprofibrate. Male C57BL/6JCrI mice were stained for BrdU labelling as shown in Section 3.1.1.3.

The number of labelled cells in the periportal or centrilobular areas were determined as described in Section 2.2.4.2. The mice were dosed orally with $100 \text{ mg kg}^{-1} \text{ day}^{-1}$ ciprofibrate or

corn oil for two and four days Figure 3.23. There was no significant difference in the number of labelled hepatocytes found around the portal space and central vein zones for all the groups, although the mean value is higher in the PS for all the groups. Statistics were done with Student's t- test, $P < 0.05$. ($n=6$).

These results show that the hepatic zonal distribution of DNA synthesis in C57BL/6JCr1 male mice treated with corn oil/ ciprofibrate for 2 and 4 days is consistent throughout the liver zones and does not significantly accumulate in any of the periportal or perivenous zones.

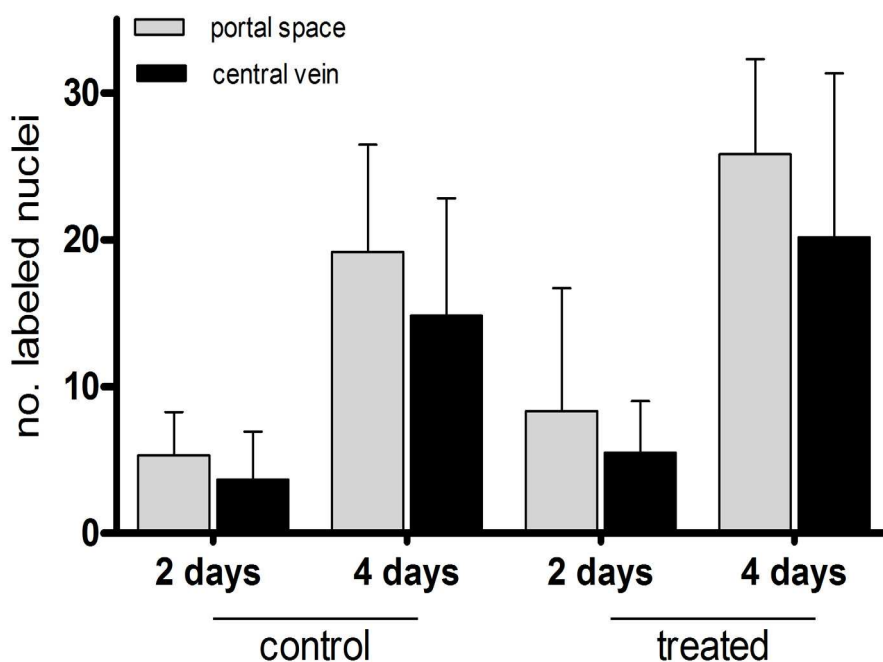


Figure 3.23 Lobular localization of hepatic DNA labelling in C57BL/6JCr1 mice treated with ciprofibrate. Male C57BL/6JCr1 mice aged 9-10 weeks were treated with BrdU as in Section 2.2.1.2. The number of cells labelled in periportal (PS- gray bars) or centrilobular (CV- black bars) is shown for animals treated orally with control (corn oil) or $100 \text{ mg kg}^{-1} \text{ day}^{-1}$ ciprofibrate for 2 or 4 days. The bars show the mean value, and the error bars depict one standard deviation for 6 animals. There was no difference between portal space and central vein for all the groups with Student's t- test ($P < 0.05$).

Section 3.2.2.2.4 Zonation in DBA/2JCrI mice treated with ciprofibrate.

This experiment was designed to determine the lobular distribution of hepatocellular proliferation in the liver of male DBA/2JCrI mice treated with ciprofibrate.

Male DBA/2JCrI mice were stained for BrdU labelling as shown in Section 3.1.1.2.2. The number of labelled cells in the periportal or centrilobular areas were determined as described in Section 2.2.4.2. The mice were dosed orally with corn oil for two and six days or $100 \text{ mg kg}^{-1} \text{ day}^{-1}$ ciprofibrate for two, four and six days Figure 3.24. There was no significant difference in the number of labelled hepatocytes found around the portal space and central vein zones for all the groups.

These results show the hepatic zonal distribution of DNA synthesis in DBA/2JCrI male mice treated with corn oil (vehicle)/ ciprofibrate for two, four and six days is consistent throughout the liver zones and does not accumulate significantly in any of the periportal or perivenous zones.

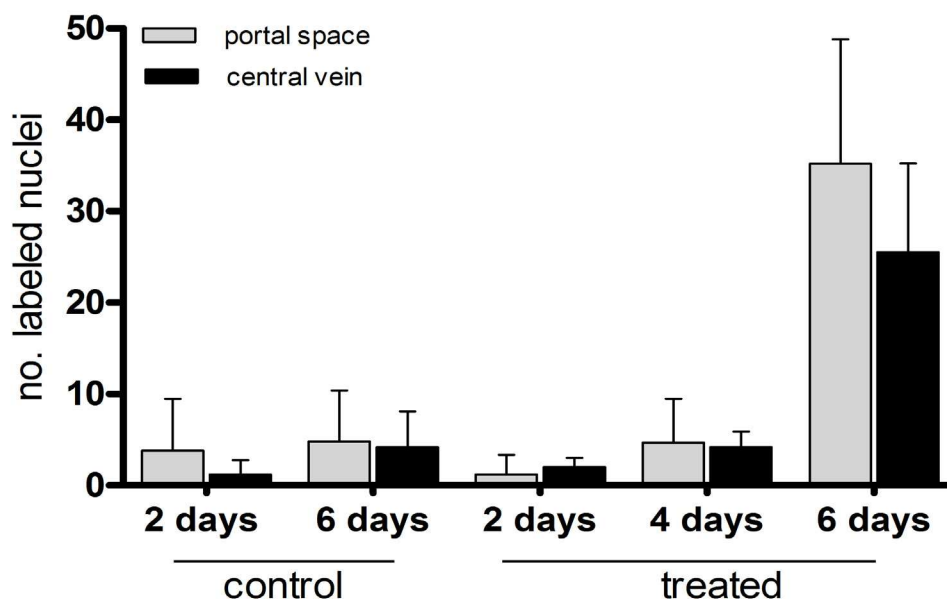


Figure 3.24 Lobular localization of hepatic DNA labelling in DBA/2JCrI mice treated with ciprofibrate. Male DBA/2JCrI mice aged 9-10 weeks were treated with BrdU as in Section 2.2.1.2. The number of cells labelled in periportal (PS- gray bars) or centrilobular (CV- black bars) is shown for animals treated orally with control (corn oil) for 2 and 6 days or 100 mg kg⁻¹day⁻¹ ciprofibrate for 2, 4 or 6 days. The bars show the mean value, and the error bars depict one standard deviation for 6 animals. No difference between portal space and central vein for all the groups with Student's t- test ($P < 0.05$).

Section 3.2.2.3 Induction of hepatic DNA synthesis by the PPAR α methylclofenapate.

From the results before, it was possible that the failure to observe lobular localization of hepatocyte DNA synthesis was due to an idiosyncratic response to ciprofibrate. Therefore, the peroxisome proliferator methylclofenapate (MCP) was used, as an agent that has previously been shown to lead to zoned DNA synthesis induction in rats (Barrass et al., 1993).

Section 3.2.2.3.1 Zonation in 129S4/SvJae mice treated with MCP

This experiment was designed to determine the lobular distribution of hepatocellular proliferation in the liver of male 129S4/SvJae mice treated with 25 mg kg⁻¹day⁻¹ MCP for 3 and 4 days. Male 129S4/SvJae mice aged 9-10 weeks were stained for BrdU labelling as shown in Section

2.2.1.2. The number of labelled cells in the periportal or centrilobular areas were determined (as described in Section 2.2.4.2).

The mice were dosed orally with corn oil for four days or $25 \text{ mg kg}^{-1} \text{ day}^{-1}$ MCP for 3 and 4 days Figure 3.25. There was no significant difference in the number of labelled hepatocytes found around the portal space and the central vein zones for all the 129S4/SvJae male mice groups when treated with $25 \text{ mg kg}^{-1} \text{ day}^{-1}$ MCP for 3 and 4 days.

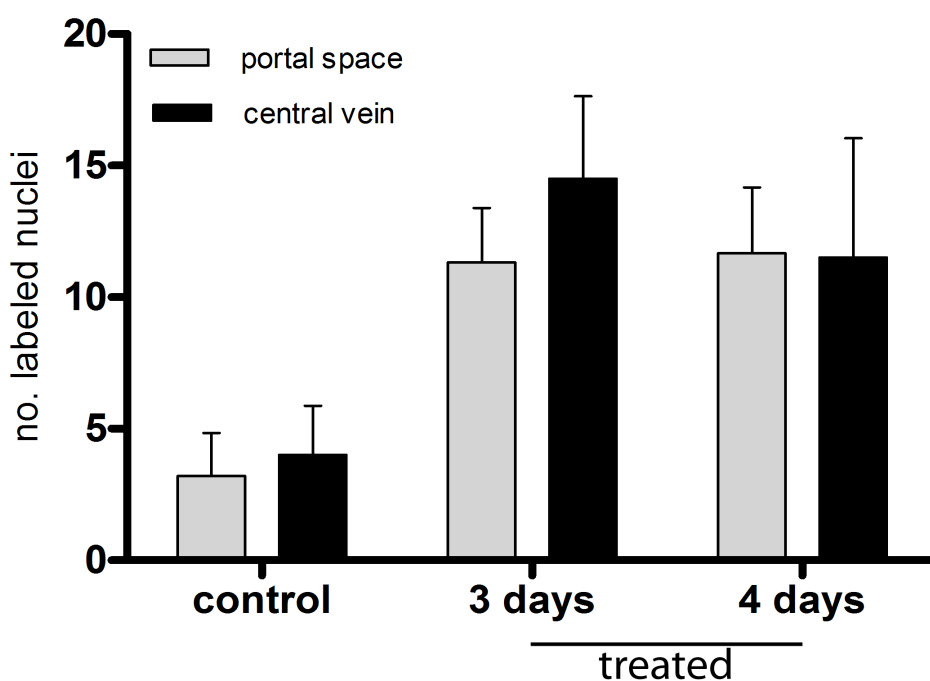


Figure 3.25 Zonal distribution of hepatic DNA synthesis in 129S4/SvJae mice treated with MCP. Male 129S4/SvJae mice aged 9-10 weeks were treated with BrdU as in Section 2.2.1.2 The number of cells labelled in periportal (PS- gray bars) or centrilobular (CV- black bars) is shown for control for 4 days (the number of animals is 5), or treated with $25 \text{ mg kg}^{-1} \text{ day}^{-1}$ MCP methylclofenapate for 3 and 4 days (the number of animals is 6). Visualization of BrdU-stained hepatocyte nuclei was undertaken as mentioned in Section 2.2.4.2. The bars show the mean value, and the error bars depict one standard deviation. There was no significant difference between portal space and central vein for all the groups according to Student's t- test ($P < 0.05$).

Section 3.2.2.3.2 Zonation in Alderley park (AP) mice treated with MCP.

This experiment was designed to determine the lobular distribution of hepatocellular proliferation in liver of male Alderley park [AP] mice treated orally with $25 \text{ mg kg}^{-1}\text{day}^{-1}$ MCP for 3 and 4 days.

Male Alderley park mice aged 9-10 weeks were stained for BrdU labelling as shown in Section 2.2.1.2. The number of labelled cells in the periportal or centrilobular areas were determined.

It was found in male AP mice treated with $25 \text{ mg kg}^{-1}\text{day}^{-1}$ MCP that there was a significant difference in the number of labelled hepatocytes between the periportal and the perivenous zones (statistics used was with Student's t- test, $P < 0.005$) for the control and the treated group for 4 days with MCP. The results show a larger distribution of hepatic DNA synthesis at the periportal. As for the group treated for 3 days, there was no significant difference between the number of labelled cells situated around the PS and CV (Figure 3.26).

From this result it is established that the hepatic zonal distribution between the periportal and perivenous regions is not significantly different for male AP mice dosed with $25 \text{ mg kg}^{-1}\text{day}^{-1}$ MCP for 3 days. As for the AP mice dosed for 4 days with MCP and the control group, the distribution of hepatic DNA synthesis was primarily periportal.

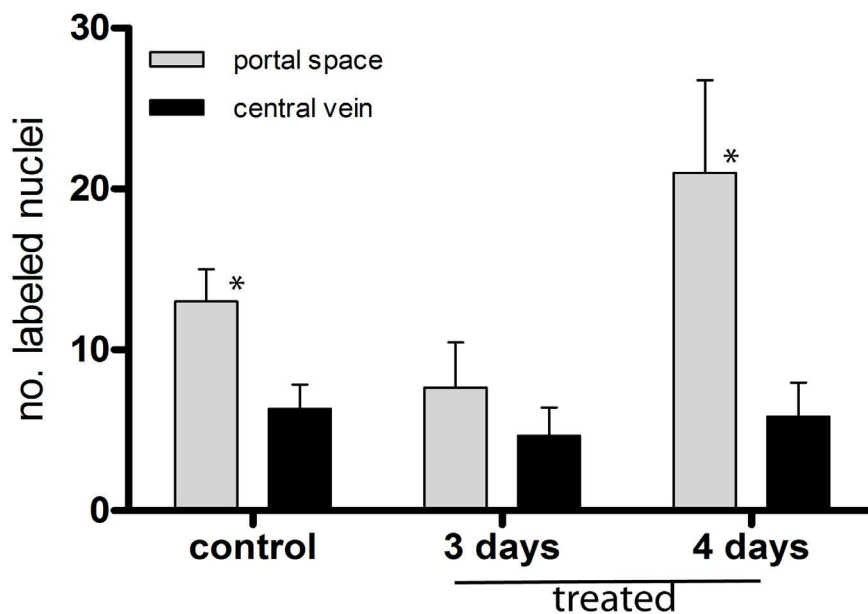


Figure 3.26 Zonal distribution of hepatic DNA synthesis in AP mice treated with MCP. Male AP mice aged 14-15 weeks old were treated with BrdU as described earlier in Figure 3.25. The number of cells labelled in periportal (PS- gray bars) or centrilobular (CV- black bars) is shown for control (4 days) or treated with $25 \text{ mg kg}^{-1} \text{ day}^{-1}$ MCP methyleclofenapate for 3 and 4 days (the number of animals is 6). Visualization of BrdU-stained hepatocyte nuclei was undertaken as mentioned in Section 2.2.4.2. The bars show the mean value, and the error bars depict one standard deviation. Statistically significant difference from the control group is indicated by an asterisk according to Student's t- test ($P < 0.05$).

Section 3.3 Microarray analysis

Knowledge from the previous results Section 3.1.2 that hepatic DNA synthesis occurs 24 hours after dosing, and the fact that cells need from 18-20 hours to undergo DNA synthesis, suggests that the transcriptional signal is sensed within the first five hours after dosing. From this, the 1-5 hour window was targeted in order to specify the early genes that may be induced, and which could be responsible for hepatic DNA synthesis at 24 hours.

The microarray analysis was applied as in Section 2.2.5 to measure the expression levels of genes induced by 50 mg kg⁻¹ ciprofibrate in livers of F-344NHsd rats after 1, 3, 5 and 24 hours in comparison with the control.

A number of controls were used;

- Corn oil (vehicle) dosed rats were used as negative controls.
- Rats dosed with PXR ligands (cyproterone acetate CPA) were used as positive controls, to compare genes that are involved in hepatic DNA synthesis, but which might not be specific for the PPAR α ligands (ciprofibrate).
- 24 hour points were used to confirm the induction happens at 24 hours by the immunohistochemical protocol and to rule out genes that would be induced after 24 hours as these genes are not of our interest.
- A general or common reference/control which is the most commonly used design of microarray experiments. This also has the benefit of a competent comparison of samples (Cherkaoui-Malki et al., 2001).

In Figure 3.27 is a diagram that clarifies the experiment design. This experiment was done in parallel with the time course done in Section 3.1.2.



Figure 3.27 Time course of hepatic DNA synthesis in F-344NHsd rats. Groups of F-344NHsd rats were dosed by gavage with 50 mg kg⁻¹ ciprofibrate or 100mg kg⁻¹ cyproterone acetate or corn oil, then killed after 1, 3, 5 and 24 hours. The groups killed after 24 hours were dosed intraperitoneally two hours before killing with BrdU and immunohistochemical protocol was undertaken (Section 2.1.3). Each point represents the number of animals is 4 for the time points 1, 3 and 5 hours and the number of animals is 5 for the group treated with ciprofibrate after 24 hour. The number of animals used is 6 for the groups treated with CPA and corn oil killed after 24 hours. Microarray analysis was done on all the time points at 1, 3, 5 and 24 hours in 4 biological replicates and 2 technical replicates.

Figure 3.28 shows that in this experiment there was a statistically significant induction of hepatic DNA synthesis after 24 hours for the group treated with ciprofibrate and even higher for the group treated with cyproterone acetate, according to the measurements of hepatic labelling index with immunohistochemistry.

Liver weight to body weight ratios were calculated for all the time points (Figure 3.29), and there was no significant difference between the control and treated (ciprofibrate/ CPA) groups for all the time points (1, 3, 5, and 24 hours).

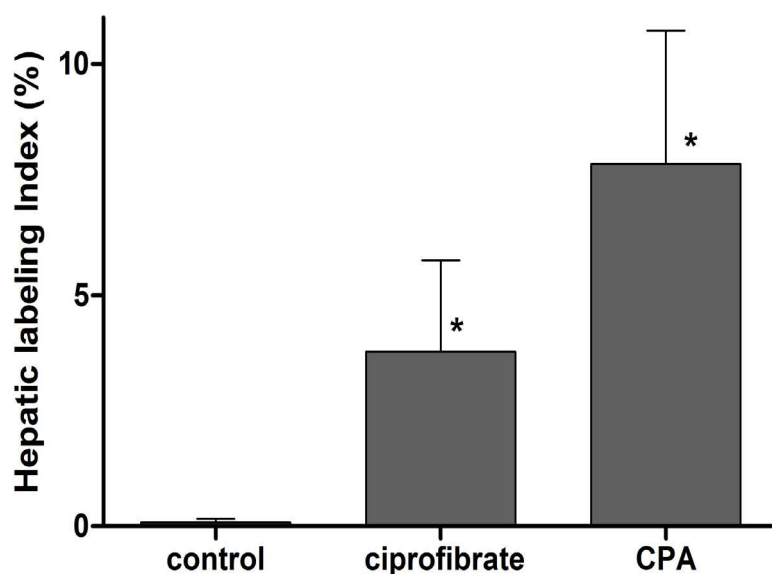


Figure 3.28 Effects of ciprofibrate and CPA on hepatic DNA synthesis in F-344NHsd rats after 24 hours of dosing. Male F-344NHsd rats aged 14-15 weeks were dosed with corn oil (vehicle), 50 mg kg⁻¹ ciprofibrate or 100 mg kg⁻¹ cyproterone acetate CPA, then killed and assayed after 24 hours (Section 2.1.1). All data are expressed as mean±SD for 6 animals. Statistically significant differences from the control group are indicated by an asterisk. Statistics were performed with Dunnett's multiple comparison test (P<0.05), exception for the number of animals in the ciprofibrate group was 5.

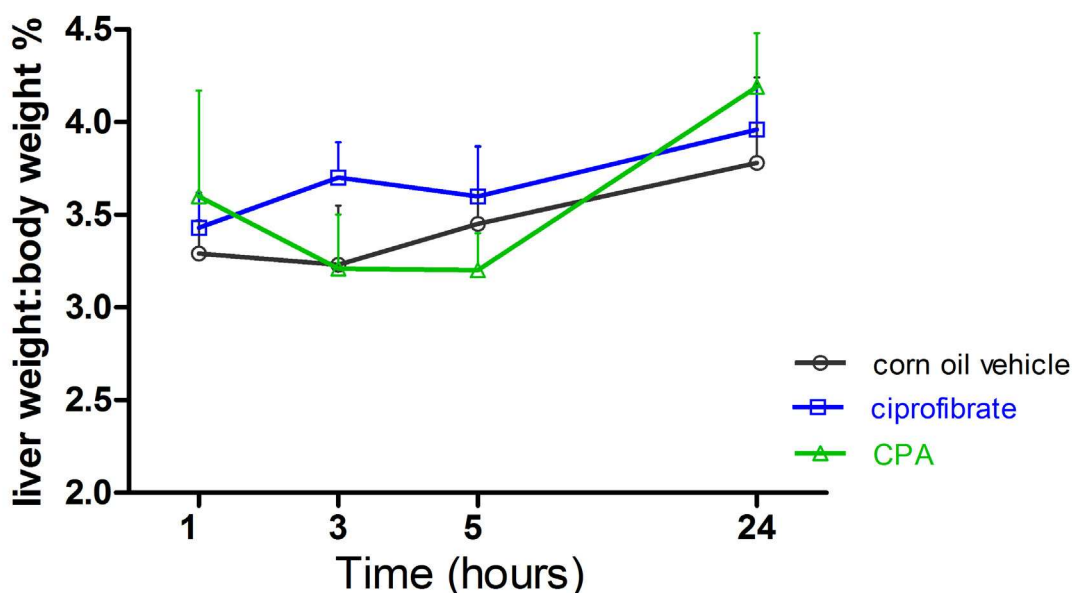


Figure 3.29 Effects of ciprofibrate and CPA on liver weight in F-344NHsd rats. Male F-344NHsd rats aged 14-15 weeks were dosed with corn oil (vehicle), 50 mg kg⁻¹ ciprofibrate or 100 mg kg⁻¹ cyproterone acetate (CPA), then killed after 1, 3, 5 and 24 hours. Liver weight to body weight ratio was determined as in Section 2.2.2. All data are expressed as mean±SD. There was no significant difference between the treated and control groups with Dunnett's multiple comparison test (P<0.05). n=4 for the time points 1, 3 and 5 hours, n=6 for time point 24 hours corn oil and CPA, n=5 for the time point 24 hours treated with ciprofibrate (n=number of animals).

Section 3.3.1 Assessment of RNA quality and quantity

The liver tissue sampled at intervals after dosing was snap frozen in liquid nitrogen and stored in a -80°C freezer until use. RNA isolation was processed with TRI reagent as in Section 2.2.5.1 for all the liver tissues. Preliminary experiments with a mini prep method and the TRI reagent method showed that the higher quality RNA preparations were established with the TRI reagent, as shown in Figure 3.30A where the 28S and 18S rRNA bands are sharper in lanes 3 and 4 while the RNA in lanes 6 and 7 (the mini prep) was more smeared. Also the TRI reagent method (when overloading the RNA on the agarose gel; Figure 3.30B) showed the 5S band indicating that the microRNA was intact.

The quality of the total RNA for all the samples was assessed by the relative intensities of the 28S and 18S rRNA bands visualized by electrophoresis on 2% agarose gels with a 1kb Plus DNA ladder and stained with ethidium bromide. Figure 3.31 and Figure 3.32 show intact total RNA with sharp 28S and 18S ribosomal RNA bands. The 28S rRNA band is approximately twice as intense as the 18S rRNA band. The 5S rRNA was determined by comparing with the literature and with the ladder and appears as a faint band (in most of the lanes that have been slightly overloaded) indicating the microRNAs.

The yield of the RNA was measured by a CECIL CE9500 spectrophotometer, and the total yield of each of the RNAs from the liver tissues were in a range between $8.2 \pm 3.9 \mu\text{g mg}^{-1}$. A reference RNA control was made by mixing aliquots of all the RNAs to be used in the microarray hybridization -to remove any bias from the 555 and 647 Alexa dyes.

From the above analysis it was established that the TRI reagent method works well, yielding intact microRNAs. All the RNAs used were of good quality and had a yield sufficiently high for microarray analysis.

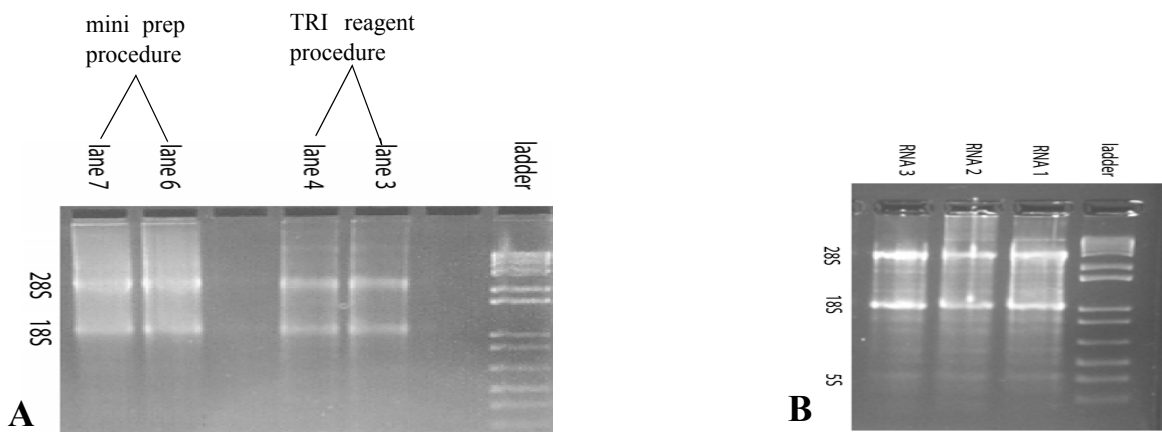


Figure 3.30 Optimizing RNA isolation from rat liver. Agarose gels (2%) stained with ethidium bromide as shown in Section 2.2.8. In A, the ladder is a 1kb Plus DNA ladder, lanes 3 and 4 have RNA made by the TRI reagent procedure; lanes 6 and 7 have RNA made with the mini prep procedure. In B, the ladder is a 1kb Plus DNA ladder, RNAs in lanes 1, 2 and 3 were made with TRI reagent method. A larger amount of the RNA was loaded to show the microRNA band at 5S as a faint band. The major RNA bands are shown at 28S and 18S.

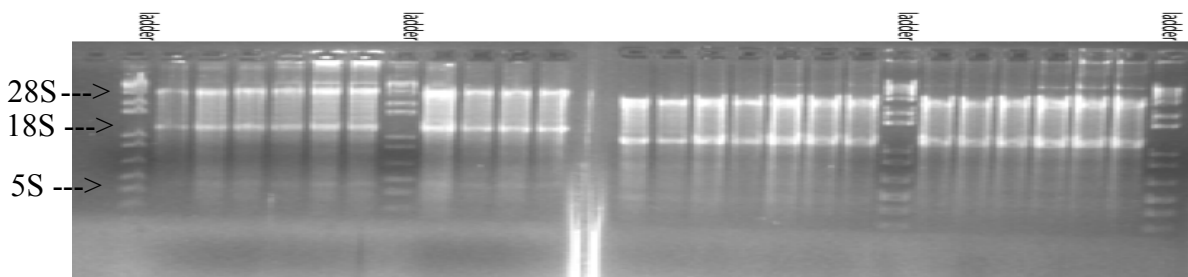


Figure 3.31 Agarose gel with RNA from F-344NHsd rat liver. Agarose gel electrophoresis (2%) was performed as described in Section 2.2.8 with a 1kb Plus DNA ladder to mark the nuclear weight of the RNA and dyed with ethidium bromide. The lanes were loaded with RNA samples from F-344NHsd rat liver treated with ciprofibrate for 24 hours.

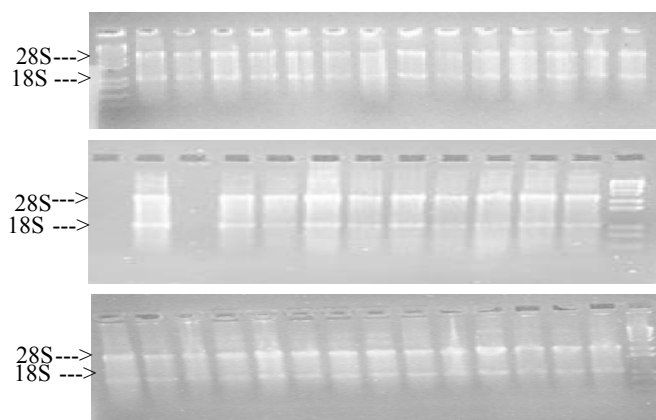


Figure 3.32 More examples of agarose gels loaded with RNAs from F-344NHsd rats. Agarose gel electrophoresis was made as described in Figure 3.30 with a 1kb Plus DNA ladder to mark the nuclear weight of the RNA and dyed with ethidiome bromide. Intact total RNA with sharp 28S and 18S ribosomal RNA bands are shown. The lanes were loaded with RNA samples from F-344NHsd rat liver treated with CPA.

Section 3.3.2 Incorporation of Alexa dyes in the cDNA

RNAs were made to cDNAs labelled with Alexa dyes 555 and 647. The Amino allyl cDNA labelling used a two step process, and all the steps were made with a negative control (sample with no RNA) and a positive control (RNA supplied in the kit). All the samples' incorporation of the Alexa dyes and the cDNA quantities were determined using a nanodrop machine, and the cDNA amounts were around $80 \pm 20 \text{ ng } \mu\text{l}^{-1}$ (and might go up to $200 \text{ ng } \mu\text{l}^{-1}$), and the incorporated dye was approximately $8 \pm 4 \text{ pmol } \mu\text{l}^{-1}$ (and might go up to $20 \text{ pmol } \mu\text{l}^{-1}$) for each of the Alexa dye 555 and 647 (Appendix I). The Amino allyl cDNA labelling kit -from Ambion- was used for the rest of the analysis, and optimizing was done to dilute the cDNA, and also to decrease the costs as will be described in Section 3.3.3.

Section 3.3.3 Optimization of the microarray technique

Figure 3.33 shows a typical outcome of the scanned "MEEBO" -Mouse genome set- microarray slide. The scanner functions using sequential scanning to create 3 images: the slide is scanned twice; first using the green 532 nm laser that scans the 555 Alexa dye (Figure 3.33A), and then the red 635 nm laser and this scans the 647 Alexa Dye (Figure 3.33B). The third slide with a ratio image where the second wavelength image is composed on the first as in Figure 3.33C, where a green spot indicates the treatment for this specific gene has less activity than the reference control; a red spot indicates the treatment would have more activity than the reference control; and a yellow spot means that there is no change in the level of activity between the two populations of test and reference control would be where the control and treated are the same. Notice for each of the red and the green scans there are white spots indicating saturating fluorescence.

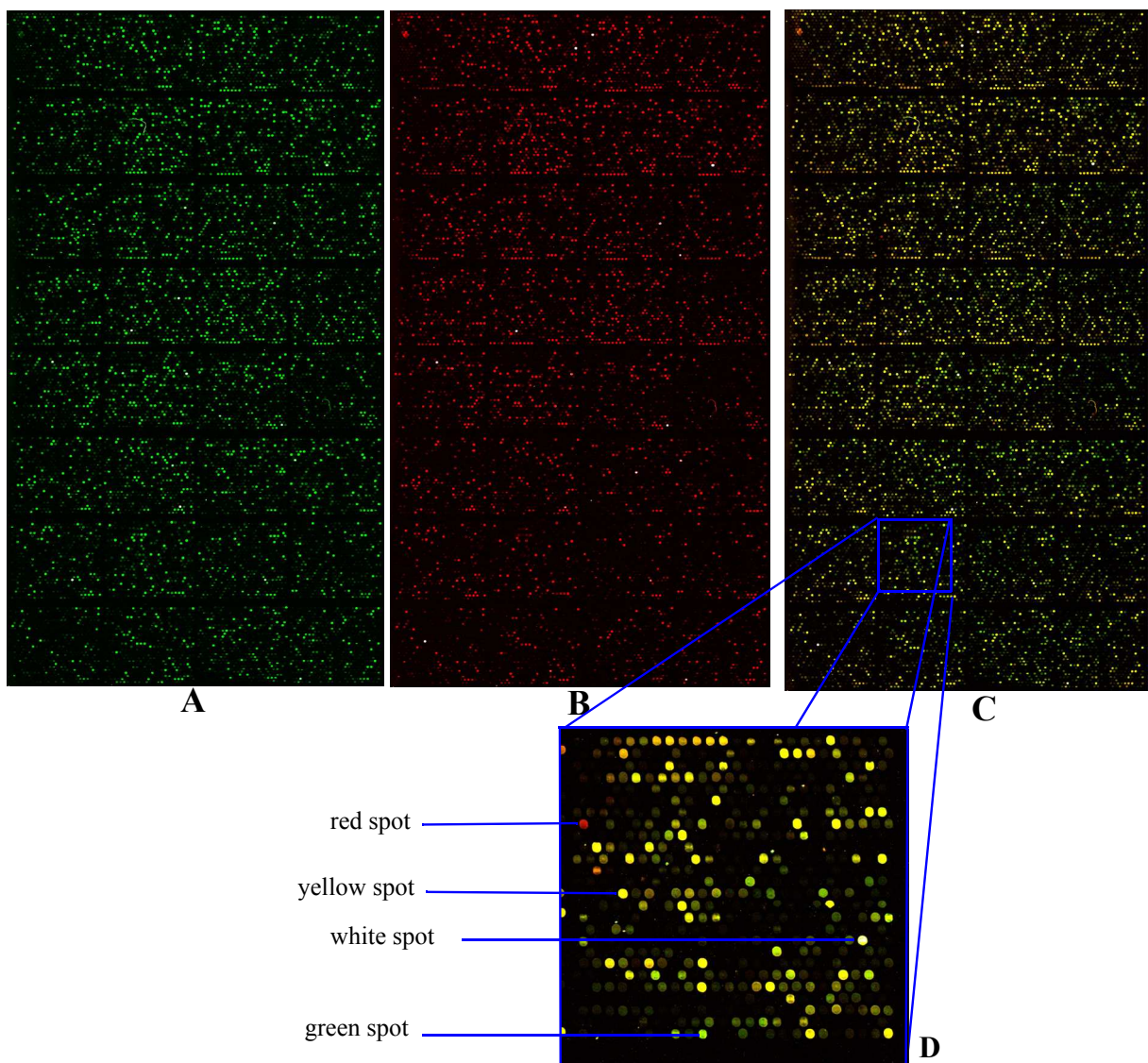


Figure 3.33 Images of the scanned microarray slide. Outcome of the Axon 4200 scanner, **A** shows the image with the green 555 fluorescent. **B** shows the image with the red 674 fluorescent. **C+D** shows the combined colours, where a green spot indicates the treatment for this specific gene has less activity than the reference control; a red spot indicates the treatment would have more activity than the reference control; and a yellow spot means that there is no change in the level of activity between the two groups control and treated. Notice for each of the red and the green scans there are white spots indicating a large intensity of the fluorescent.

It was of importance to validate the microarray technique to ensure the results are reliable and repeatable with the least expense possible. So, the following issues were addressed in an independent experiment: (1) Section 3.3.3.1- Reproducibility of the scanning. (2) Section 3.3.3.2- Reproducibility of the hybridizing. (3) Section 3.3.3.3- Testing of the concentration of cDNA. For optimizing the microarray technique, RNA extracted from liver of F-344NHsd rats treated with ciprofibrate at 24 hours was used. Groups of the same cDNA were labelled and hybridized as described in Section 2.1.4.2. They were then all scanned. A random group was chosen to be scanned repeatedly. All groups then went through the analyzing process, then normalised using the NORTT program, and then compared against each other.

Section 3.3.3.1 Reproducibility of scanning

The aim of this experiment was to determine that the scanning section of the microarray technique is reliable and the results are repeatable. Slides were chosen at random and each slide was scanned twice. The data from the scanner was normalised and one set of scan results was compared against the other.

The results in Figure 3.34 shows the linear regression between the first and the second scan at a 1:1 ratio, which means that a specific gene that is high in one scan is also high in the other scan, and the ones that are low in one, would be low in the other scan. As the $r^2 =$ from 0.9 to 0.6, it can be said that the two scans of the same slide are 60-90% reproducible.

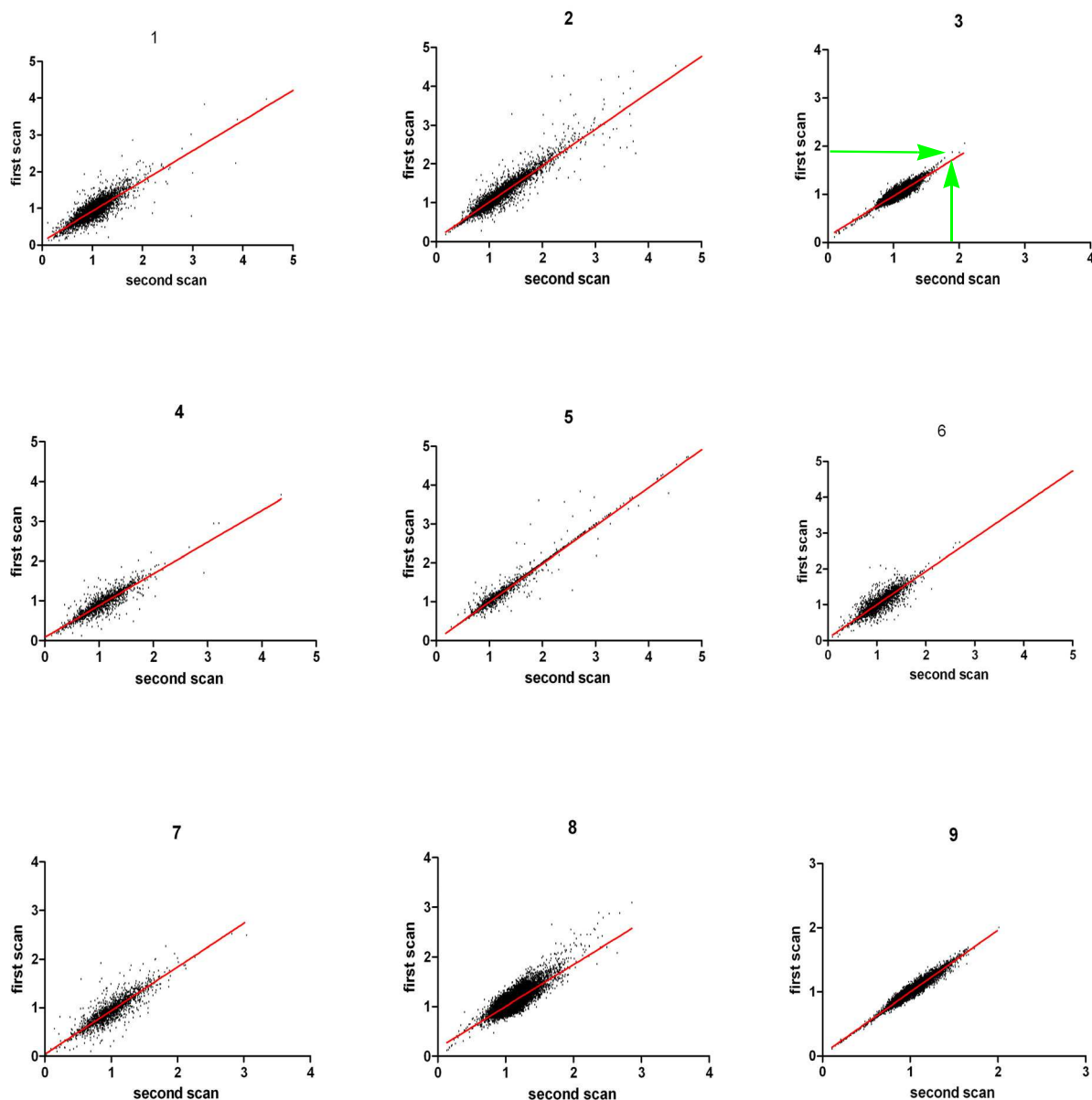


Figure 3.34 Reproducibility of scanning. Each plot shows a comparison between two set of genes of the same slide when scanned twice. On the Y axis is a set of the fold-inducion of the normalised log2 ratio of median for the first scan, and the X axis is the same for the second scan, the linear regression line is at a angle at $\sim 45^\circ$, $r^2=0.8\pm 0.1$. The 2 green arrows are pointing to the same spot (gene) indicating that this gene’s fold induction is just under 2 in both of the scans, and this is the same for most of the genes.

Section 3.3.3.2 Reproducibility of hybridization

It was necessary to ensure that the hybridization in this protocol was repeatable. In this section cDNAs were hybridized on three different sets of slides, then the slides were scanned, and the results from the scans were normalised as shown in Section 2.2.5. The results were then compared. Preliminary experiments showed that, when the same cDNA is hybridized on different slides, they did not give repeatable results as in Figure 3.35. The $r^2=0.2\pm 0.1$ meaning; when comparing a gene from slide 1 against the same gene in slide 2 it gives a different fold induction. Thus the same cDNA when hybridized on different slides gives a 20-30% similarity. This was a problem, and results from this hybridizations were not reliable.

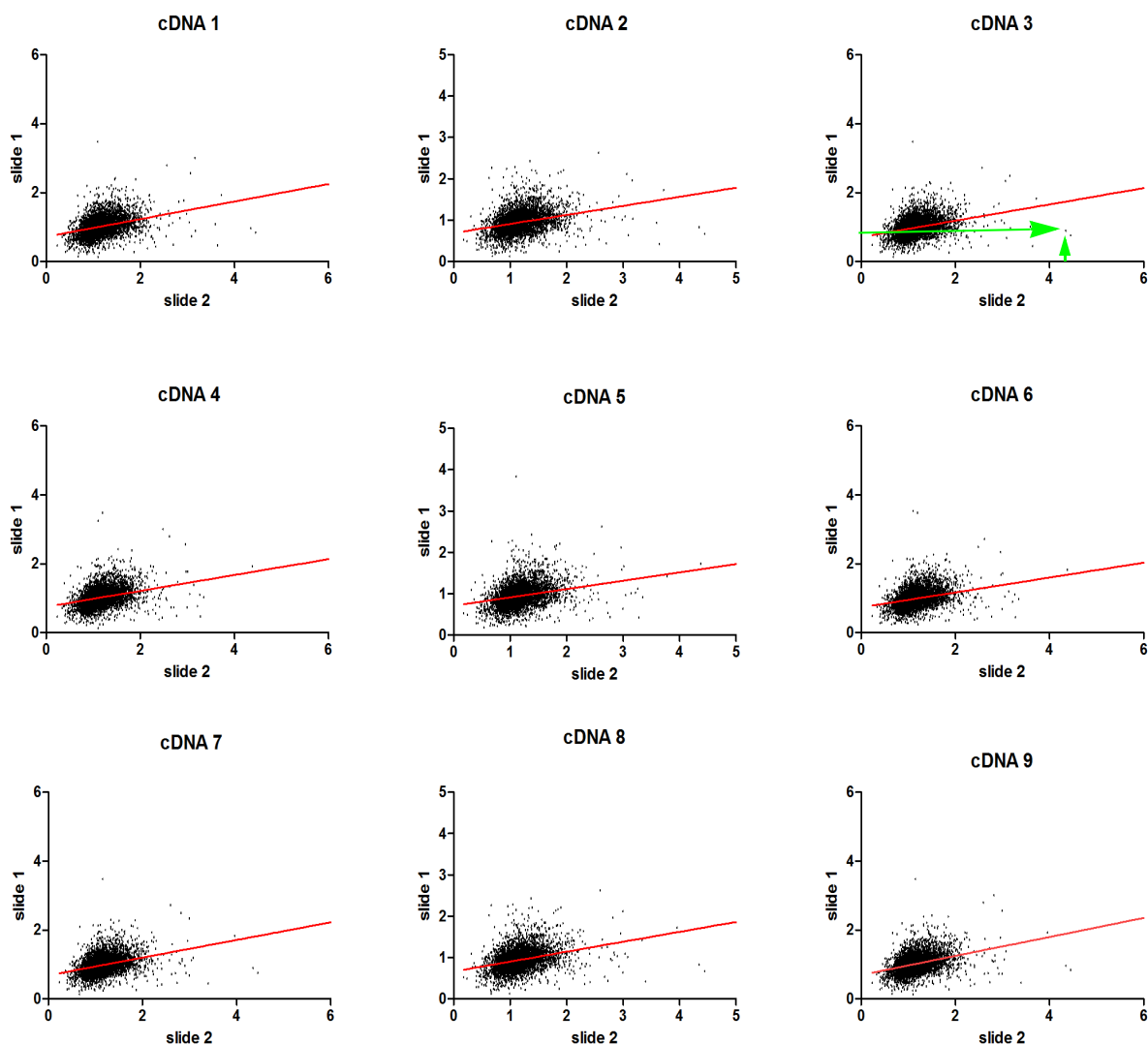


Figure 3.35 Reproducibility of hybridization. Each plot shows two sets of genes from the same cDNA hybridized on different microarray slides, processed as in Figure 3.34. The genes then plotted, the Y axis is a set of the fold-induction of the normalised log₂ ratio of median for the first slide, and the X axis is the same for the second slide. The line presents the linear regression line, the $r^2=0.2\pm 0.1$. The green arrows are pointing to the same spot (gene) indicating that the genes fold induction is different in each of the slides (under 1 in the first slide and over 4 in the second slide).

It was essential to address the issue of not being able to get the same results from the same cDNA hybridized on different slides before being able to consider using the microarray technique for further analysis. Therefore looking into the microarray procedure systematically by checking all the steps (hybridizing, scanning and fitting features) was a reliable technique and the results were comparable to each other and gave a higher r^2 (Figure 3.36).

The main area where the error could be made was misplacing the gene names on the gene spots of the slides (human error) when featuring the scan to the GAL file. The improvement here is that when fitting the features greater care was taken and there were also some markers on the slides to indicate where exactly each gene is situated. From the above information it was decided that all the hybridizations have to have technical replicates that were at least 60% similar to be considered in the analysis.

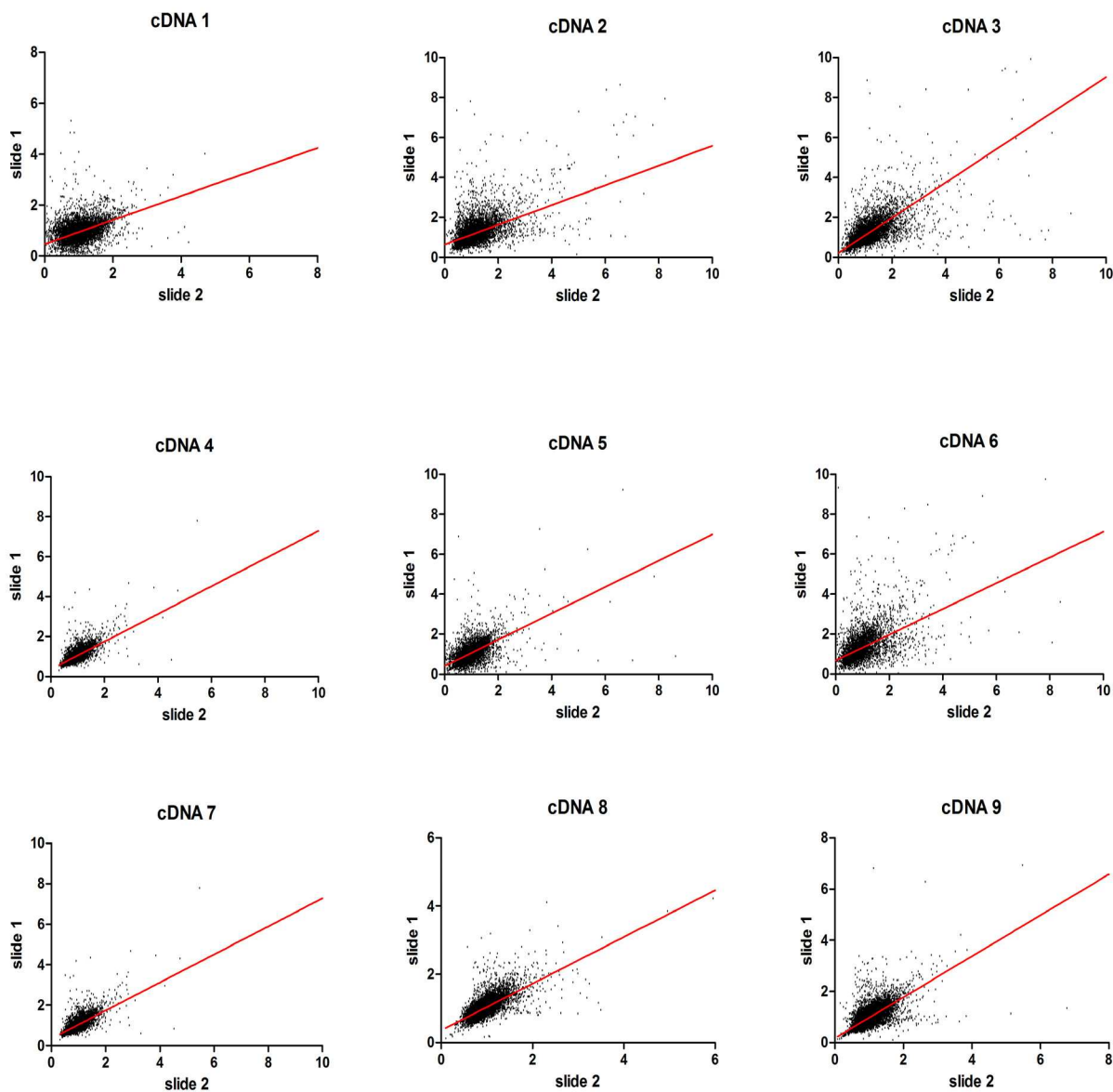


Figure 3.36 Reproducibility of hybridization (improved). Each of the figures show two sets of genes of the same cDNA hybridized on different slides, then processed as in Figure 3.34. The genes were then plotted; on the Y axis is a set of the fold-induction of the normalised log2 ratio of median for the first slide, and on X the same for the second slide. The linear regression line is shown in red and had a $r^2=0.6\pm 0.1$. The improvement here is when fitting the features more care was taken and there were also some markers on the slides to indicate where exactly each gene is situated on the slide.

Section 3.3.3.3 Testing of the cDNA concentration

Because of the expense of the cDNA labelling kit it was desirable to try to decrease these expenses to make it possible to continue with the rest of the planned experiments. So, it was decided to cut the costs in half by diluting the cDNA and spreading what is usually spread on one set of slides to be spread on two sets of slides (to be used as technical replicates). Figure 3.37 shows a comparison between the undiluted cDNA (concentrated 50:1:50 μ l of 80ng μ l⁻¹ labelled-cDNA: tRNA: hybridization buffer) and the diluted cDNA (50:30:2:80 μ l of 80ng μ l⁻¹ labelled-cDNA:DEPC H₂O:tRNA: hybridization buffer).

An experiment was designed where 4 replicates of the same cDNA was labelled (the cDNA was from a sample of liver treated with ciprofibrate and killed after 24 hours). cDNA 1, 2 and 3 were hybridized as concentrated, cDNA 4 was diluted and hybridized on 2 sets of slides. The slides were then compared, the diluted against the undiluted.

When comparing the technical replicates, diluted vs. diluted, non-diluted vs. non-diluted and diluted vs. non-diluted, they all gave a similar results of 60% similarity (the mean and standard deviation of all the $r^2=0.6\pm 0.1$). This means that the dilution of the cDNA sample did not affect its quality. This dilution method was used for the rest of the experiments.

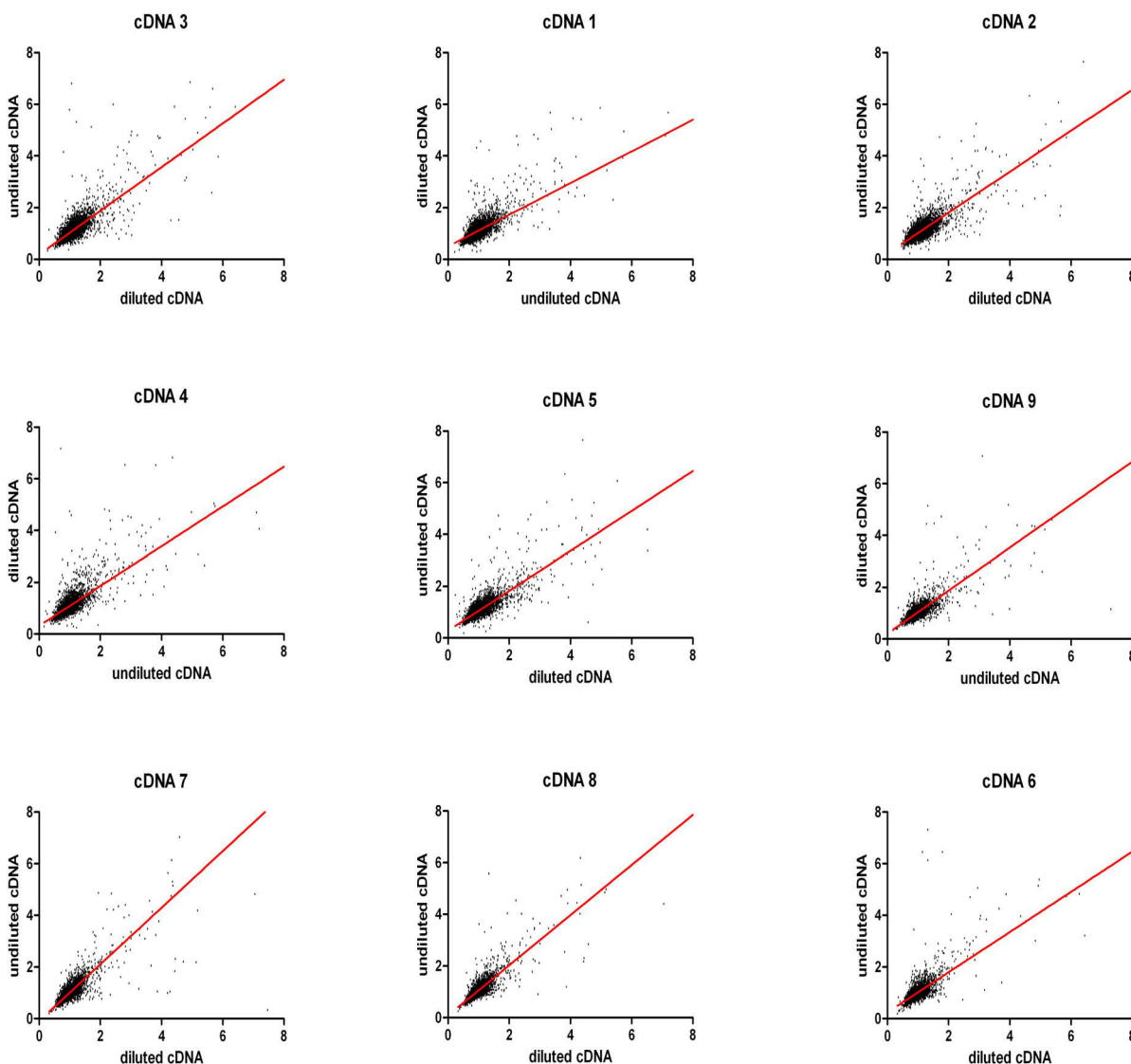


Figure 3.37 Testing of the cDNA concentration. Each of the figures shows a comparison between the diluted cDNA and undiluted cDNA. Hybridized on different slides, then processed as in Figure 3.34. The genes were then plotted as in Figure 3.36. The linear regression line is shown in red and the mean and standard deviation of all the samples is $r^2=0.6\pm0.1$.

From these results, it was found that the optimal way to resolve inconsistently issues in microarray procedures was to:

- 1- Use more replicates, so 4 biological replicates and 2 technical replicates were used.
- 2- Compare the technical replicates against each other and accept only the results that the r^2 is more than 0.6.

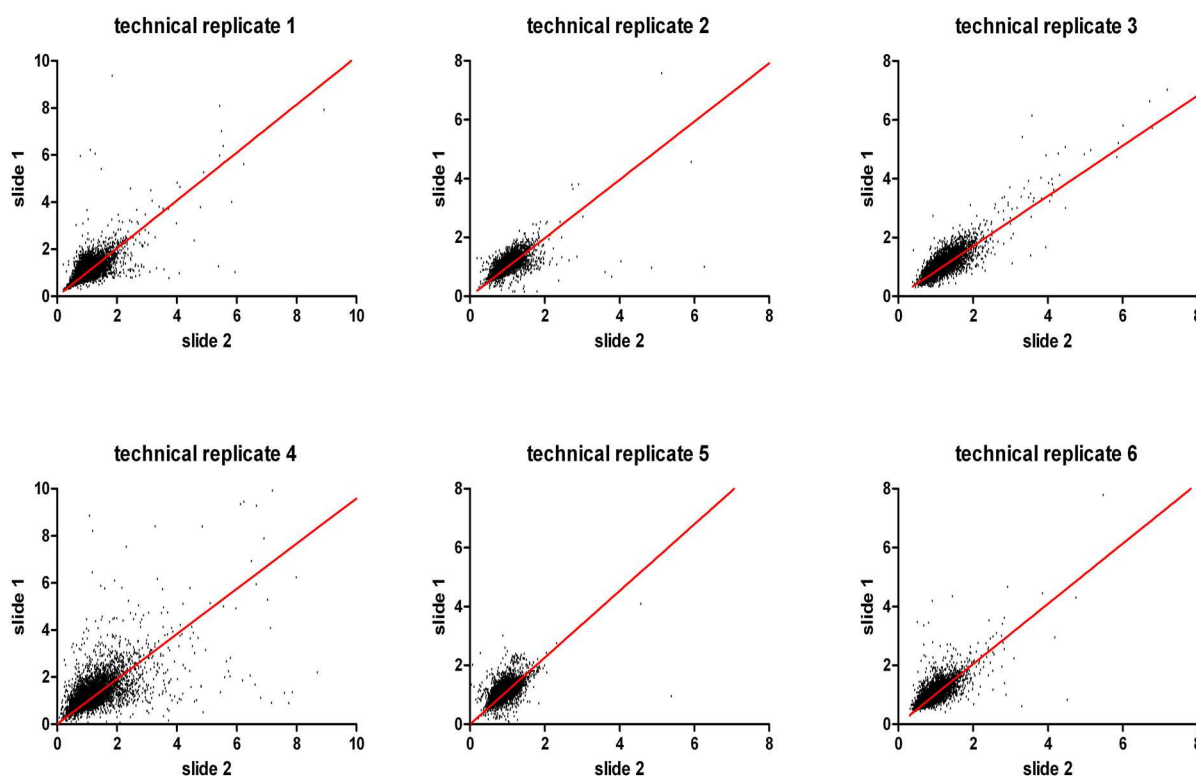


Figure 3.38 Examples of technical replicates . Each plot shows a comparison of technical replicates from the same diluted cDNA. The samples were hybridized on different slides, then processed as in Figure 3.34. The genes were then plotted as in Figure 3.36. The linear regression line is shown in red and the mean and standard deviation of the $r^2=0.6\pm 0.1$.

Section 3.3.4 Microarray analysis of dosed rats after 24 hours

In order to measure the expression levels of early genes that may be up-regulated after dosing of ciprofibrate, and might be responsible for the induction of hepatic DNA synthesis, microarray analysis was applied as in Section 2.2.5 on F-344 fisher rats dosed with 50 mg kg⁻¹ ciprofibrate, or 100 mg kg⁻¹ CPA, or corn oil and killed after 1, 3, 5 and 24 hours in comparison with the control.

Preliminary results were established from microarray analysis done on F-344 fisher rats treated with ciprofibrate, CPA and corn oil for 24 hours. This was to help ensure that the analysis and processing of the data is reliable, by checking for induction of specific genes (used as markers) that are known from the literature to be induced after 24 hours of dosing with ciprofibrate and CPA.

Analysis of the 24 hour groups with R software revealed 1597 genes whose expression was significantly changed (FDR p-value <0.05). The whole list is attached in a CD at the end of the Thesis. Table 3.2 shows a list of up-regulated genes for samples treated with ciprofibrate for 24 hours, that have a fold induction of >1.2 fold change. These include Cyp4A1 (CYP4A10), a gene that is a known marker for PPAR α agonists treatments. Also an up-regulation of GstYb4 (Gstm3) gene was detected for the F-344HNsd rats dosed with CPA as shown in Table 3.3. This gene is known to be induced by CPA (Krebs et al., 1998). Significantly down-regulated genes for the samples treated with ciprofibrate for 24 hours are listed in Table 3.4, and the samples treated with CPA for 24 hours are listed in Table 3.5. The cut-off for choosing the fold change depended on the number of genes in each group. From this we can say that the microarray method and the analysis used to achieve these results were all reliable and reproducible, and that we could proceed with the analysis of the early hour samples at 1, 3 and 5 hours treated with ciprofibrate and CPA.

Table 3.2 Known up-regulated genes at 24 hour in ciprofibrate dosed Fisher rats. The list of up-regulated genes for male F-344NHsd rats aged 14-15 weeks dosed with 50 mg kg⁻¹ ciprofibrate then killed after 24 hours. Microarray analysis was carried out as in Section 2.2.5.

Description	Name	Mean fold change	FDR p-value
Mus musculus 3-ketoacyl-CoA thiolase B	MGC29978	3.51	0.000001
Cytochrome P450, family 4, subfamily a, polypeptide 1	Cyp4A1 (CYP4A10)	2.97	0.00001
Cytosolic thioesterase 1	Cte1	2.39	0.000001
Member RAS oncogene family	Rab19	2.26	0.00001
Solute carrier family 9 (sodium/hydrogen exchanger), member 2	Slc9a2	1.78	0.00001
Tensin 1	E030037J05Rik	1.70	0.000001
Mitochondrial ribosomal protein S18A	18S	1.62	0.00001
Lactate dehydrogenase C	Ldh3	1.61	0.00001
NMDA receptor regulated 1	Narg1	1.53	0.0001
RIKEN cDNA 5430402E10 gene	5430402E10Rik	1.46	0.00001
Polymerase (RNA) II (DNA directed) polypeptide A	Polr2a	1.46	0.00001
Succinate-CoA ligase, alpha subunit	Suclg1	1.41	0.0001
	LOC14433	1.39	0.00011
Talin	Tln	1.37	0.0001
RIKEN cDNA 2300009N04 gene	2300009N04Rik	1.33	0.0001
Heat shock protein 90, alpha (cytosolic),	Hspca	1.28	0.000001
UDP glucuronosyltransferase 1 family, polypeptide A5	Ugt1A5	1.27	0.000001
Adenosine A2B receptor	Adora2b	1.26	0.000001
Putative homeodomain transcription factor 1	Phtf1	1.21	0.00001

Table 3.3 Known up-regulated genes at 24 hour in CPA dosed Fisher rats. The list of up-regulated genes for male F-344NHsd rats aged 14-15 weeks dosed with 100 mg kg⁻¹ CPA then killed after 24 hours. Microarray analysis was carried out as in Section 2.2.5

Description	Name	Mean fold change	FDR p-value
Glutathione S-transferase Alpha 1	Gsta1	2.01	0.0001
Heat Shock Protein A8	Hspa8	1.88	0.0001
Glutathione S-transferase Mu 3	GstYb4 (Gstm3)	1.87	0.0001
Udp Glucuronosyltransferase 1 Family, Polypeptide A5	Ugt1a5	1.67	0.0001
Major Urinary Protein 3	Mup3	1.47	0.0001
Eukaryotic Translation Elongation Factor 1 Alpha 1	Eef1a1	1.42	0.0001
Ribosomal Protein S6	Rps6	1.27	0.0001
Metallothionein 1a	Mt1	1.26	0.0001
Ubiquitin C	Ubc	1.18	0.00001
Apolipoprotein A-i	Apoa1	1.16	0.0001

Table 3.4 Known down- regulated genes at 24 hour in ciprofibrate dosed Fisher rats. The list of the down-regulated genes for male F-344NHsd rats aged 14-15 weeks dosed with 50 mg kg⁻¹ ciprofibrate, then killed after 24 hours. Microarray analysis was determined as in Section 2.2.5.

Description	Name	Mean fold change	FDR p-value
Major urinary protein 2	Mup2	-2.47	0.0001
Zinc finger protein 598	Zfp598	-1.85	0.002
Fibrinogen gamma chain	Fgg	-1.77	0.0001
Murinoglobulin 2	Mug2	-1.54	0.0001
Cytochrome P450, family 2, subfamily c, polypeptide 29	CYP2c29	-1.44	0.0001
Polymerase (RNA) II (DNA directed) polypeptide A	Polr2a	-1.38	0.0001
Brain protein 44-like	Brp44l	-1.27	0.0001
Cysteine dioxygenase, type I	Cdo1	-1.23	0.0001
Ornithine carbamoyltransferase	Otc	-1.21	0.0001
	Sepina3a	-1.21	0.0001

Table 3.5 Known down- regulated genes at 24 hour in CPA dosed Fisher rats. The list of the down-regulated genes for male F-344NHsd rats aged 14-15 weeks dosed with 100 mg kg⁻¹ CPA, then killed after 24 hours. Microarray analysis was determined as in Section 2.2.5.

Description	Name	Mean fold change	FDR p-value
RIKEN cDNA 2300009N04 gene	2300009N04Rik	-1.52	0.00001
Major urinary protein 2	Mup2	-1.29	0.00001
Ribonuclease, RNase A family 4	Rnase4	-1.27	0.00001
Transferrin	Trf	-1.19	0.00001
Cytochrome P450, family 2, subfamily d, polypeptide 26	CYP2d26	-1.14	0.00001

Figure 3.39 shows a heatmap depicting the hepatic genes that were affected by the ciprofibrate after 24 hour treatment: red indicates highly- induced, and blue indicates down- regulation of the genes. These were established with High-Throughput GoMiner (Zeeberg, 2005)). Also the Figure shows the different pathways that are changed with the ciprofibrate dose after 24 hours. The main changed pathways are the fatty acid metabolic process where 5 genes from 14 total genes involved in this process were over expressed (35.7%), and the monocarboxylic acid metabolic process with 5 out of 19 genes (26.3%), also, *fatty acid β oxidation* and *acyl-CoA metabolic process* had both 50% up- regulation.

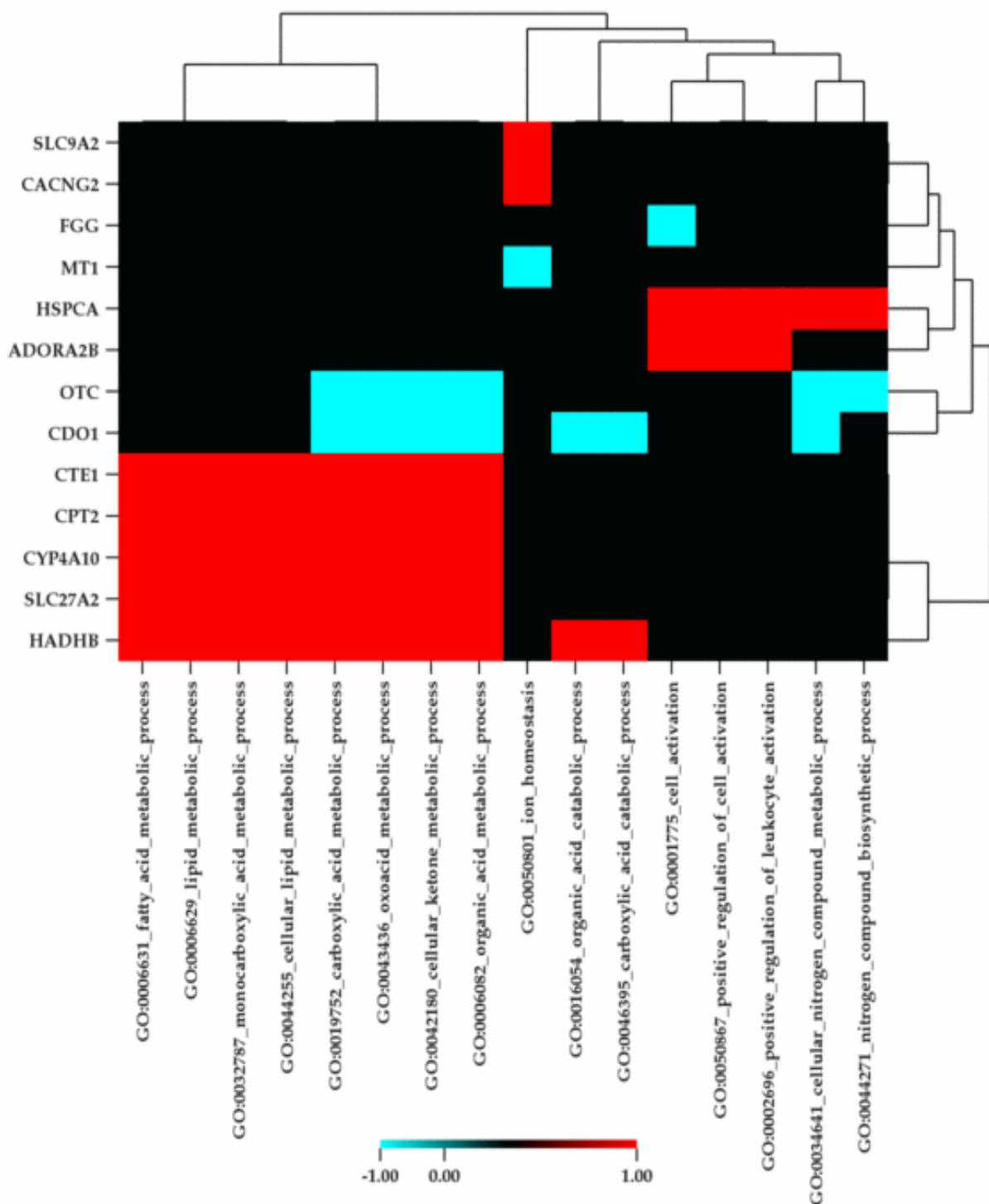


Figure 3.39 Effect of ciprofibrate on gene expression after 24 hour. Male F-344NHsd rats aged 14-15 weeks were dosed with 50 mg kg⁻¹ ciprofibrate, then killed after 24 hours. Microarray analysis was carried out as in Section 2.2.5, this heatmap was established with High-Throughput GoMiner (Zeeberg, 2005). Red demonstrates up-regulated, and blue demonstrates down-regulated (-1 down-regulated, +1 up-regulated).

In Figure 3.40 is a heatmap showing the affected hepatic genes up/down- regulated with the CPA after 24 hours: red describes highly induced, and blue demonstrates down- regulated induction of the genes. The Figure shows the different pathways that are changed with the CPA dose after 24 hours. Nevertheless, the highly changed processes are the glutathione transferase activity where 2 of 3 genes are over expressed (66.7%), the other process is related to the inter-phase of mitotic cell cycle which is also 2 out of 3 over expressed genes (66.7%).

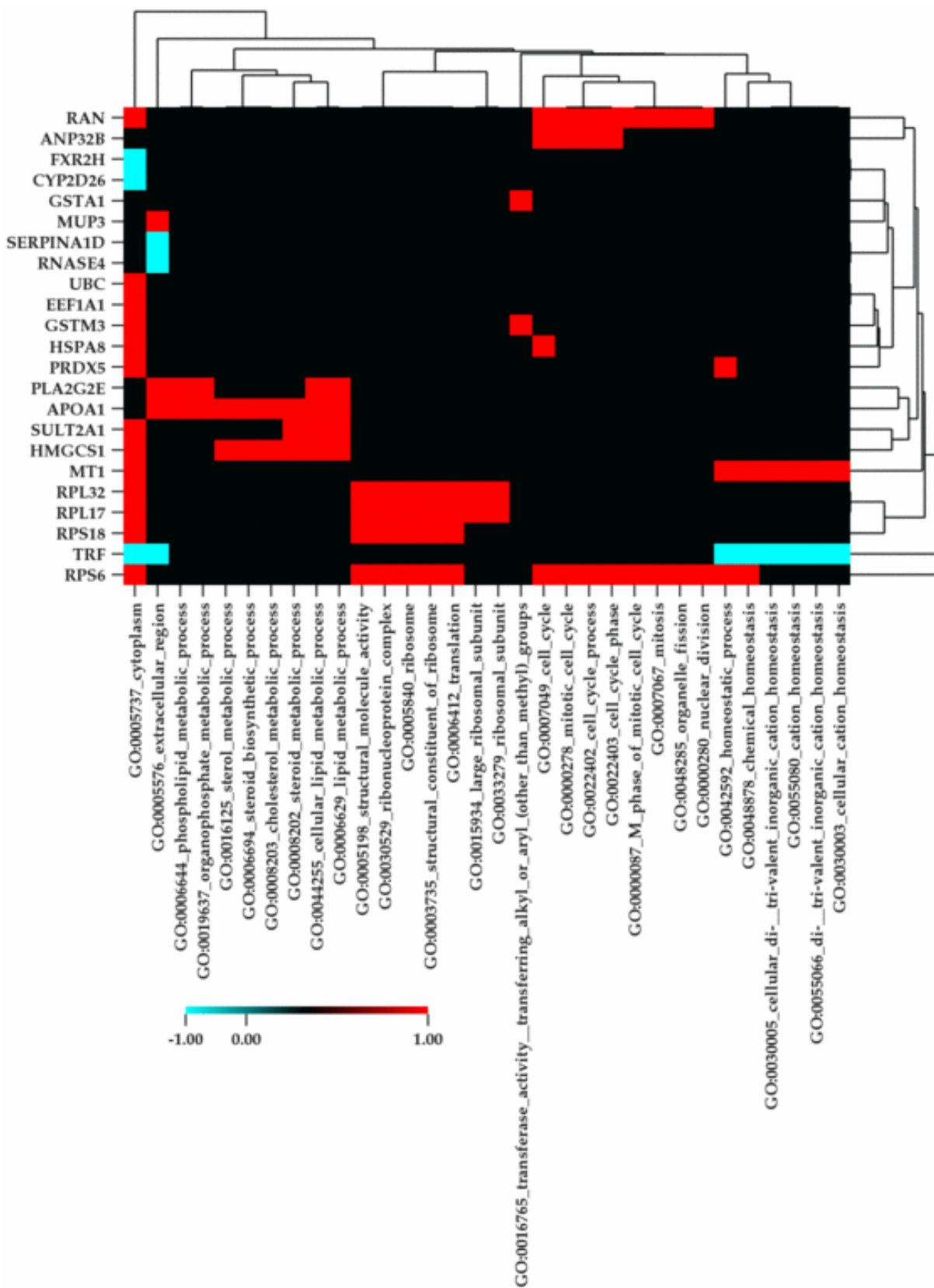


Figure 3.40 Effect of CPA after 24 hour on gene expression. Male F-344NHsd rats aged 14-15 weeks dosed with 100 mg kg⁻¹ CPA, then killed after 24 hours. Microarray analysis was carried out as in Section 2.2.5, this heatmap was established with High-Throughput GoMiner (Zeeberg, 2005). Red demonstrates up- regulated, and blue demonstrates down- regulated (-1 down- regulated, +1 up- regulated).

Section 3.3.5 Microarray analysis of ciprofibrate dosed rats after 1, 3, and 5 hours

Having performed preliminary analysis at 24 hours (Section 3.3.4). The microarray analysis of the early hour samples was undertaken. This was done collectively at 1, 3 and 5 hours for the hepatic samples from F-344NHsd fisher rats dosed with 50 mg kg⁻¹ ciprofibrate, 100 mg kg⁻¹ CPA, or corn oil (Section 2.2.5).

The analysis of the 1, 3 and 5 hour groups with R software revealed 1377 significant genes for each of the groups (FDR p-value <0.05) out of 5826 genes, (the whole list is attached in a CD at the end of the Thesis). A list of the up- regulated genes for the samples treated with ciprofibrate for 1 hour with a fold induction of >2.6 fold change is presented in Table 3.6 which include *Igf1*, *Sstr4*, *Abcc2*, *mup1*, *mmu* and *Adamts*, and the down- regulated genes after 1 hour is found in Table 3.7, while a list of up and down- regulated genes for 3 hour samples are shown in Table 3.8 (fold change >1.7) and Table 3.9 respectively which include, a number of the up- regulated for 3 hours are *Seprina1b*, *mup4*, *Mug2*, *Gsta1* and *Atp8b3*. A list of the genes up and down- regulated with ciprofibrate after 5 hours are listed in Table 3.10 (fold change >2.0) and Table 3.11 respectively, the up- regulated genes include *Seprina1b*, *Mug1*, *Sema5a*, *My16*, *Mup4*, and *Mug4*. The different cut-off for the fold changes depends on the number of genes in each of the groups.

Table 3.6 Up- regulated genes at 1 hour in ciprofibrate dosed Fisher rats. The list of up- regulated genes for male F-344NHsd rats aged 14-15 weeks dosed with 50 mg kg⁻¹ ciprofibrate then killed after 1 hours. Microarray analysis was carried out as in Section 2.2.5.

Description	Name	Mean fold change	FDR p-value
FXYP domain-containing ion transport regulator 2	Fxyd2	3.86	0.0001
Insulin-like growth factor 1	Igf1	3.39	0.005
Olfactory receptor 976	Olf976	3.37	0.0001
Serine/threonine kinase 3	Stk3	3.29	0.0001
N-myc downstream regulated gene 1	Ndr91	3.19	0.0001
Somatostatin receptor 4	Sstr4	3.18	0.003
Adaptor-related protein complex 1, sigma 1 subunit	Ap1s1	3.13	0.0001
Thymidylate synthetase	Tyms	3.09	0.0001
Carboxypeptidase A5	Cpa5	3.08	0.001
Zinc finger protein 64	Zfp64	3.07	0.006
Complement factor H	Cfh	3.06	0.002
Zinc finger protein 180	Zfp180	2.99	0.0001
PDZ domain containing 3	Pdzk2	2.92	0.003
DNA segment, Chr 11, Brigham & Women's Genetics 0517 expressed	D11Bwg0517e	2.91	0.0001
CD68 antigen	Cd68	2.90	0.0001
Sema domain, seven thrombospondin repeats, transmembrane domain and short cytoplasmic domain 5A	Sema5a	2.89	0.00011
Serum amyloid A 3	Saa3	2.89	0.0001
Chloride intracellular channel 1	Clic1	2.86	0.008
Dihydropyrimidinase-like 2	Dpysl2	2.86	0.008
Proteolipid protein 1	Plp1	2.85	0.0001
Otoancorin	Otoa	2.82	0.0001
Kinesin family member 9	Kif9	2.82	0.0001
Deoxynucleotidyltransferase, terminal	Dntt	2.79	0.0001

Table 3.6 Up- regulated genes at 1 hour in ciprofibrate dosed Fisher rats. The list of up- regulated genes for male F-344NHsd rats aged 14-15 weeks dosed with 50 mg kg⁻¹ ciprofibrate then killed after 1 hours. Microarray analysis was carried out as in Section 2.2.5.

Description	Name	Mean fold change	FDR p-value
Transmembrane protein 140	1110007F12Rik	2.77	0.0001
Telomeric repeat binding factor 2	Terf2	2.77	0.003
PWWP domain containing 2A	mmu	2.76	0.001
Chemokine-like receptor 1	Cmklr1	2.74	0.008
Galactosyltransferase I	B4galt7	2.73	0.0001
Phosphoinositide binding specific) member 3	Plekha3	2.73	0.0001
Ganglioside-induced differentiation-associated protein 1-like 1	Gdap111	2.73	0.001
BTB (POZ) domain containing 9	Btbd9	2.72	0.0001
ATP-binding cassette, sub-family C member 2	Abcc2	2.72	0.0001
Major urinary protein 1	Mup1	2.71	0.001
ADAM metallopeptidase with thrombospondin type 1 motif, 2	Adamts2	2.71	0.0001
Mucin 13, cell surface associated	Ly64	2.70	0.003
Endoplasmic reticulum protein 44	Txndc4	2.70	0.005

Table 3.7 Down-down- reg regulated genes at 1 hour in ciprofibrate dosed Fisher rats. The list of down-regulated genes for male F-344NHsd rats aged 14-15 weeks dosed with 50 mg kg⁻¹ ciprofibrate then killed after 1 hours. Microarray analysis was carried out as in Section 2.2.5.

Description	Name	Mean fold change	FDR p-value
Fibroblast growth factor 21	Fgf21	-6.62	0.0001
Regulator of G-protein signalling like 2	Rgs12	-5.89	0.0001
Ubiquitin specific peptidase 28	Usp28	-5.74	0.0001
Chloride channel 6	Clcn6	-5.48	0.0001
Aspartyl-tRNA synthetase	Dars	-5.28	0.0001
Zinc finger, DHHC-type containing 9	Zdhhc9	-5.20	0.0001
ArfGAP with RhoGAP domain, ankyrin repeat and PH domain 1	Centd2	-4.99	0.0001
Origin recognition complex, subunit 6 like	Orc6l	-4.36	0.0001
Vanin 3	Vnn3	-4.03	0.0001
Zinc finger protein 212	Zfp212	-4.03	0.0001
Poliovirus receptor-related 4	Pvr14	-3.98	0.0001
Frizzled homolog 7	Fzd7	-3.93	0.00011
Transforming growth factor, beta 1	Tgfb1	-3.90	0.0001
Proprotein convertase subtilisin/kexin type 4	Pcsk4	-3.85	0.022
UDP glucuronosyltransferase 1 family, polypeptide A6	Ugt1a6	-3.84	0.0001
Rho GDP dissociation inhibitor (GDI) alpha	Arhgdia	-3.83	0.0001
PRP4 pre-mRNA processing factor 4 homolog	Prpf4	-3.75	0.0001
Immunoglobulin superfamily, DCC subclass, member 3	Punc	-3.65	0.0001
MAP/microtubule affinity-regulating kinase 2	Mark2	-3.65	0.012
WD repeat and HMG-box DNA binding protein 1	Wdhd1	-3.60	0.0001
Tubulin folding cofactor B	Ckap1	-3.32	0.0001
Keratinocyte associated protein 2	Krtcap2	-3.29	0.0001
Homeobox A3	Hoxa3	-3.22	0.0001

Table 3.7 Down-down- reg regulated genes at 1 hour in ciprofibrate dosed Fisher rats. The list of down-regulated genes for male F-344NHsd rats aged 14-15 weeks dosed with 50 mg kg⁻¹ ciprofibrate then killed after 1 hours. Microarray analysis was carried out as in Section 2.2.5.

Description	Name	Mean fold change	FDR p-value
Cystathionase (cystathionine gamma-lyase)	Cth	-3.09	0.0001
ASF1 anti-silencing function 1 homolog B (S. cerevisiae)	Asf1b	-3.02	0.001
E2F transcription factor 2	E2f2	-2.84	0.002
Rho-related BTB domain containing 2	Rhobtb2	-2.82	0.004
Uracil DNA glycosylase	Ung	-2.75	0.001
CD97 antigen	Cd97	-2.74	0.0001
Pro-platelet basic protein	Cxcl7	-2.72	0.0001
Mannosyl (alpha-1,6-)-glycoprotein beta-1,2-N-acetylglucosaminyltransferase	GnT	-2.69	0.005
Glucosamine-6-phosphate deaminase 2	Gnpda2	-2.69	0.019
Jagged 2	Jag2	-2.64	0.016
Immunoglobulin kappa chain variable 28	IGKV2	-2.64	0.008
HIV-1 Rev binding protein 2	Hrb2	-2.55	0.003
Antigen identified by monoclonal antibody Ki-67	Mki67	-2.55	0.0001
Vomer nasal 1 receptor, H3	V1rh3	-2.47	0.001
Ubiquitin-like modifier activating enzyme 1	Ube1x	-2.34	0.0001
Formiminotransferase cyclodeaminase	Ftcd	-2.30	0.0001
Phenylalanyl-tRNA synthetase 2, mitochondrial	Fars1	-2.26	0.001
Zinc finger protein 364	Zfp364	-2.24	0.0001
RAR-related orphan receptor B	Rorb	-2.20	0.0001
TNFAIP3 interacting protein 1	Tnip1	-2.17	0.035
Cas-Br-M (murine) ecotropic retroviral transforming sequence b	Cblb	-2.16	0.001
Splicing factor, arginine/serine-rich 9	Sfrs9	-2.12	0.0001
Papillary renal cell carcinoma	Prcc	-2.04	0.007
Receptor-interacting serine-threonine kinase 2	Ripk2	-2.04	0.003

Table 3.7 Down-down- reg regulated genes at 1 hour in ciprofibrate dosed Fisher rats. The list of down-regulated genes for male F-344NHsd rats aged 14-15 weeks dosed with 50 mg kg⁻¹ ciprofibrate then killed after 1 hours. Microarray analysis was carried out as in Section 2.2.5.

Description	Name	Mean fold change	FDR p-value
Phosphate regulating endopeptidase homolog, X-linked	Phex	-2.02	0.021
Xylosyltransferase II	Xylt2	-2.02	0.045
Pumilio homolog 1	Pum1	-2.01	0.001
Phosphodiesterase 4D interacting protein (myomegalin)	Usmg4	-2.01	0.003

Table 3.8 Up- regulated genes after 3 hours in ciprofibrate dosed Fisher rats. The list of up- regulated genes for male F-344NHsd rats aged 14-15 weeks dosed with 50 mg kg⁻¹ ciprofibrate then killed after 3 hours. Microarray analysis was carried out as in Section 2.2.5.

Description	Name	Mean fold change	FDR p-value
Proteasome (prosome, macropain) subunit, beta type 9	Psmb9	3.65	0.0001
Inositol polyphosphate phosphatase-like 1	Inpp1l	3.07	0.001
Opioid growth factor receptor	Ogfr	2.95	0.0001
Serum amyloid A 3	Saa3	2.71	0.0001
Transmembrane protein 222	D4Ertd196e	2.47	0.0001
Otoancorin	Otoa	2.43	0.0001
Histone cluster 1, H2af	Hist1h2af	2.40	0.0001
Hemopexin	Hpxn	2.24	0.005
Olfactory receptor 1395	Olfr1395	2.23	0.000
Serine/threonine kinase 25	Stk25	2.17	0.003
UDP-GlcNAc:betaGal beta-1,3-N-acetylglucosaminyltransferase 7	B3gnt7	2.17	0.0001
Iroquois homeobox 1	Irx1	2.06	0.001
WW domain binding protein 1	Wbp1	2.05	0.002
Serine (or cysteine) preptidase inhibitor, clade A, member 1B	Serpina1b	2.05	0.001
Major urinary protein 4	Mup4	2.03	0.006
Insulin 1	Ins1	2.02	0.000
Group specific component	Gc	2.02	0.000
FLYWCH-type zinc finger 1	E030034P13Rik	2.01	0.000
Extracellular matrix component; may play a role in fibrosis and tumour metastasis	Fn1	1.96	0.000
Testis expressed 264	Tex264	1.92	0.005
Metallothionein 1, pseudogene 1	Mt1	1.90	0.001
DAZ associated protein 1	Dazap1	1.89	0.000

Table 3.8 Up-regulated genes after 3 hours in ciprofibrate dosed Fisher rats. The list of up-regulated genes for male F-344NHsd rats aged 14-15 weeks dosed with 50 mg kg⁻¹ ciprofibrate then killed after 3 hours. Microarray analysis was carried out as in Section 2.2.5.

Description	Name	Mean fold change	FDR p-value
DNA (cytosine-5-)-methyltransferase 3-like	Dnmt3l	1.87	0.007
Serpin peptidase inhibitor, clade A 1	Serpina1	1.87	0.001
Ventral anterior homeobox 2	Vax2	1.87	0.0001
Receptor (G protein-coupled) activity modifying protein 2	Ramp2	1.83	0.0001
Deiodinase, iodothyronine, type III	Dio3	1.81	0.006
Coiled-coil domain containing 146	4930528G09Rik	1.80	0.004
Ribosomal protein S17	Rps17	1.78	0.001
Cathepsin Z	Ctsz	1.76	0.0001
Glutathione S-transferase alpha 1	Gsta1	1.75	0.0001
Murinoglobulin 2	Mug2	1.75	0.0001
ATPase, Class I, type 8B, member 3	Atp8b3	1.74	0.0001
RIKEN cDNA 1700048F04 gene	1700048F04Rik	1.74	0.002
Interferon, alpha-inducible protein 27 like 1	D12Ert647e	1.74	0.004
Coiled-coil domain containing 89	1700019B01Rik	1.72	0.001
Synaptogyrin 4	Syng4	1.70	0.0001

Table 3.9 Down-regulated genes after 3 hours in ciprofibrate dosed Fisher rats. The list of down-regulated genes for male F-344NHsd rats aged 14-15 weeks dosed with 50 mg kg⁻¹ ciprofibrate then killed after 3 hours. Microarray analysis was carried out as in Section 2.2.5.

Discription	Name	Mean fold change	FDR p-value
6-phosphofructo-2-kinase/fructose-2,6-biphosphatase 2	Pfkfb2	-3.55	0.0001
Intraflagellar transport 122 homolog	Wdr10	-2.97	0.0001
Cystathionase (cystathionine gamma-lyase)	Cth	-2.62	0.0001
KH domain containing, RNA binding, signal transduction associated 2	Khdrbs2	-2.45	0.0001
C-type lectin domain family 4, member e	Clecsf9	-2.39	0.011
Casein kinase 2, alpha 1 polypeptide	Csnk2a1	-2.37	0.025
Transportin 1	Tnpo1	-2.04	0.002
Potassium voltage-gated channel, KQT-like subfamily, member 2	Kcnq2	-2.02	0.008

Table 3.10 Up- regulated genes after 5 hours in ciprofibrate dosed Fisher rats. The list of up- regulated genes for male F-344NHsd rats aged 14-15 weeks dosed with 50 mg kg⁻¹ ciprofibrate then killed after 5 hours. Microarray analysis was carried out as in Section 2.2.5.

Description	Name	Mean fold change	FDR p-value
Group specific component	Gc	3.64	0.0001
Apolipoprotein B	Apob	2.92	0.0001
Dual specificity phosphatase 14	Dusp14	2.88	0.0001
Mucin 2, oligomeric mucus/gel-forming	Muc2	2.88	0.0001
Glucosidase, alpha; neutral C	5830445O15Rik	2.86	0.004
Serine (or cysteine) preptidase inhibitor, clade A, member 1B	Serpina1b	2.81	0.001
Kininogen 1	Kng1	2.79	0.0001
Murinoglobulin 1	Mug1	2.76	0.0001
Hemopexin	Hpxn	2.57	0.005
Metallothionein 1	Mt1	2.55	0.001
Microtubule-associated protein 7 domain containing 1	BC019977	2.54	0.0001
Potassium voltage-gated channel, subfamily H, member 1	Kcnh1	2.48	0.001
Hyaluronan and proteoglycan link protein 4	Hapln4	2.42	0.0001
Per-hexamer repeat gene 5	Phxr5	2.41	0.003
Peptidylprolyl isomerase A (cyclophilin A)	Ppia	2.41	0.0001
Sortilin-related VPS10 domain containing receptor 2	Sorcs2	2.40	0.002
Serine (or cysteine) proteinase inhibitor, clade A, member 3M	Serpina3m	2.39	0.001
Gulonolactone (L-) oxidase	Gulo	2.39	0.0001
Transmembrane protein 41a	5730578N08Rik	2.37	0.001
RIKEN cDNA 1700001G17 gene	1700001G17Rik	2.32	0.0001
RIKEN cDNA 4930543D07 gene	4930543D07Rik	2.31	0.005

Table 3.10 Up- regulated genes after 5 hours in ciprofibrate dosed Fisher rats. The list of up- regulated genes for male F-344NHsd rats aged 14-15 weeks dosed with 50 mg kg⁻¹ ciprofibrate then killed after 5 hours. Microarray analysis was carried out as in Section 2.2.5.

Description	Name	Mean fold change	FDR p-value
Sema domain, seven thrombospondin repeats transmembrane domain 5B	Sema5b	2.31	0.004
Sema domain, seven thrombospondin repeats transmembrane domain 5A	Sema5a	2.29	0.0001
Fructose-1,6-bisphosphatase 1	Fbp1	2.27	0.0001
Myosin, light chain 6	Myl6	2.27	0.001
Collagen, type XI, alpha 2	Col11a2	2.26	0.005
Major urinary protein 4	Mup4	2.26	0.006
Human immunodeficiency virus type I enhancer binding protein 1	Hivep1	2.26	0.0001
Serum amyloid P-component	Apcs	2.24	0.0001
B9 protein domain 1	Eppb9	2.22	0.0001
Ankyrin repeat domain 40	Gcap15	2.22	0.00011
Propionyl-coenzyme A carboxylase, alpha polypeptide	Pcca	2.20	0.001
Murinoglobulin 2	Mug2	2.17	0.0001
Metallothionein 4	Mt4	2.17	0.009
Ubiquitin A-52 residue ribosomal protein fusion product 1	Uba52	2.16	0.002
Ceruloplasmin	Cp	2.16	0.0001
Prostaglandin E synthase 2	Ptges2	2.16	0.0001
Fetuin B	Fetub	2.16	0.001
Glutathione transferase zeta 1	Gstz1	2.13	0.0001
RIKEN cDNA 4930444A02 gene	4930444A02Rik	2.13	0.0001
Vitamin K epoxide reductase complex, subunit 1	Vkorc1	2.12	0.006
RIKEN cDNA 2010107G23 gene	2010107G23Rik	2.11	0.007
Olfactory receptor 123	Olf123	2.11	0.003

Table 3.11 Down-regulated genes after 5 hours in ciprofibrate dosed Fisher rats. The list of down-regulated genes for male F-344NHsd rats aged 14-15 weeks dosed with 50 mg kg⁻¹ ciprofibrate then killed after 5 hours. Microarray analysis was carried out as in Section 2.2.5.

Description	Name	Mean fold change	FDR p-value
NmrA-like family domain containing 1	1110025F24Rik	-4.91	0.00001
Neural precursor cell expressed, developmentally down-regulated 4	Nedd4	-4.49	0.00001
Zinc finger, MIZ-type containing 1	BC065120	-4.31	0.00001
RAS (RAD and GEM)-like GTP-binding 1	Rem1	-3.79	0.00001
Adenylate cyclase 8 (brain)	Adcy8	-3.57	0.00001
Taste receptor, type 2, member 116	Tas2r116	-3.50	0.026
Phosphofructokinase, platelet	Pfcp	-3.41	0.0001
Rab40c, member RAS oncogene family	Rab40c	-3.35	0.027
Mucin 5AC, oligomeric mucus/gel-forming	Muc5ac	-3.19	0.0001
Tubulin folding cofactor B	Ckap1	-3.16	0.0001
Cas-Br-M (murine) ecotropic retroviral transforming sequence b	Cblb	-3.10	0.005
Acyl-Coenzyme A dehydrogenase, short/branched chain	Acadsb	-2.93	0.0001
Cyclin B3	Ccnb3	-2.89	0.00001
RAN binding protein 1	Ranbp1	-2.89	0.004
RAB interacting factor	Rabif	-2.87	0.001
Usher syndrome 1C binding protein 1	Ushbp1	-2.81	0.0001
Zinc finger protein 212	Zfp212	-2.70	0.0001
Diazepam binding inhibitor-like 5	Dbil5	-2.70	0.030
Immunoglobulin lambda chain complex	Igl	-2.68	0.00011
Minichromosome maintenance complex component 5	Mcm5	-2.55	0.0001
TPX2, microtubule-associated, homolog	Tpx2	-2.50	0.004
Olfactory receptor 609	Olf609	-2.46	0.026
Casein kinase 2, alpha 1 polypeptide	Csnk2a1	-2.43	0.025

Table 3.11 Down- regulated genes after 5 hours in ciprofibrate dosed Fisher rats. The list of down-regulated genes for male F-344NHsd rats aged 14-15 weeks dosed with 50 mg kg⁻¹ ciprofibrate then killed after 5 hours. Microarray analysis was carried out as in Section 2.2.5.

Description	Name	Mean fold change	FDR p-value
Sonic hedgehog	Shh	-2.38	0.005
Transglutaminase 5	Tgm5	-2.25	0.012
Nuclear transcription factor-Y alpha	Nfya	-2.15	0.00001
Poly (ADP-ribose) polymerase family, member 12	Zc3hdc1	-2.13	0.036

Figure 3.41 shows a heatmap of the hepatic genes that were highly induced with ciprofibrate after 5 hours. In addition, the trees at the top and right show how these genes and pathways cluster together. Apparently there was not enough data to be able to generate figures for the other time points (as indicated by the software). Genes of the acute phase response were found to be up-regulated in both the 3 and 5 hour treated groups. For the 3 hour group there was 4 out of 23 genes responsible for the acute phase response (17.4%) and in the 5 hour group there was 8 out of 23 genes (34.8%).

This also was the case for the genes of the acute inflammatory response, where for the 3 hour group there was 4 out of 43 genes and for the 5 hours there was 9 out of 43 genes (20.9%). As for the genes of the response to wounding in the 3 hour samples there was 7 out of 193 (3.6%) and 19 out of 193 genes (9.8%) for the 5 hour samples.

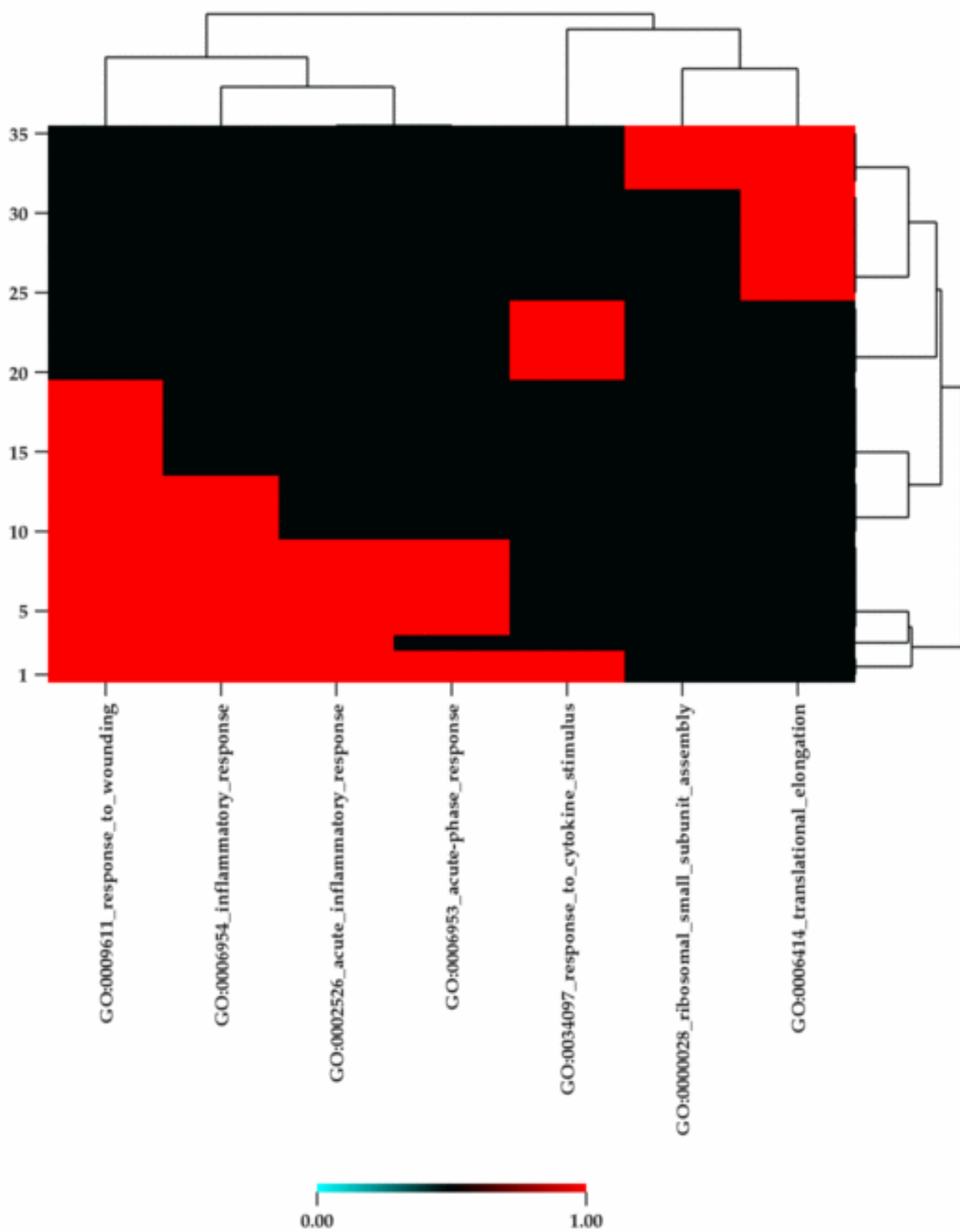


Figure 3.41 Heatmap for microarray analysis of ciprofibrate gene expression after 5 hours.

Male F-344/NHsd rats aged 14-15 weeks dosed with 50 mg kg⁻¹ ciprofibrate, then killed after 24 hours. The numbers at the left side refer to genes 1= SERPINA1A, 5= HP, 10= SERPINC1, 15= UCN2, 20= STATG, 25= TUBB4, 30=CDC20, 35=RPS27. Microarray analysis was carried out as in Section 2.2.5, this heatmap was established with High-Throughput GoMiner (Zeeberg, 2005). Red demonstrates up-regulated, and blue demonstrates down-regulated (-1 down-regulated, +1 up-regulated).

Section 3.3.6 Microarray analysis of CPA dosed rats after 1, 3, and 5 hours

Microarray analysis was performed with hepatic samples from F-344NHsd fisher rats dosed with 100 mg kg⁻¹ CPA (Section 2.2.5). A list of the up-regulated genes for the samples treated with CPA for 1 hour with a fold induction of >1.2 fold change is presented in Table 3.12 including *Cd97*, *Sstr4* and *Rnfl41*. A list of down-regulated genes after 1 hour is presented in Table 3.13, while a list of up-regulated genes for 3 hour samples were found in Table 3.14 (fold change >1.0) including *Hpxn*, *Cyp2c29*, and *Seprina3m* and Table 3.15 lists the down-regulated genes after 3 hours CPA treatment. The genes up-regulated with ciprofibrate after 5 hours were listed in Table 3.16 (fold change >1.5) including *Scd1*, *Mug2*, *Ppp1cb* and *Hspd1*, and the down-regulated genes were listed in Table 3.17.

Table 3.12 Up- regulated genes at 1 hour in CPA dosed Fisher rats. A list of up- regulated genes for male F-344NHsd rats aged 14-15 weeks dosed with 100 mg kg⁻¹ CPA then killed after 1 hours. Microarray analysis was determined as in Section 2.2.5.

Description	Name	Mean fold change	FDR p-value
CD97 molecule	Cd97	2.50	0.0001
Lix1-like	D130027M04Rik	1.47	0.005
Somatostatin receptor 4	Sstr4	1.47	0.003
Transient receptor potential cation channel, subfamily C, member 4 associated protein	Trpc4ap	1.29	0.003
E2F transcription factor 3	E2f3	1.86	0.039
Sirtuin (silent mating type information regulation 2 homolog) 2	Sirt2	1.80	0.013
MCF.2 cell line derived transforming sequence-like	Mcf2l	1.22	0.042
Kinesin family member 3B	Kif3b	1.39	0.010
Dual specificity phosphatase 28	0710001B24Rik	1.16	0.002
Neurexin 3	Nrxn3	1.71	0.038
Ring finger protein 141	Rnf141	1.22	0.0001

Table 3.13 Down- regulated genes at 1 hour in CPA dosed Fisher rats. A list of down- regulated genes for male F-344NHsd rats aged 14-15 weeks dosed with 100 mg kg⁻¹ CPA then killed after 1 hours. Microarray analysis was determined as in Section 2.2.5.

Description	Name	Mean fold change	FDR p-value
Phospholipase A2, group IIC	Pla2g2c	-3.09	0.006
RIKEN cDNA 4930507D10 gene	4930507D10Rik	-3.03	0.021
Caspase recruitment domain family, member 10	Card10	-2.62	0.001
Kruppel-like factor 15	Klf15	-2.60	0.002
Protein phosphatase 1, regulatory (inhibitor) subunit 14A	Ppp1r14a	-2.58	0.021
Retinoic acid induced 1	Rai1	-2.38	0.006
RIKEN cDNA 4933433G15 gene	4933433G15Rik	-2.31	0.012
RIKEN cDNA 7030407O06 gene	7030407O06Rik	-2.30	0.015
Hepatocyte nuclear factor 4, alpha	Hnf4a	-2.25	0.021
GS homeobox 1	Gsh1	-2.23	0.0001
RIKEN cDNA 9530085L11 gene	9530085L11Rik	-2.13	0.0001
RIKEN cDNA 1810046J19 gene	1810046J19Rik	-2.05	0.033

Table 3.14 Up- regulated genes after 3 hours in CPA dosed Fisher rats. A list of up- regulated genes for male F-344NHsd rats aged 14-15 weeks dosed with 100 mg kg⁻¹ CPA then killed after 3 hours. Microarray analysis was determined as in Section 2.2.5.

Description	Name	Mean fold change	FDR p-value
RIKEN cDNA E130101E03 gene	E130101E03Rik	1.16	0.003
Hemopexin	Hpxn	1.14	0.005
COP9 constitutive photomorphogenic homolog subunit 4	Cops4	1.30	0.012
Hairy/enhancer-of-split related with YRPW motif 1	Hey1	1.12	0.028
Cytochrome P450, family 2, subfamily c, polypeptide 29	Cyp2c29	1.08	0.039
Serine (or cysteine) proteinase inhibitor, clade A, member 3M	Serpina3m	1.12	0.001

Table 3.15 Down- regulated genes after 3 hours in CPA dosed Fisher rats. A list of down- regulated genes for male F-344NHsd rats aged 14-15 weeks dosed with 100 mg kg⁻¹ CPA then killed after 3 hours. Microarray analysis was determined as in Section 2.2.5

Description	Name	Mean fold change	FDR p-value
Membrane-associated ring finger	Axot	-2.35	0.006
Aminolevulinate, delta-, synthase 1	Alas1	-1.84	0.0001
Cytochrome P450, family 3, subfamily a, polypeptide 44	Cyp3a44	-1.46	0.001
UDP glycosyltransferase 1 family, polypeptide A10	Ugt1a13	-1.13	0.002
ER degradation enhancer, mannosidase alpha-like 1	Edem1	-1.01	0.002
Serum induced transcript 1	Sif1	-1.01	0.020

Table 3.16 Up- regulated genes after 5 hours in CPA dosed Fisher rats. A list of up- regulated genes for male F-344NHsd rats aged 14-15 weeks dosed with 100 mg kg⁻¹ CPA then killed after 5 hours. Microarray analysis was determined as in Section 2.2.5

Description	Name	Mean fold change	FDR p-value
Cathepsin E	C920004C08Rik	3.48	0.0001
Metallothionein	mt	2.87	0.0001
Murinoglobulin 1	Mug1	2.47	0.0001
Stearoyl-Coenzyme A desaturase 1	Scd1	2.21	0.0001
ADAM metallopeptidase with thrombospondin type 1 motif, 1	Adamts1	2.17	0.001
Group specific component	Gc	2.18	0.0001
RIKEN cDNA 2810455D13 gene	2810455D13Rik	2.12	0.003
Frizzled homolog 7	Fzd7	2.10	0.0001
Murinoglobulin 2	Mug2	2.09	0.0001
Zinc finger protein 335	1810045J01Rik	2.02	0.006
Beta-2 microglobulin	B2m	2.01	0.004
cDNA sequence BC018242	BC018242	1.95	0.006
RIKEN cDNA 2600006L11 gene	2600006L11Rik	1.91	0.0001
Protein phosphatase 1, catalytic subunit, beta isoform	Ppp1cb	1.91	0.002
Predicted gene	IGKV12	1.87	0.0001
Transmembrane BAX inhibitor motif containing 6	Tegt	1.81	0.003
C-type lectin domain family 5, member a	Clecsf5	1.80	0.0001
SET binding protein 1	Setbp1	1.66	0.005
RIKEN cDNA 4930538K18 gene	4930538K18Rik	1.64	0.0001
Ribosomal protein L5	Rpl5	1.63	0.006
Sulfotransferase family 2A, member 2	Sult2a2	1.61	0.008
Pregnancy upregulated non-ubiquitously expressed CaM kinase	Pnck	1.59	0.0001
Heat shock protein 1	Hspd1	1.58	0.005

Table 3.17 Down- regulated genes after 5 hours in CPA dosed Fisher rats. A list of down- regulated genes for male F-344NHsd rats aged 14-15 weeks dosed with 100 mg kg⁻¹ CPA then killed after 5 hours. Microarray analysis was determined as in Section 2.2.5

Description	Name	Mean fold change	FDR p-value
Acyl-Coenzyme A dehydrogenase, short/branched chain	Acadsb	-3.56	0.0001
Diazepam binding inhibitor-like 5	Dbil5	-2.76	0.030
Splicing factor, arginine/serine-rich 16	Sfrs16	-2.53	0.007
Inositol polyphosphate phosphatase-like 1	Inpp1l	-2.50	0.001
ArfGAP with dual PH domains 1	Centa1	-2.43	0.027
RIKEN cDNA 1700034B16 gene	1700034B16Rik	-2.37	0.001
Calcineurin-like phosphoesterase domain containing 1	C530044N13Rik	-2.23	0.033
RIKEN cDNA 1700126L10 gene	1700126L10Rik	-2.22	0.006
Mitochondrial ribosomal protein S18A	18S	-2.19	0.0001
RIKEN cDNA 2300009N04 gene	2300009N04Rik	-2.16	0.0001

Figure 3.42 shows a heatmap of the hepatic genes that were highly induced with CPA after 3 hours. In addition, the trees at the top and right show how these genes and pathways cluster together. There was not enough data to be able to generate figures for the other time points, or for the down- regulated genes, and there was not any related pathways that were repeated with the 1, 3 and 5 hour time points.

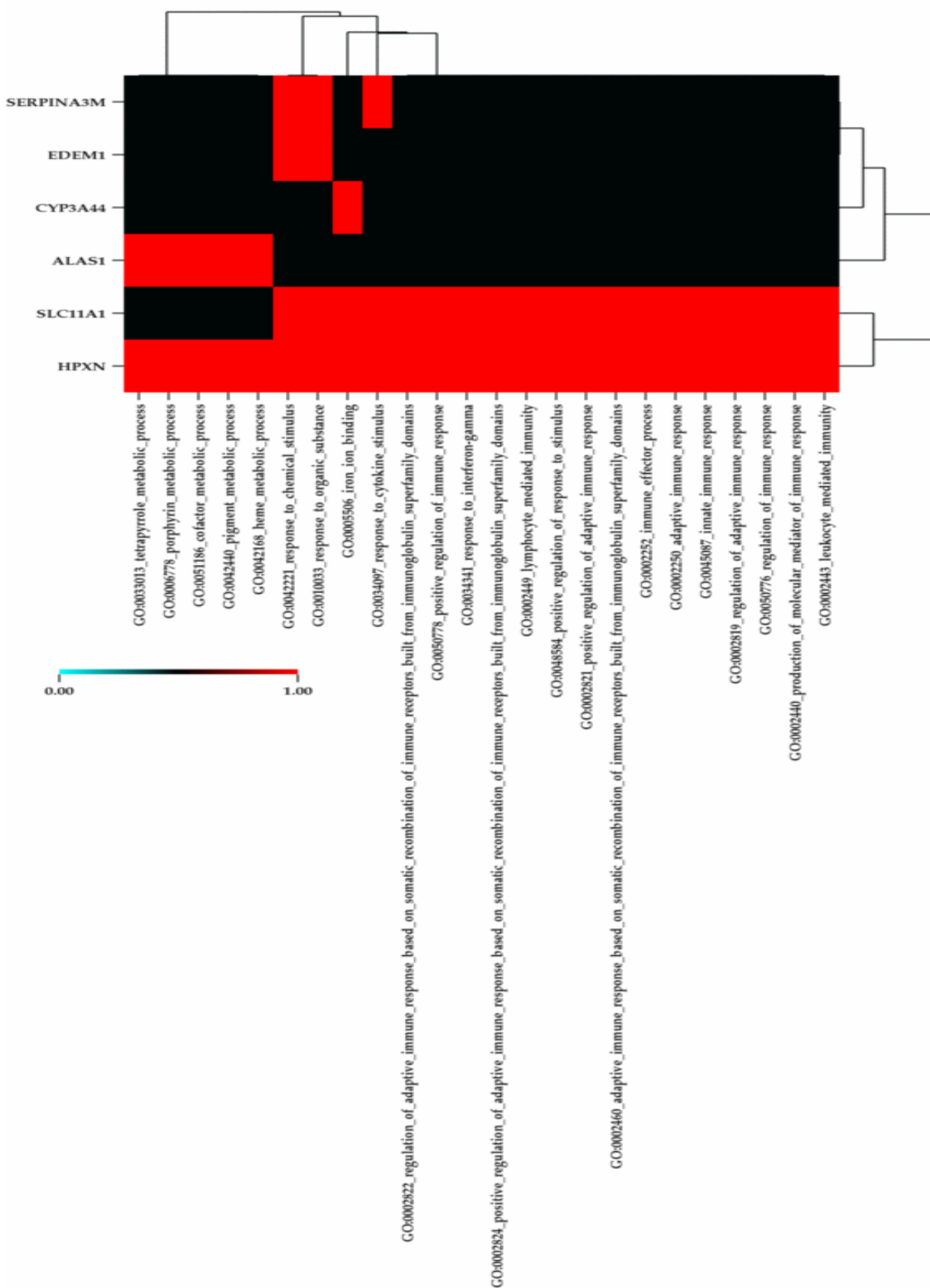


Figure 3.42 Heatmap for microarray analysis of CPA treated samples after 3 hours. Male F-344NHsd rats aged 14-15 weeks dosed with 100 mg kg⁻¹ CPA, then killed after 24 hours. Microarray analysis was determined as in Section 2.2.5, this heatmap was established with High-Throughput GoMiner (Zeeberg et al., 2005). Red demonstrates up-regulated, and blue demonstrates down-regulated (-1 down-regulated, +1 up-regulated).

Section 3.4 Transcriptome sequencing analysis

To study the early genes whose expression may be linked to the induction of hepatic DNA synthesis, RNA sequencing (whole transcriptome sequencing) was applied to the mRNAs isolated from F-344 fisher rats dosed with 50mg kg⁻¹ ciprofibrate or corn oil and killed after 3 hours. This was done with the Next Generation sequencing facilities, at the Deep seq research facility at the University of Nottingham (The AB SOLID 3 platform) as described in Section 2.2.6. The number of genes that resulted from this analysis was 24962 genes, with 705 genes that have a FDR p-value correction of <0.05, and 4245 genes that have the p-value is <0.05. In this analysis reads per kilobase of exon model per million mapped reads (RPKM) was used, which normalises for the difference in number of mapped reads between the samples (4 treated and 4 control samples).

Section 3.4.1 Comparing the data in a scatter plot

To show the effect of the ciprofibrate against the control on the genes a scatter plot was applied, this represents the number and distribution of the genes after the analysis. Shown in Figure 3.43 the spread of the control normalised genes against the treated normalised genes is illustrated The original data can be found at http://spldeepseq.nottingham.ac.uk/~aziz/wtp_rat/.

The points closest to the linear regression plot (the x=y line in the plot) are the genes which are least affected by the treatment. Genes of interest would be represented by the points that are most dispersed along an axis perpendicular to the linear regression plot. It is clarified in Figure 3.43 that the genes are behaving in a similar way because most of the genes are close to the x=y line, but if they are affected by the ciprofibrate as there is a drift away from the main line, and these should be the differently induced genes that are of interest.

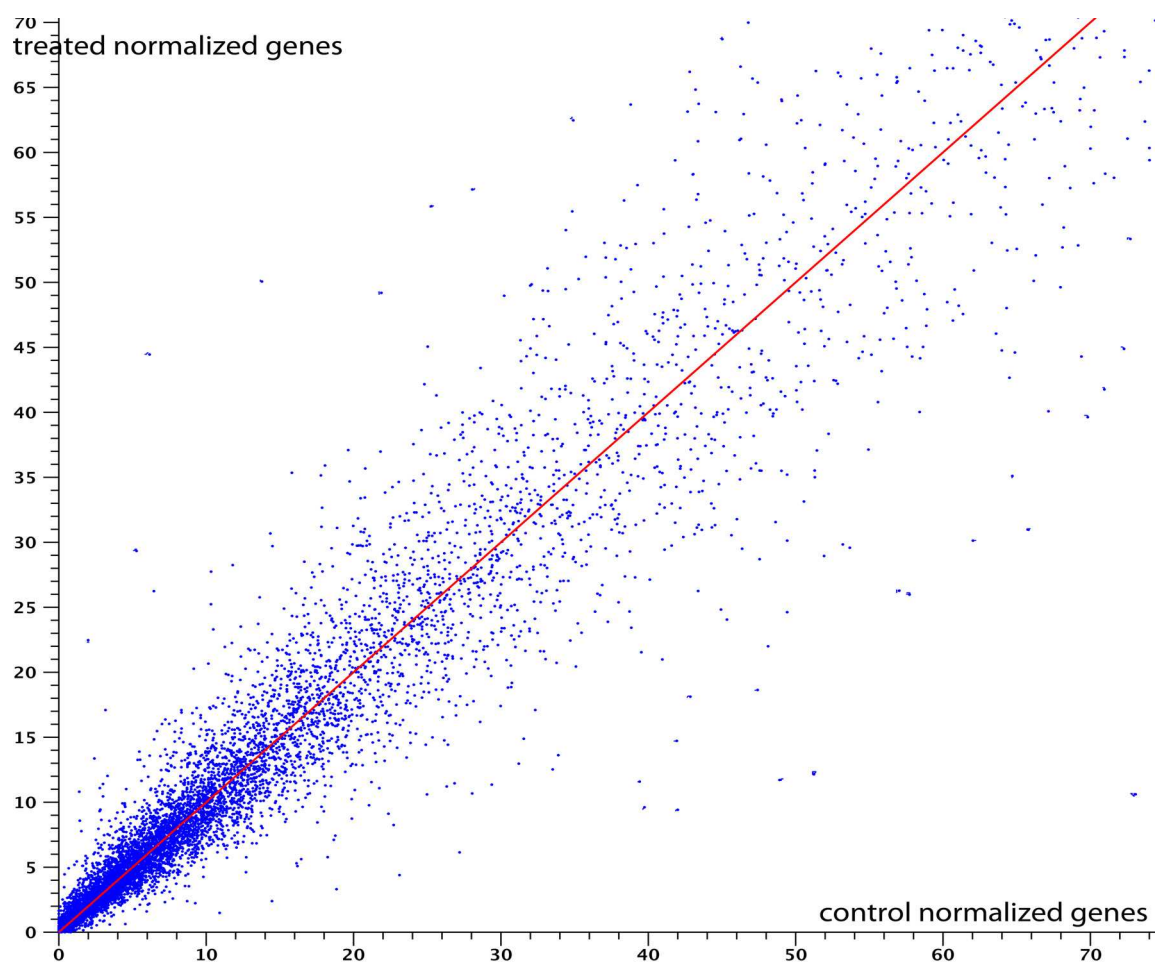


Figure 3.43 Scatter plot view of transcriptome data. A scatter plot view of the treated normalised mean data for each gene against the control normalised means, from the liver-RNA of F-344NHsd fisher rats dosed with 50 mg kg⁻¹ ciprofibrate (or corn oil) for 3 hours. Shown in red is a linear regression plot of the points. Each point indicates the relationship between the control and treated samples. (Software used The CLC Genomics Workbench).

Section 3.4.2 Checking between and within the groups variability

The objective here is to compare and examine the general distributions of the data within the same group. This is important to exclude any flaws in the preparation or other defects in the control and treated groups, so box plots were used.

The normalised expression values of the control data are shown in Figure 3.44 where the 4 samples are similar in spread (the upper and lower bars are aligned) and the median lines in all the boxes are aligned. Similar results were observed in the normalised treated data (Figure 3.45) and the data for the whole experiment (when comparing the treated against the control) (Figure 3.46).

From the spread of the samples in the figures it can be concluded that the samples are equally distributed within the control/ treated groups and in the whole experiment, and this indicates that the preparation and processing of the samples were correctly labelled and assigned in each of the groups.

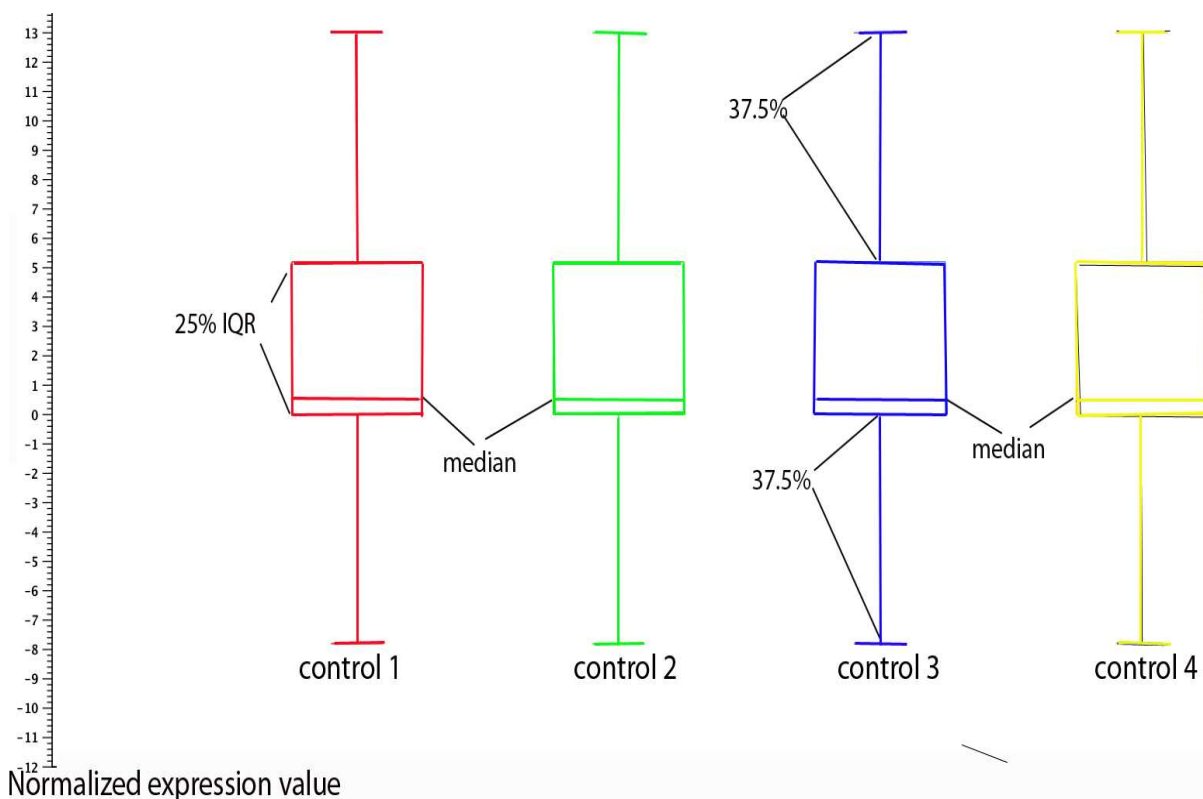


Figure 3.44 Analysing distributions of the control data. Four box plots of the normalised expression values of each of the 4 samples from control livers of F-344 fisher rats dosed with corn oil for 3 hours. Each box plot refers to the distribution of the data, and includes a square box indicating the IQR (Inter quartile Range 25%) value for each sample’s distribution (from the lower to the upper quartile), the lines displayed in the square boxes are the median and the upper and lower bars extend 1.5 (75%) the height of the box.(Software used The CLC Genomics Workbench).

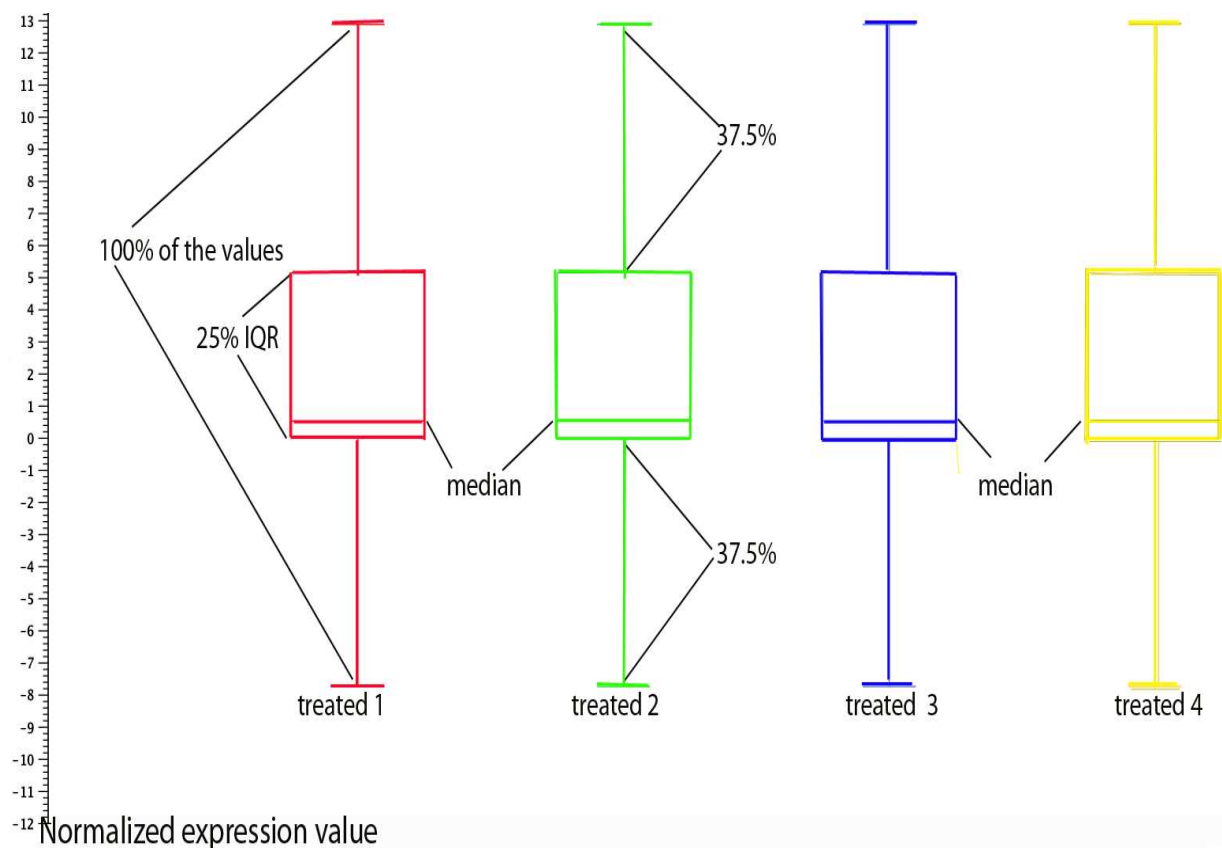


Figure 3.45 Analysing distributions of the treated data. Four box plots of the normalised expression values of each of the 4 samples from livers of F-344 fisher rats treated with 50 mg kg⁻¹ ciprofibrate for 3 hours. Each box plot refers to the distribution of the data, and includes a square box indicating the IQR (Inter quartile Range 25%) value for each sample’s distribution (from the lower to the upper quartile), the lines displayed in the square boxes are the median and the upper and lower bars extend 1.5 (75%) the height of the box (Software used The CLC Genomics Workbench).

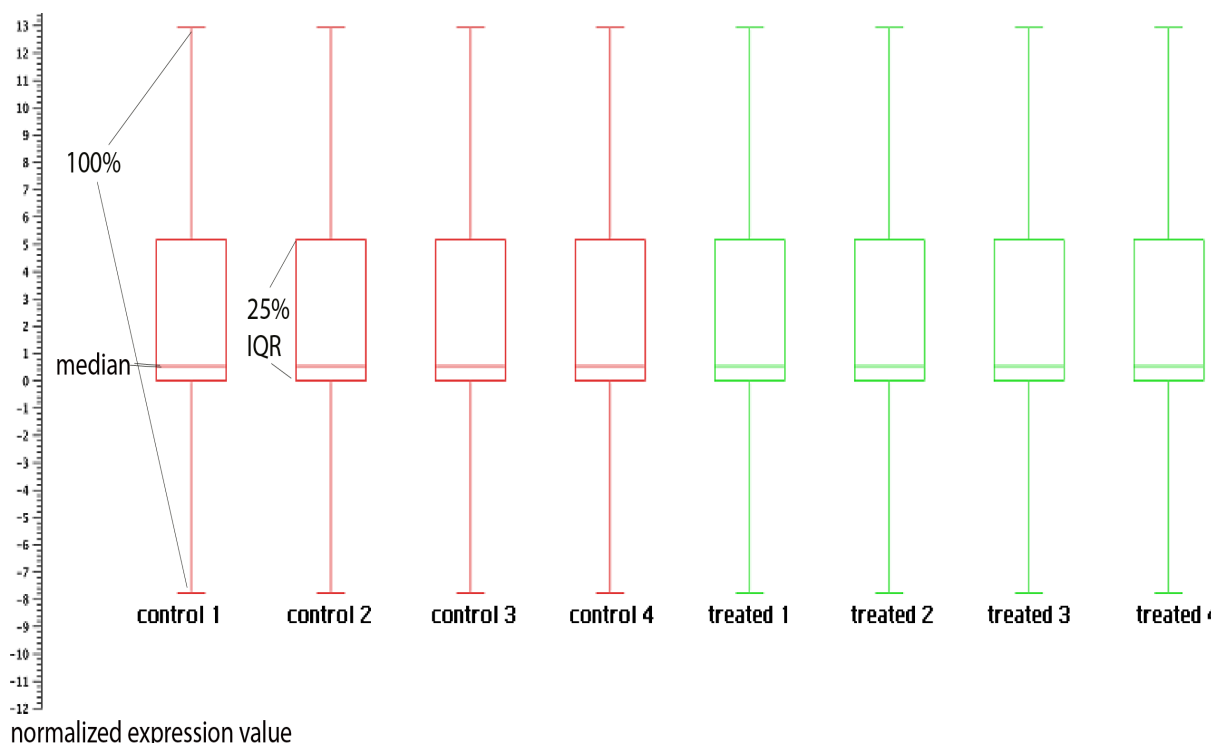


Figure 3.46 Analysing distributions of all the transcriptome data. 8 box plots of the normalised expression values of all 8 samples from livers of F344 fisher rats treated with 50 mg kg^{-1} ciprofibrate/corn oil for 3 hours. Red are the 4 control (corn oil) samples and green are the 4 treated samples. Each box plot refers to the distribution of the data, and includes a square box indicating the IQR (Inter quartile Range 25%) value for each sample's distribution (from the lower to the upper quartile), the lines displayed in the square boxes are the median and the upper and lower bars extend 1.5 (75%) the height of the box. (Software used The CLC Genomics Workbench).

Section 3.4.3 A volcano plot view of the data

This was done to see the genes on the basis of the p-value and the fold change differences between the control and the ciprofibrate treated samples for 3 hours, by applying the volcano plot view of the data.

A volcano plot of the results from the RNAseq analysis is illustrated in Figure 3.47 showing the \log_{10} of the p-value (of the Student's t- test of the difference between the control and treated samples) and the \log_2 of the fold change difference between the control and treated samples. The plots closest to the zero value for \log_2 fold change are the genes with least change in expression, while the genes with significantly changed expression are indicated within the plots in

red (statistics done was with Students' t- test) data used, and had a false discovery rates FDR p-values of < 0.05 , and a fold change of ± 2 .

The volcano plot shows the relationship between the p-values and the variation in expression values of the control and treated groups. Points for gene expressions with statistically significant differences will be positioned in the upper left and upper right of the volcano plot specified in red in Figure 3.47. This shows that the samples are mostly similar with only a small number of genes that are differently (up or down) regulated. These genes are the genes of interest for this study.

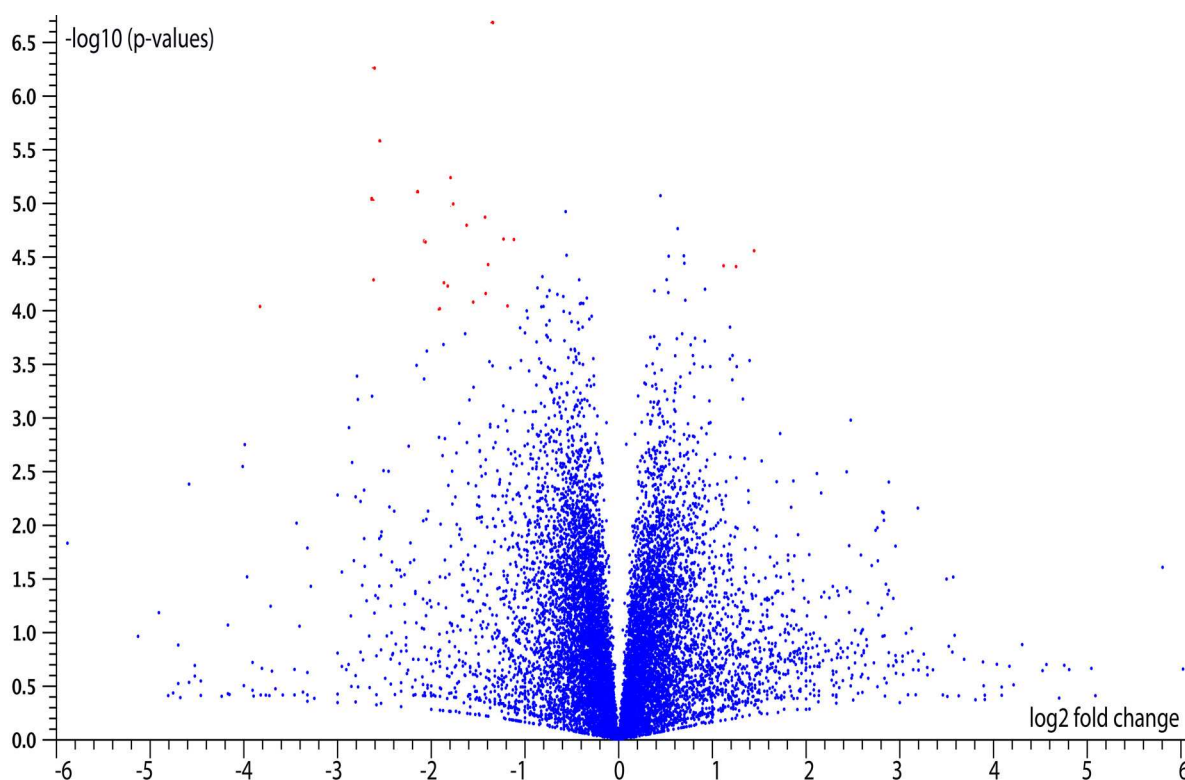


Figure 3.47 volcano plot view of transcriptome data (Student's t- test). A volcano plot view of the log₁₀ p-value versus the log₂ fold change difference between the control and treated samples, from the hepatic RNA of F-344NHsd fisher rats dosed with 50 mg kg⁻¹ ciprofibrate (or corn oil) for 3 hours. Red points are the features with false discovery rates (FDR) p-values of < 0.05 , and a fold change of ± 2 . (Software used The CLC Genomics Workbench).

Section 3.4.4 Hierarchical clustering of samples

Hierarchical clustering was presented to see the comparative similarities of the control and treated samples, and to check if they cluster homogeneously or not.

The hierarchical cluster is a heat map with the clustering of the samples at the bottom. The tree was arranged by placing each group of related genes as a cluster and calculating the pair-wise space between all clusters, then combining each two closest clusters to be made into one new cluster and then the next closest joined and so on until there was only one cluster left (which contains all samples). In the tree the spaces between the clusters reflect the distance of the branches in the tree. Therefore, samples that have closely resembled gene expressions have short distances between them, and those that are more different are separated further.

In Figure 3.48 the samples on the left are the treated samples and the samples on the right are the controls, the red and blue colours express the induction of the genes in the groups (red indicates high gene expression and blue presents low expression of the genes), the hierarchical clustering of the samples clarifies the relationship in between the groups, and shows that the treatment has affected the gene induction and that it is different from the untreated.

The tree cluster at the bottom of Figure 3.48 shows that the samples similar in their gene expression are together, and also indicates that the ciprofibrate treatment replicates had a similar effect on the genes at 3 hours (the red colour is more abundant than the blue). This is different from the control replicates that are all clustered together.

As a conclusion the treated replicates show similar effects on genes that appear highly induced, and the same for the control replicates, which also show a difference between the treated and control groups.

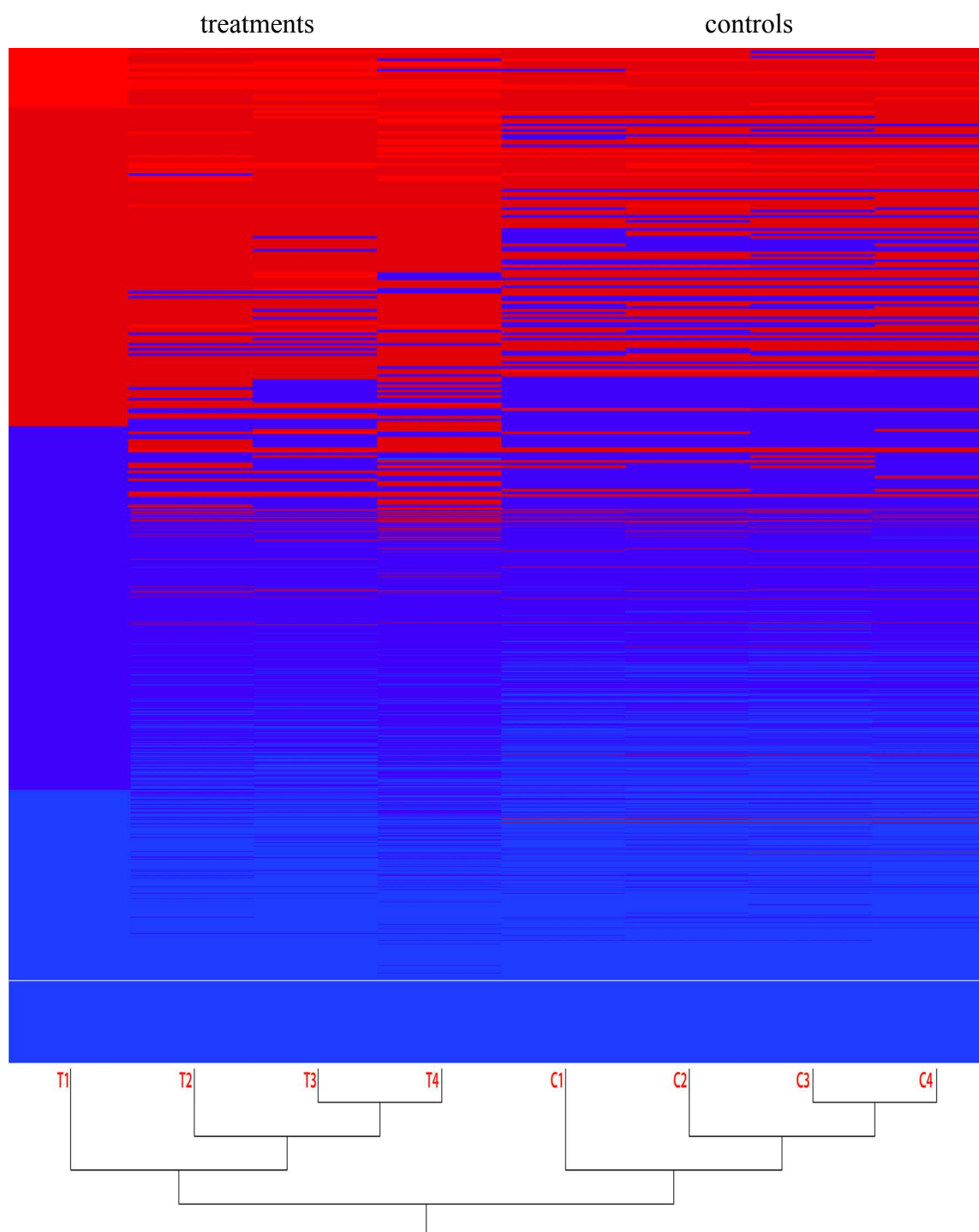


Figure 3.48 A hierarchical cluster of samples. Male Fisher rats were treated with 50mg kg⁻¹ ciprofibrate or corn oil for 3 hours, then RNAseq analysis was applied on the hepatic RNA samples as described in Section 2.2.6. At the top of the figure are listed the names of the samples the treatments (4 groups) on the left and the controls (4 groups) on the right. The horizontal lines present the features (genes) and are coloured according to the expression level. Red presents the maximum level of expression and blue shows the minimal level of expression. The tree at the bottom of the heat-map visualizes the clustering. The features are sorted by the expression intensity in the first sample on the left. (Software used The CLC Genomics Workbench).

Section 3.4.5 Hierarchical and K-means clusters of features

Feature clustering was performed with the results from the RNAseq analysis to recognize and cluster together genes with related gene expression values from the controls or samples treated with 50 mg kg⁻¹ ciprofibrate for 3 hours. Features that are clustered together are expected to be involved in similar biological procedures.

Hierarchical clustering of features was a tree arrangement of the similar expressed genes over the set of control and treated samples (or groups). In the hierarchical cluster each feature is a cluster. Then the pairwise distance between the clusters was evaluated and the two closest clusters combined into one new cluster. Then the clusters were grouped together depending on the relationship of the gene induction until there was only one cluster left. This feature clustering was performed on a subset of the data after filtering away the genes that have low fold difference values or have little difference between the samples (large p-value).

With the hierarchical clustering a number of the genes were found by the feature clustering to be more expressed in the treated than that in the control group, e.g. *Cyp4A1*, *G0/G1switch 2* and *Scd1* genes (sterol-Coenzyme A desaturase 1), and the last two genes are more closely clustered together than the *Cyp4A1* gene.

The feature clustering was also performed as a k-means clustering, where the features (gene expressions) are clustered together in groups depending on the expression response to the control and treated samples. This clustering resulted in presenting how some genes were down-regulated in the control samples and up-regulated in the treated samples, or presenting how the gene expression behaves in a different way, where the genes were up-regulated in the control samples then the same gene expression goes down in the treated samples, or how the gene expression in the control and treated samples have similar expression values.

Section 3.4.6 Induced genes by ciprofibrate after 3 hours with RNAseq

The objective from this is to find genes that are differential expressed in hepatic cells of F344NHsd Fisher rats dosed with 50mg kg⁻¹ ciprofibrate after 3 hours see (Section 2.2.1). This should highlight genes whose expression is correlated with the hepatic DNA synthesis that occurs at 24 hours.

This was done by Transcriptome Sequencing analysis of the hepatic RNA detailed in Section 2.2.6. Stastical analysis was done with Student t- test and genes were defined as significant if the FDR p-value was < 0.05. A whole list of all the ~25000 genes and associated data analysis is provided in the CD at the end of this thesis.

Section 3.4.6.1 Down- regulated genes with RNAseq

Genes were considered down- regulated if the FDR p-value was > 0.05 and there was a fold change in expression of < -1.5 (0.67 fold). Table 3.18 shows a list of down- regulated known (named) genes (e.g. *Igfbp2*, *ppargcla*, *Cyp1a2*, *AhR*.....), and in Table 3.19 is a list of the novel genes that have not been named to date.

Table 3.18 Named down- regulated genes in ciprofibrate dosed rats treated for 3 hours. A list of down-regulated genes for male F-344NHsd rats aged 14-15 weeks dosed with 50 mg kg⁻¹ ciprofibrate then killed after 3 hours. RNAseq analysis was done as described in Section 2.2.6. Expressed genes with a fold change of <-1.5 and a FDR p-value of <0.05 were listed.

Description	Gene name	Ensembl Feature ID	Fold Change <-1.5	FDR P-value <0.05
Insulin-like growth factor-binding protein 2 Precursor (IGF-binding protein 2)	Igfbp2	ENSRNOT00000023068	-5.0	0.01
Peroxisome proliferator-activated receptor gamma, coactivator 1 alpha	Ppargc1a	ENSRNOT00000006071	-2.1	0.01
Cytochrome P450, family 1, subfamily a, polypeptide 2 Gene	Cyp1a2	ENSRNOT00000058571	-1.7	0.03
Acyl-CoA synthetase medium-chain family member 2	Acsm2	ENSRNOT00000020587	-3.3	0.02
Hepatocyte nuclear factor 6 (One cut domain family member 1)	HNF-6	ENSRNOT00000010738	-2.2	0.03
Aryl hydrocarbon receptor	Ahr	ENSRNOT00000006618	-2.0	0.04
Immediate early response 2- immediate early response gene 2 protein	Ier2	ENSRNOT00000003793	-1.8	0.02
Forkhead box D1 (mediates the coordinated expression of hepatocyte-specific genes)	Foxd1	ENSRNOT00000010201	-1.9	0.04
Death-associated kinase 2	dapk2	ENSRNOT00000023372	-1.7	0.04
Solute carrier family 2 (facilitated glucose transporter), member 1	Slc2a1	ENSRNOT000000064452	-2.4	0.04
Protein kinase-like protein SgK493	Sgk493	ENSRNOT00000005664	-2.2	0.04
Oncogene-serine threonine protein kinase	pim-1	ENSRNOT00000000637	-2.0	0.03
Claudin 9 (predicted); LOC287099; protein-coding	Cldn9	ENSRNOT000000004848	-2.1	0.03
Claudin 6 (protein binding)	Cldn6	ENSRNOT000000004845	-1.9	0.03
Claudin 4	Cldn4	ENSRNOT000000002003	-2.0	0.04

Table 3.18 **Named down- regulated genes in ciprofibrate dosed rats treated for 3 hours.** A list of down-regulated genes for male F-344NHsd rats aged 14-15 weeks dosed with 50 mg kg⁻¹ ciprofibrate then killed after 3 hours. RNAseq analysis was done as described in Section 2.2.6. Expressed genes with a fold change of <-1.5 and a FDR p-value of <0.05 were listed.

Description	Gene name	Ensembl Feature ID	Fold Change <-1.5	FDR P-value <0.05
Sialidase 2 (cytosolic enzyme that cleaves sialic acid; involved in the degradation of glycolipids and glycoproteins)	Neu2	ENSRNOT00000022818	-2.1	0.02
Proline rich 16 similar to mesenchymal stem cell protein DSC54 (predicted)	Prr16	ENSRNOT00000026724	-1.9	0.04
Zink finger protein 36-acts as a transcriptional activator	Zfp36	ENSRNOT00000026661	-1.9	0.03
SH3-domain binding protein 5 (BTK-associated)-may bind c-Jun N-terminal kinase (JNK); may play a role in JNK and Bruton's tyrosine kinase (Btk) mediated signaling pathways	Sh3bp5	ENSRNOT00000026389	-1.8	0.03
UDP-Gal:betaGlcNAc beta 1,3-galactosyl transferase, polypeptide 1	B3galt1	ENSRNOT00000010018	-1.8	0.01
Lipopolysaccharide binding protein-binds lipopolysaccharide on outer membrane of gram negative bacteria; involved in immune response	Lbp	ENSRNOT00000019787	-1.8	0.04
Leukocyte cell-derived chemotaxin 2	Lect2	ENSRNOT00000016252	-1.8	0.04
Small nuclear ribonucleoprotein 25 (U11/U12)	Snrnp25	ENSRNOT00000067079	-1.8	0.03
Tripartite motif-containing 36	Trim36	ENSRNOT00000022294	-1.8	0.04
Sprouty homolog 2 (Drosophila)	Spry2	ENSRNOT00000013342	-1.7	0.04
Sec61 gamma subunit, pseudogene 1 gama subunit-like Q7Tq10_RAT	SEC61g-ps1	ENSRNOT00000043104	-1.7	0.02
Phosphodiesterase 4C, cAMP-specific (phosphodiesterase E1 dunce homolog, Drosophila- putative nucleotide phosphodlerase E1 dunce homolog, drosophila)	Pde4c	ENSRNOT00000026457	-1.6	0.04
UDP-N-acteylglucosamine pyrophosphorylase 1-like 1.	Uap1/1	ENSRNOT00000017373	-1.6	0.02

Table 3.18 **Named down- regulated genes in ciprofibrate dosed rats treated for 3 hours.** A list of down-regulated genes for male F-344NHsd rats aged 14-15 weeks dosed with 50 mg kg⁻¹ ciprofibrate then killed after 3 hours. RNAseq analysis was done as described in Section 2.2.6. Expressed genes with a fold change of <-1.5 and a FDR p-value of <0.05 were listed.

Description	Gene name	Ensembl Feature ID	Fold Change <-1.5	FDR P-value <0.05
Hermansky-Pudlak syndrome 4 homolog (human)-human homolog is involved in the biogenesis of lysosomes and related cellular vesicles; mutations in the human gene are associated with Hermansky-Pudlak syndrome	Hps4	ENSRNOT00000000824	-1.6	0.02
Sodium-dependent phosphate transport protein 1	Slc17a1	ENSRNOT00000066313	-1.6	0.04
Translocase of outer mitochondrial membrane 7 homolog (yeast)	Tomm7	ENSRNOT00000065962	-1.6	0.02
Transmembrane protein 38a	Tmem38a	ENSRNOT00000015877	-1.5	0.01
High mobility group nucleosomal binding domain 3	Hmgn3	ENSRNOT00000039917	-1.5	0.04
ADP- ribosylation factor- like 4A - may play role in GTP binding and hydrolysis	arl4a	ENSRNOT00000005800	-1.6	0.04
ADP- ribosylation factor-like	arl11	ENSRNOT00000019646	-1.5	0.04
Ribosomal protein S27-ribosomal protein subunit	Rps27	ENSRNOT00000022897	-1.7	0.02
Ribosomal protein S14-structural component of the 40S subunit of the ribosome, the organelle responsible for protein synthesis	Rps14	ENSRNOT00000059501	-1.8	0.05
Ribosomal protein L10A 60S ribosomal subunit protein	Rpl10a	ENSRNOT00000000603	-1.5	0.02
Ribosomal protein L11	Rpl11	ENSRNOT00000030043	-1.5	0.01
Ribosomal protein L23a	Rpl23a	ENSRNOT00000035657	-1.5	0.03
SnRNA no protein product	U8	ENSRNOT00000054056	-5.3	0.01

Table 3.19 Novel down- regulated genes in rats dosed with ciprofibrate for 3 hours. A list of novel down-regulated genes for male F-344NHsd rats aged 14-15 weeks, dosed with 50 mg kg⁻¹ ciprofibrate then killed after 24 hours. RNAseq analysis was done as described in Section 2.2.6. Expressed genes with a fold change of <-1.5 and a FDR p-value of <0.05 are shown.

Description	Gene name	Ensembl Feature ID	Fold change >1.5	FDR P-value <0.05
Pseudogene (a noncoding sequence similar to an active protein) no protein product	novel	ENSRNOT00000011532	-9.0	0.05
Pseudogene (no protein product)	novel	ENSRNOT00000025854	-1.9	0.04
Novel MiRNA microarray (a single-stranded RNA, typically 21-23 base pairs long), thought to be involved in gene regulation (especially inhibition of protein expression)	novel	ENSRNOT00000054415	-1.7	0.04
Protein coding- a protein coding transcript is a spliced mRNA that leads to a protein product	novel	ENSRNOT00000002046	-1.8	0.05
Protein coding	novel	ENSRNOT000000065817	-1.6	0.03
Protein coding	novel	ENSRNOT000000060517	-1.6	0.04
Protein coding	novel	ENSRNOT000000004213	-1.5	0.03
Protein coding transcript	RGD1563551	ENSRNOT000000047507	-3.1	0.04
Protein coding transcript	RGD1562755	ENSRNOT000000051559	-2.5	0.01
Protein coding transcript	LOC691255	ENSRNOT000000036330	-1.9	0.03
Pseudogene gene a noncoding sequence similar to an active protein (no protein product)	LOC365595	ENSRNOT00000000346	-2.1	0.03
Pseudogene	LOC367016	ENSRNOT000000030115	-1.8	0.05
Ribosomal protein P1-like	LOC10036522	ENSRNOT00000015893	-1.5	0.03
Similar to ribosomal protein L30	RGD1562397	ENSRNOT000000044756	-1.5	0.04
Similar to 60S acidic ribosomal protein P1	RGD1565054	ENSRNOT000000034714	-1.6	0.01

Table 3.19 Novel down- regulated genes in rats dosed with ciprofibrate for 3 hours. A list of novel down-regulated genes for male F-344NHsd rats aged 14-15 weeks, dosed with 50 mg kg⁻¹ ciprofibrate then killed after 24 hours. RNAseq analysis was done as described in Section 2.2.6. Expressed genes with a fold change of <-1.5 and a FDR p-value of <0.05 are shown.

Description	Gene name	Ensembl Feature ID	Fold change >1.5	FDR P-value <0.05
Similar to dystonin isoform 1	LOC680875	ENSRNOT00000014191	-1.5	0.04
Similar to 60S ribosomal protein L23a	LOC689899	ENSRNOT00000040647	-1.5	0.01
Similar to ribosomal protein S27a	RGD1564290	ENSRNOT00000027780	-1.6	0.03
Similar to ribosomal protein S27a	RGD1560997	ENSRNOT00000051805	-1.6	0.01

Section 3.4.6.2 Up- regulated genes with RNAseq

Genes were considered up- regulated if the FDR p-value was more than 0.05 and there was a fold change in expression of > 2.5 (because the interest is in the up-regulated genes). Table 3.20 shows a list of up- regulated known (named) genes and in Table 3.21 is a list of the novel genes that have not been named to date.

Table 3.20 Named up- regulated genes in ciprofibrate dosed rats treated for 3 hours. A list of up- regulated genes for male F-344NHsd rats aged 14-15 weeks dosed with 50 mg kg⁻¹ ciprofibrate then killed after 3 hours. RNAseq analysis was done as described in Section 2.2.6. Expressed genes with a fold change of >2.5 and a FDR p-value of <0.05 were listed.

Description	Gene name	Ensembl Feature ID	Fold Change >2.5	FDR p-value <0.05
Cytochrome P450, family 4, subfamily a, polypeptide 10-member of the cytochrome P450 monooxygenase enzyme superfamily; plays a role in lipid metabolism; involved in androgen mediated signalling	Cyp4a10	ENSRNOT00000051385	3.2	0.01
Cytochrome P450 4X1 cytochrome P450 protein; may play a role in neurovascular function cytochrome P450, family 4, subfamily x, polypeptide 1- cytochrome P450 protein; may play a role in neurovascular function	Cyp4x1	ENSRNOT00000011985	3.1	0.04
Peroxisome proliferator-activated receptor gamma, coactivator 1 beta	Ppargc 1b	ENSRNOT00000023661	2.9	0.03
G0/G1switch 2	G0s2	ENSRNOT00000007879	5.8	0.01
Arylacetamide deacetylase-like 1 - exhibits catalytic activity and hydrolase activity; involved in metabolism	Aadacl1	ENSRNOT00000017805	6.1	0.02
Acyl-CoA thioesterase 5	Acot5	ENSRNOT00000013760	2.8	0.01
Acyl-CoA thioesterase 6	Acot6	ENSRNOT00000058101	4.0	0.01
Acid phosphatase, prostate-	Acpp	ENSRNOT00000016222	3.2	0.03
Angiotensin-like 4 -a circulating protein which causes an increase in plasma very low density lipoprotein by inhibition of lipoprotein lipase activity	Angptl4	ENSRNOT00000010031	2.7	0.01
Apolipoprotein L 9a	Apol9a	ENSRNOT00000031951	2.8	0.02
Arrestin domain containing 4	Arrdc	ENSRNOT00000051402	3.5	0.01
Calcium channel, voltage-dependent, alpha 2/delta sub-unit 4	Cacna2d4	ENSRNOT00000010746	3.3	0.01
Chemokine binding protein 2 CC-chemokine receptor; may have a role in placental immunity or hematopoiesis	Ccbp	ENSRNOT00000026343	4.5	0.05

Table 3.20 Named up- regulated genes in ciprofibrate dosed rats treated for 3 hours. A list of up- regulated genes for male F-344NHsd rats aged 14-15 weeks dosed with 50 mg kg⁻¹ ciprofibrate then killed after 3 hours. RNAseq analysis was done as described in Section 2.2.6. Expressed genes with a fold change of >2.5 and a FDR p-value of <0.05 were listed.

Description	Gene name	Ensembl Feature ID	Fold Change >2.5	FDR P-value <0.05
Cyclin D1 a cell cycle protein involved in the regulation of cell proliferation; associated with many cancers and other diseases	Ccnd1	ENSRNOT00000028411	2.9	0.03
Carnitine palmitoyltransferase 1b, muscle - muscle isoform of enzyme that catalyses the transfer of long chain fatty acids to carnitine for translocation across the mitochondrial inner membrane	Cpt1b	ENSRNOT00000013985	4.2	0.02
Carnitine palmitoyltransferase 2 - inner mitochondrial membrane protein that converts acylcarnitine to acyl-CoA	Cpt2	ENSRNOT00000016954	2.6	0.01
Eph receptor A2	Epha2	ENSRNOT00000066072	3.3	0.04
Ets variant 3-like	Etv3l	ENSRNOT00000057490	2.9	0.04
Ets variant 6 human homolog is an ETS family transcription factor; may be involved in hematopoiesis and maintenance of the vascular network	Etv6	ENSRNOT00000007889	4.6	0.01
Fin bud initiation factor homolog (zebra fish)	Fiin	ENSRNOT00000006203	3.2	0.04
Inhibin beta E cytokine; involved in cell fate determination	Inhbe	ENSRNOT00000010106	3.7	0.03
Kelch repeat and BTB (POZ) domain containing 11	Kbtbd11	ENSRNOT00000016420	6.5	0.01
Known pseudogene potassium channel tetramerisation domain containing 12	Kctd12	ENSRNOT00000037179	2.9	0.02
Keratin 23 (histone deacetylase inducible)	Krt23	ENSRNOT00000016657	3.9	0.01
MHC I like leukocyte 2	Mill2	ENSRNOT00000048149	3.7	0.01
Myotubularin related protein 7	Mtmr7	ENSRNOT00000065327	4.2	0.03
Myeloblastosis oncogene-like 1	Mybl1	ENSRNOT00000066911	2.6	0.01

Table 3.20 Named up- regulated genes in ciprofibrate dosed rats treated for 3 hours. A list of up- regulated genes for male F-344NHsd rats aged 14-15 weeks dosed with 50 mg kg⁻¹ ciprofibrate then killed after 3 hours. RNAseq analysis was done as described in Section 2.2.6. Expressed genes with a fold change of >2.5 and a FDR p-value of <0.05 were listed.

Description	Gene name	Ensembl Feature ID	Fold Change >2.5	FDR P-value <0.05
Myc myelocytomatosis viral oncogene homolog 1, lung carcinoma derived (avian) mouse lung carcinoma myc related oncogene 1 (Lmyc1) is a DNA binding protein with cell cycle related transcriptional activities	Mycl1 v	ENSRNOT00000019101	3.6	0.02
Nuclear receptor subfamily 4, group A, member 2- nuclear receptor and transcription factor; plays a role in development and maintenance of neurons synthesizing the neurotransmitter dopamine	Nr4a2	ENSRNOT00000041394	2.8	0.01
Progesterin and adipo Q receptor family member VII	Paqr7	ENSRNOT00000022713	5.7	0.01
Peptidase M20 domain containing 2; aminoacylase 1-like 2; aminoacylase 1-like 2 (predicted); acy112; LOC313130; Acy112; Acy112_predicted	Pm20d2	ENSRNOT00000060914	3.8	0.05
Protein phosphatase 1K (PP2C domain containing)	Ppm1k	ENSRNOT00000009202	2.5	0.03
PR domain containing 6	Prdm6	ENSRNOT00000047755	2.6	0.03
Prospero homeobox 2	Prox2	ENSRNOT00000006704	3.2	0.04
Proline rich Gla (G-carboxyglutamic acid) 4 (trans membrane)	Prrg4	ENSRNOT00000038464	7.9	0.02
Putative neuronal cell adhesion molecule	Punc	ENSRNOT00000051353	6.3	0.04
RAS, dexamethasone-induced 1- interacts with neuronal NO synthase adaptor protein CAPON and is involved in nitric oxide mediated signalling	Rasd1	ENSRNOT00000004475	2.6	0.03
Ras responsive element binding protein 1	Rreb1	ENSRNOT00000021007	2.5	0.02
Sterile alpha and TIR motif containing 1	Sarm1	ENSRNOT00000013639	2.8	0.04
Stearoyl-Coenzyme A desaturase 1-enzyme involved in the synthesis and regulation of unsaturated fatty acids	Scd1	ENSRNOT00000051086	13.2	0.01
Stearoyl-coenzyme A desaturase 4	Scd4	ENSRNOT00000017834	15.6	0.02

Table 3.20 Named up- regulated genes in ciprofibrate dosed rats treated for 3 hours. A list of up- regulated genes for male F-344NHsd rats aged 14-15 weeks dosed with 50 mg kg⁻¹ ciprofibrate then killed after 3 hours. RNAseq analysis was done as described in Section 2.2.6. Expressed genes with a fold change of >2.5 and a FDR p-value of <0.05 were listed.

Description	Gene name	Ensembl Feature ID	Fold Change >2.5	FDR P-value <0.05
Solute carrier family 16, member 6 (monocarboxylic acid transporter 7) - human homolog is a monocarboxylate transporter	Slc16a6	ENSRNOT0000000262	6.4	0.03
Solute carrier family 22 (organic cation/carnitine transporter), member 5 mediates high-affinity sodium-dependent carnitine transport and sodium-independent organic cation transport	Slc22a5	ENSRNOT00000011340	2.7	0.01
Solute carrier family 25, member 30 kidney mitochondrial carrier protein 1; solute carrier family 25 member 30; solute carrier family 25, member 30, Kidney mitochondrial carrier protein 1	Slc25a30 / KMCP1_rat	ENSRNOT00000040316	3.0	0.04
Solute carrier family 34 (sodium phosphate), member 2- a sodium dependent phosphate transporter that may provide inorganic phosphate for the synthesis of lung surfactant and is associated with aging	Slc34a2	ENSRNOT00000048509	5.6	0.04
T-box 3 - human homolog acts as a transcriptional repressor and plays a role in the development of several organ systems	Tbx3	ENSRNOT00000011552	3.1	0.03

Table 3.21 Novel up- regulated genes in rats dosed with ciprofibrate for 3 hours with RNAseq analysis.

A list of novel down- regulated genes for male F-344NHsd rats aged 14-15 weeks, dosed with 50 mg kg⁻¹ ciprofibrate then killed after 24 hours. RNAseq analysis was done as described in Section 2.2.6. Expressed genes with a fold change of >2.5 and a FDR p-value of <0.05 are shown.

Description	Gene name	Ensembl Feature ID	Fold Change >2.5	FDR p-value <0.05
Fer-1-like 5 (C. elegans) Gene	novel	ENSRNOT00000059806	2.6	0.03
XIAP associated factor 1 Gene	novel	ENSRNOT00000056551	2.6	0.04
Known protein coding	novel	ENSRNOT00000061062	3.5	0.02
Known protein coding	novel	ENSRNOT00000041253	2.6	0.02
Protein coding	novel	ENSRNOT00000061064	3.1	0.04
Protein coding	novel	ENSRNOT00000061065	3.1	0.04
Protein coding	novel	ENSRNOT00000016611	3.5	0.03
Known protein coding similar to serine/threonine kinase Synonyms: LOC296256; RGD1565143_predicted; similar to serine/threonine kinase (predicted);	RGD1565143	ENSRNOT00000065773	3.3	0.05
rCG62747-like	LOC100361582	ENSRNOT00000050476	3.3	0.01
mCG140381-like	LOC100363013	ENSRNOT00000005583	2.9	0.03
Similar to ras homolog gene family, member f	LOC690130	ENSRNOT00000064390	2.8	0.03
Similar to hypothetical protein MGC42105	RGD1308116	ENSRNOT00000021964	2.9	0.01
Ab2-060	LOC501038	ENSRNOT00000045455	2.7	0.04
RGD1560010	RGD1560010	ENSRNOT00000013356	3.6	0.02

Section 3.4.7 Pathways for over-representative genes

The identification of a number of up-regulated genes calls for characterisation of common pathways or functions in this gene subset. This was achieved by manual comparison of the up-regulated genes with pathways found at these websites <http://rgd.mcw.edu/wg/pathway?100> and http://www.ensembl.org/Rattus_norvegicus/Info/Index.

The pathways with relationship with DNA synthesis, cell cycle or PPAR signalling were studied. 16 genes were induced out of 73 PPAR signalling pathway genes (21.9%), when compared with KEGG website. Also, 12 up-regulated genes out of 43 fatty acid metabolic pathway genes (27.9%), and 10 out of 103 genes (9.7%) were engaged in the cell cycle. Other smaller groups of genes (5-6 genes) were involved in sodium ion transport, oxidation reduction and palmitoyl-CoA hydrolase activity as demonstrated in Table 3.22.

Pathways that are not listed in the table but have a number of up-regulated genes are metabolic process and bioactivation pathway via cytochrome P450.

Table 3.22 Some pathways represented by named up-regulated genes in ciprofibrate dosed rats for 3 hours. A list of pathways and the related genes are shown for up-regulated genes for male F-344NHsd rats aged 14-15 weeks dosed with 50 mg kg⁻¹ ciprofibrate then killed after 3 hours. RNAseq analysis was done as described in Section 2.2.6. Expressed genes with a fold change of >1.5 and a FDR p-value of <0.05 were listed.

Description	gene name	fold change
PPAR signalling pathway:		
Acetyl-Coenzyme A acyl transferase 1/peroxisomal 3-ketoacyl-CoA thiolase	Acaa1	1.9
Acyl-CoA synthetase long-chain family member 1	Acs11	1.8
Acyl-CoA synthetase long-chain family member 3	Acs13	2.0
Angiopoietin-like 4	Angptl4	2.7
Apolipoprotein L 9a	Apo19a	2.8
Carnitine palmitoyltransferase 1a, liver	Cpt1a	1.7
Carnitine palmitoyltransferase 1b, muscle	Cpt1b	4.2
Carnitine palmitoyltransferase 2	Cpt2	2.6
Cytochrome P450, family 4, subfamily a, polypeptide 1	Cyp4a1	3.3
Cytochrome P450, family 4, subfamily a, polypeptide 2	Cyp4a2	2.3
Cytochrome P450, family 4, subfamily a, polypeptide 3	Cyp4a3	2.3
Cytochrome P450, family 4, subfamily b, polypeptide 1	Cyp4b1	1.8
3-hydroxy-3-methylglutaryl-Coenzyme A synthase 2	Hmgcs2	1.7
Peroxisome proliferator-activated receptor gamma, coactivator 1 beta	Ppargc1b	2.9

Table 3.22 Some pathways represented by named up-regulated genes in ciprofibrate dosed rats for 3 hours. A list of pathways and the related genes are shown for up-regulated genes for male F-344NHsd rats aged 14-15 weeks dosed with 50 mg kg⁻¹ ciprofibrate then killed after 3 hours. RNAseq analysis was done as described in Section 2.2.6. Expressed genes with a fold change of >1.5 and a FDR p-value of <0.05 were listed.

Description	gene name	fold change
Stearoyl-Coenzyme A desaturase 1	Scd1	13.2
Stearoyl-coenzyme A desaturase 4	Scd4	15.6
Fatty acid metabolic pathway:		
Acetyl-Coenzyme A acyltransferase 1	Acaa1	1.9
Acyl-CoA synthetase long-chain family member 1	Acs11	1.8
Acyl-CoA synthetase long-chain family member 3	Acs13	2.0
Aldehyde dehydrogenase 3 family, member A2	Aldh3a2	1.7
Carnitine palmitoyltransferase 1a, liver	Cpt1a	1.7
Carnitine palmitoyltransferase 1b, muscle	Cpt1b	4.2
Carnitine palmitoyltransferase 2	Cpt2	2.6
Cytochrome P450, family 4, subfamily a, polypeptide 10	Cyp4a10	3.3
Cytochrome P450, family 4, subfamily a, polypeptide 2	Cyp4a2	2.3
Cytochrome P450, family 4, subfamily b, polypeptide 1	Cyp4b1	1.8
Hydroxyacyl-Coenzyme A dehydrogenase, alpha subunit	Hadha	1.6
Stearoyl-Coenzyme A desaturase 1	Scd1	13.2

Table 3.22 Some pathways represented by named up- regulated genes in ciprofibrate dosed rats for 3 hours. A list of pathways and the related genes are shown for up- regulated genes for male F-344NHsd rats aged 14-15 weeks dosed with 50 mg kg⁻¹ ciprofibrate then killed after 3 hours. RNAseq analysis was done as described in Section 2.2.6. Expressed genes with a fold change of >1.5 and a FDR p-value of <0.05 were listed.

Description	gene name	fold change
Cell cycle:		
Cell cycle associated protein 1 Gene	Caprin1	1.8
CC-chemokine receptor	Ccbp	4.5
Cyclin D1	Ccnd1	2.9
Cyclin T1	Ccnt1	1.8
G0/G1switch 2	G0s2	5.8
Nicotinamide phosphoribosyltransferase	Nampt	2.3
Nibrin	Nbn	2.0
Par-6 (partitioning defective 6) homolog beta	Pard6b	2.0
Retinoblastoma 1	Rb1	1.6
Salt-inducible kinase 1	Sik1	2.2
Sodium ion transport:		
Solute carrier family 13 (sodium/sulfate symporters), member 4	Slc13a4	2.2
Solute carrier family 22 (organic cation/carnitine transporter), member 5	Slc22a5	2.7
Solute carrier family 23 (nucleobase transporters), member 2	Slc23a2	2.0
Solute carrier family 34 (sodium phosphate), member 2	Slc34a2	5.6

Table 3.22 Some pathways represented by named up-regulated genes in ciprofibrate dosed rats for 3 hours. A list of pathways and the related genes are shown for up-regulated genes for male F-344NHsd rats aged 14-15 weeks dosed with 50 mg kg⁻¹ ciprofibrate then killed after 3 hours. RNAseq analysis was done as described in Section 2.2.6. Expressed genes with a fold change of >1.5 and a FDR p-value of <0.05 were listed.

Description	gene name	fold change
Solute carrier family 4 (anion exchanger), member 4	Slc4a4	1.8
Palmitoyl-CoA hydrolase activity:		
Acyl-CoA thioesterase 5	Acot5	2.8
Acyl-CoA thioesterase 6	Acot6	4.0
Acyl-CoA thioesterase 7	Acot7	1.6
Acyl-CoA synthetase long-chain family member 1	Acaa1	1.8
Acyl-CoA synthetase long-chain family member 3	Acs13	2.0
Acetyl-Coenzyme A acyltransferase 1	Acaa1	1.9
Oxidation reduction:		
Cytochrome P450, family 4, subfamily a, polypeptide 10	Cyp4a1	3.3
Cytochrome P450, family 4, subfamily a, polypeptide 2	Cyp4a2	2.3
Cytochrome P450, family 4, subfamily a, polypeptide 3	Cyp4a3	2.3
Cytochrome P450, family 4, subfamily b, polypeptide 1	Cyp4b1	1.8
Jumonji domain containing 1C	Jmjd1c	1.5

Section 3.5 Confirmation of results with real- time PCR

This section was set out to confirm the results found by the RNAseq and microarray analysis with real- time PCR by choosing some genes and measuring the levels of mRNA. Samples from male F-344 fisher rats treated with 50 mg kg⁻¹ ciprofibrate, 100 mg kg⁻¹ CPA and the control/ vehicle corn oil for 1, 3, 5 and 24 hours were analysed with real- time PCR as shown in Section 2.2.7. These samples were in 4 biological replicates and 2 technical replicates.

Primers and probes were designed for chosen genes. The genes used were *CYP4A1*, *CYP3A1*, *G0s2*, *Ccnd1*, *AhR*, β -*actin* and *Scd1*. *AhR* and β -*actin* genes were chosen as normalization genes as they were found to have a minimal induction change with the RNAseq results and they are also well known housekeeping genes regularly used for this purpose (Bazzi et al., 2009).

Cytochrome P450, family 4, subfamily a, polypeptide 1 (*CYP4A1*) was used as a positive control for the samples treated with ciprofibrate, as it is known to be induced by PPAR α agonists (Bell et al., 1991)(Bars et al., 1993)(Morris and Davila, 1996). The induction of *CYP4A1* was also noted in the microarray analysis for the 24 hour samples. Cytochrome P450, family 3, subfamily a, polypeptide 1 (*CYP3A1*) was used as a positive control for CPA treated samples, as this is known to be highly induced by PXR agonists (Lehmann et al., 1998)(Hosoe et al., 2005).

The other genes tested were G0/G1switch 2 (*G0s2*) and cyclin D1 (*Ccnd1*) as they are involved in cell cycle pathway. Stearoyl-Coenzyme A desaturase 1 (*Scd1*) is involved in the PPAR signalling pathway. As for the Aryl hydrocarbon receptor (*AhR*) it was found to be down- regulated in the ciprofibrate samples treated for 3 hours.

Section 3.5.1 Efficiency of primers and probes

To test the efficiency of the real-time PCR primers and probes, standard curves were produced for the *CYP4A1*, *CYP3A1*, *G0s2*, *Ccnd1*, *AhR*, β -actin and *Scd1* genes by amplifying a 5 fold serial dilution of cDNA prepared from the rat samples treated with corn oil. This was done in triplicate and the negative controls used were NTC, -RT and -RNA as explained in Section 2.2.7.2.

Figure 3.49 shows the Cycle-Threshold (Ct) (the cycle at which the fluorescence from the samples cross the threshold) against the log₁₀ cDNA concentrations for the genes *AhR*, *Scd1* and *CYP3A1*. Figure 3.50 shows the standard curves for the genes *Ccnd1*, *CYP4A1* and *G0s2*. While the standard curves for the gene β -actin are shown in Figure 3.51.

The amplification efficiency(%) and regression coefficient (r^2) of the standard curves for the genes are represented in Table 3.23. Amplification efficiencies of the standard curves for all seven genes (*CYP4A1*, *CYP3A1*, *G0s2*, *Ccnd1*, *AhR*, β -actin and *Scd1*) were within 9% of 100% and a regression coefficient of 0.996 ± 0.003 . This is ideal, as the amplification efficiency for PCR reactions should be $100\% \pm 10\%$, and the r^2 should be close to 1.

These results show that the genes and the method used, as described in Section 2.2.7, are reliable and can be applied to analyse the samples from male F-344 fisher rats treated with 50 mg kg^{-1} ciprofibrate, 100 mg kg^{-1} CPA and the control/vehicle corn oil for 1, 3, 5 and 24 hours.

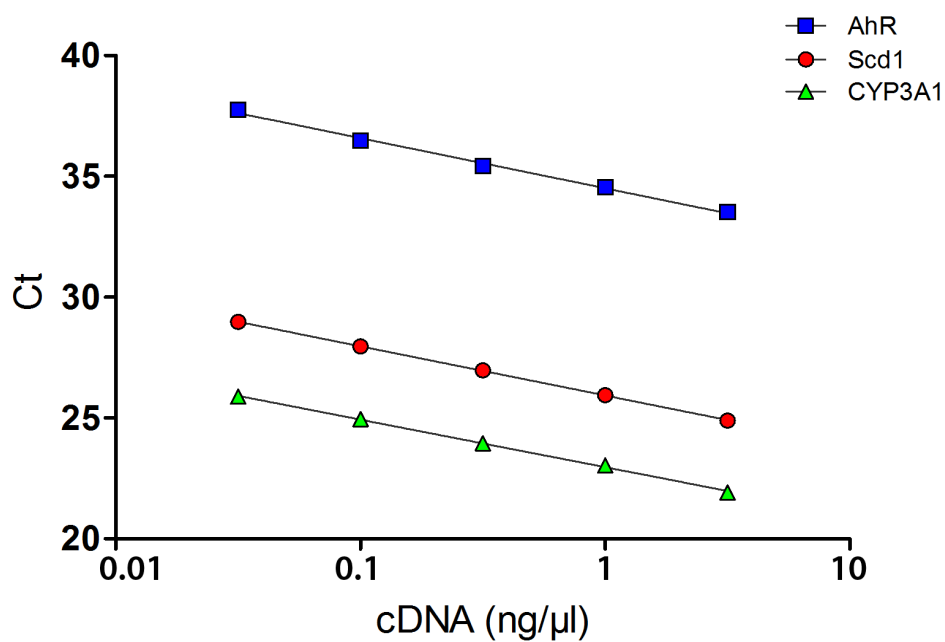


Figure 3.49 Amplification efficiencies for *AhR*, *Scd1* and *CYP3A1*. cDNA from rat hepatic RNA treated with corn oil for 30 hours was diluted 5 fold and amplified separately with each of *AhR*-with dye HEX (blue squares), *Scd1*-with dye HEX (red circles) and *CYP3A1*- with dye FAM (green triangles), then detected with real-time RT-PCR using TaqMan probe as described in Section 2.2.7. Each point represents the mean for triplicate samples. The efficiencies of each of the genes were obtained separately.

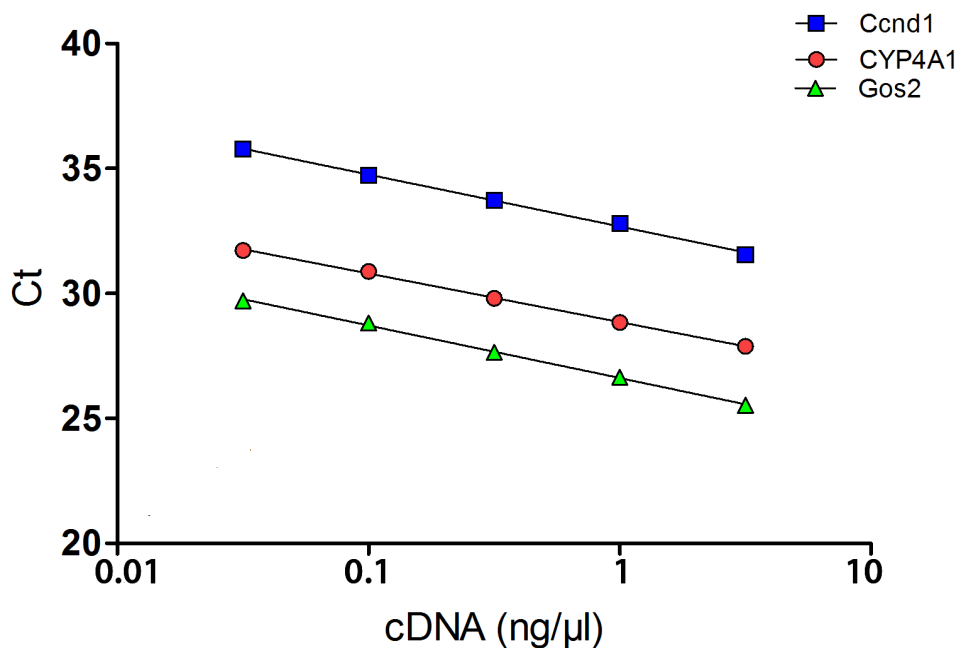


Figure 3.50 Amplification efficiencies for *Ccnd1*, *CYP4A1* and *Gos2*. cDNA from rat hepatic RNA treated with corn oil for 30 hours was diluted to 5 fold and amplified separately with each of *Ccnd1*-with dye HEX (blue squares), *CYP4A1*- with dye FAM (red circles) and *Gos2*- with dye At0647N (green triangles), then detected with real-time RT-PCR using TaqMan probe as described in Section 2.2.7. Each point represents the mean for triplicate samples. The efficiencies of each of the genes were obtained separately.

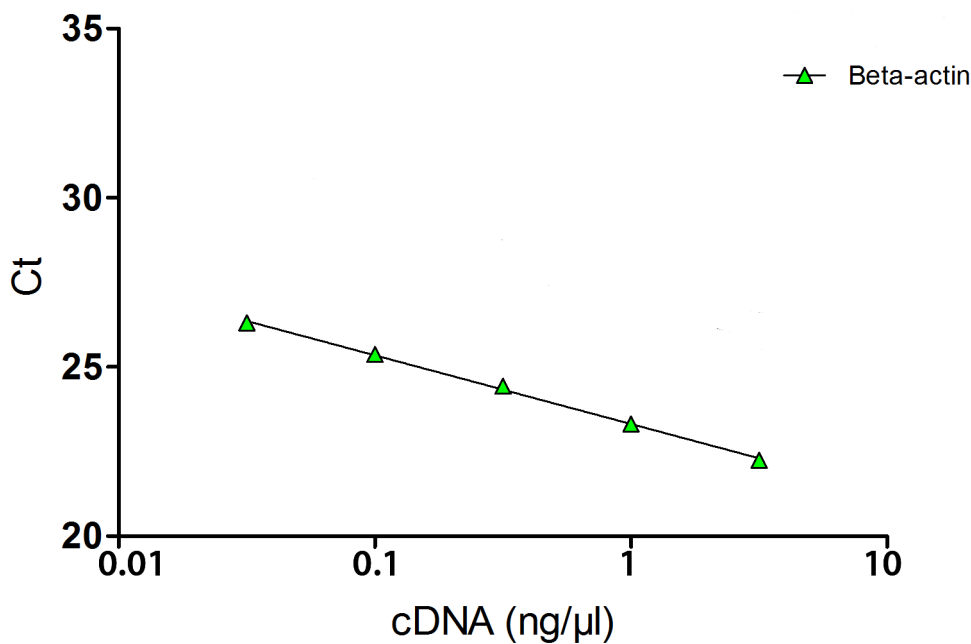


Figure 3.51 Amplification efficiencies for β -actin. cDNA from rat hepatic RNA treated with corn oil for 30 hours was diluted to 5 fold and amplified with β -actin- with dye Cy5 (green triangles), then detected with real-time RT-PCR using TaqMan probe as described in Section 2.2.7. Each point represents the mean for triplicate samples.

Table 3.23 Quantitation data of real- time RT-PCR for the genes used. Regression and efficiencies of CYP4A1, CYP3A1, G0s2, Ccnd1, AhR, beta actin and Scd1. The regression correlation coefficient value r^2 were determined from the equation line. The amplification efficiencies were derived as percentages and were calculated from the slope.

Gene	r^2	Efficiency%
<i>CYP3A1</i>	0.998	102.3
<i>CYP4A1</i>	0.997	104.1
<i>G0s2</i>	0.998	93.4
<i>Scd1</i>	1.0	97.7
<i>Ccnd1</i>	0.992	94.98
<i>AhR</i>	0.992	95.1
<i>β-actin</i>	0.998	97.9

Section 3.5.2 Real- time PCR analysis of the samples

The aim of this section was to measure the levels of mRNA of the genes *CYP4A1*, *CYP3A1*, *G0s2*, *Ccnd1* and *Scd1* with real- time PCR. To correct for loading differences, *AhR* and *β -actin* were both used as reference genes to normalise the measurements of mRNA.

F344 Fisher rats were treated with 50 mg kg⁻¹ ciprofibrate, 100 mg kg⁻¹ CPA or corn oil/vehicle for 1, 3, 5 and 24 hours. Hepatic RNA was extracted from frozen livers and reverse transcribed into cDNA as explained in Section 2.2.5.1 and. The cDNA was amplified with real- time PCR and the Ct levels were determined as shown in Section 2.2.7.3.

Section 3.5.2.1 Effects of ciprofibrate and CPA on *CYP4A1*

This experiment was done to test the induction levels of *CYP4A1* mRNA at different time points on ciprofibrate/CPA treated samples in comparison with controls, the *CYP4A1* was used as a positive control because it was known to be one of the first markers to the induction with PPA- $R\alpha$ agonists (Bell et al., 1991)(Bars et al., 1993)(Morris and Davila, 1996).

The results for the samples treated with ciprofibrate showed statistically significant induction of *CYP4A1* after 3, 5 and 24 hours of treatment, as shown in the time course in Figure 3.52 (red line). The samples from the 24 hour treatment showed a mean of ~30 fold induction higher than the control at the same time point. These results were consistent with the results from RNAseq analysis of the 3 hour samples treated with ciprofibrate which showed 3.2 fold induction of *CYP4A1*, while the real- time PCR gave a 2.4 fold induction at 3 hours (Section 3.4.6.2).

The induction of *CYP4A1* in fisher rats treated with CPA was not affected at all the time points as demonstrated as a blue line in Figure 3.52. Statistics was done with one-way analysis of variance (Bonferroni's multiple comparison test), $P < 0.05$.

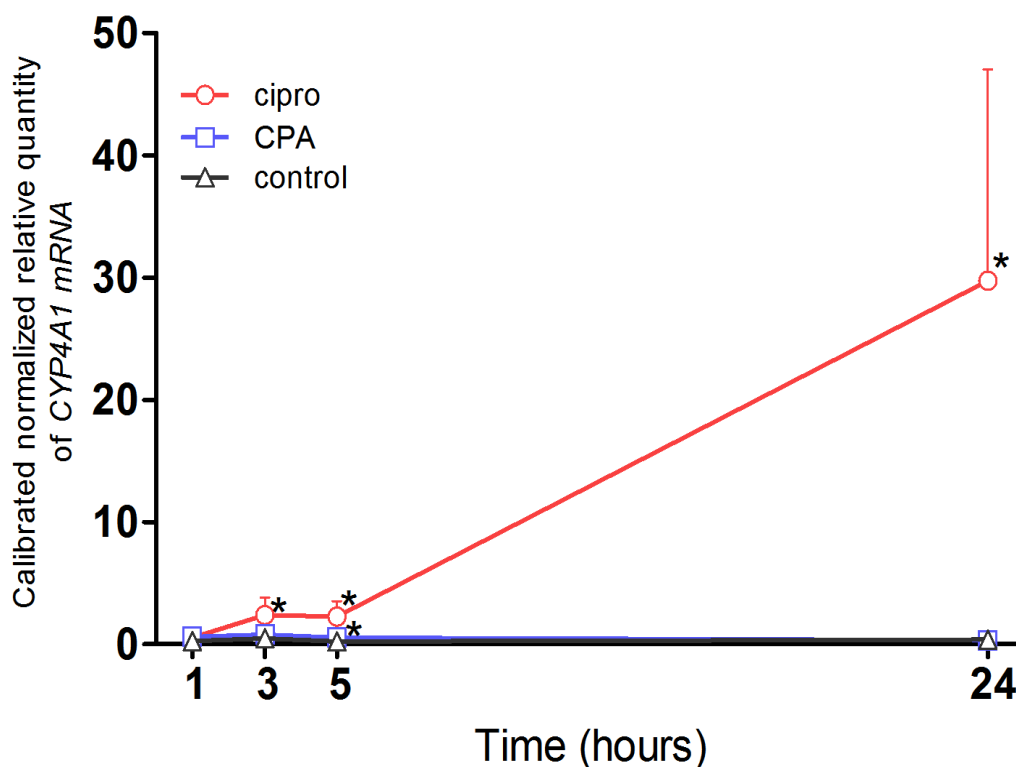


Figure 3.52 Effect of treatments on *CYP4A1* mRNA expression. The effects of treatments on *CYP4A1* mRNA expression, on cDNA from hepatic RNA of rats treated for 1, 3, 5 and 24 hours with 50 mg kg⁻¹ ciprofibrate (red circles), 100 mg kg⁻¹ CPA (blue squares) or corn oil (black triangles). Real-time PCR was performed as in Section 2.2.7.3 and analysed as in Section 2.2.7.4. Points are the mean value, and the error bars depict one standard deviation. Statistically significant difference from the control group was indicated by an asterisk. Statistics was done with one-way analysis of variance (Bonferroni's multiple comparison test), $P < 0.05$.

Section 3.5.2.2 Effects of ciprofibrate and CPA on *CYP3A1*

The objective of this section was to test the induction levels of *CYP3A1* in ciprofibrate and CPA treated samples in comparison with control samples at different time points. *CYP3A1* was used as a positive control for CPA treated samples, because the *CYP3A1* was reported to be highly induced by PXR agonists (Lehmann et al., 1998)(Hosoe et al., 2005).

The results for the samples treated with CPA showed a statistically significant induction of *CYP3A1* after 24 hours treatment; > 30 fold difference as shown in Figure 3.53/ blue line.

In contrast, the expression of *CYP3A1* in Fisher rats treated with ciprofibrate was not different from the control at any of the 1, 3, 5 and 24 hour samples.

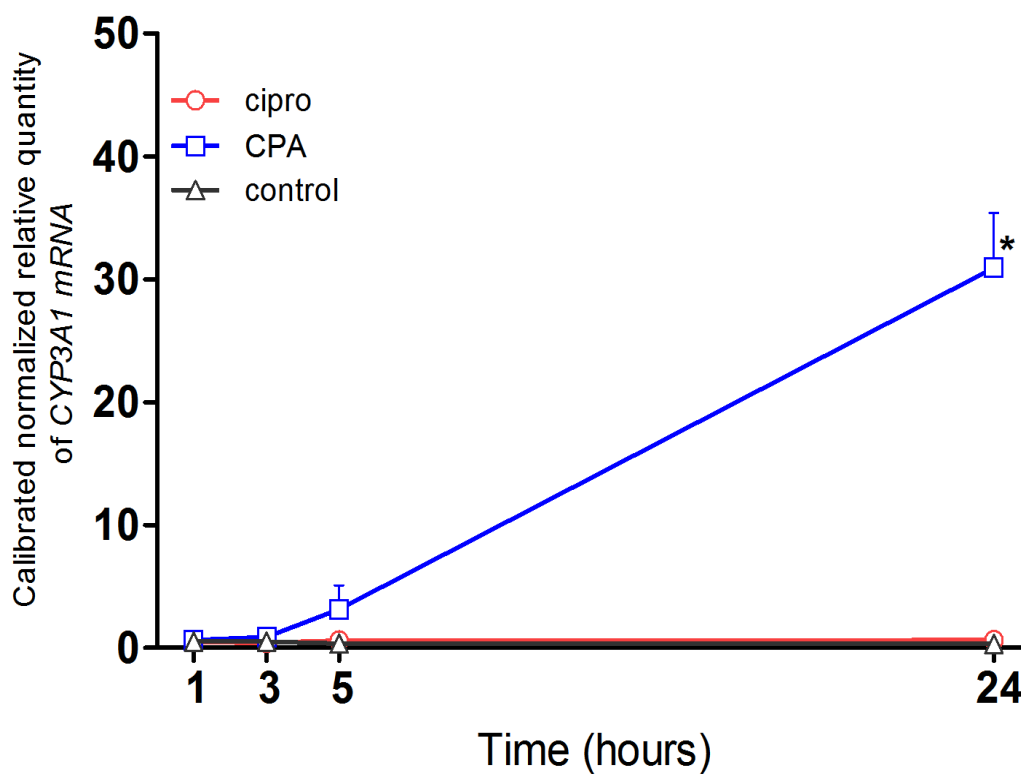


Figure 3.53 Effect of treatments on *CYP3A1* mRNA expression. The effects of treatments on *CYP3A1* mRNA expression, on cDNA from hepatic RNA of rats treated for 1, 3, 5 and 24 hours with 50 mg kg⁻¹ ciprofibrate (red circles), 100 mg kg⁻¹ CPA (blue squares) or corn oil (black triangles). Real-time PCR was performed as in Section 2.2.7.3 and analysed as in Section 2.2.7.4. Points are the mean value, and the error bars depict one standard deviation. Statistically significant difference from the control group is indicated by an asterisk. Statistics was done with one-way analysis of variance (Bonferroni's multiple comparison test), P<0.05.

Section 3.5.2.3 Effects of ciprofibrate and CPA on *G0s2*

This experiment was carried out to test the effects of ciprofibrate/CPA on the induction levels of G0/G1switch 2 (*G0s2*) mRNA (involved in cell cycle control), at the early time points and at 24 hours after treatment in comparison with the control samples.

Figure 3.54 shows the results for the samples treated with ciprofibrate (in red) where the expression of *G0s2* at 3 hours was significantly higher (5.4- fold) than the control at the same time point but declines by 24 hours. This result was similar to the results from RNAseq at 3 hours where the *G0s2* gene was found to be 5.8 fold higher than control samples (Section 3.4.6.2).

The induction of *G0s2* in fisher rats treated with CPA showed no significant difference from the control in all the samples treated at 1, 3 and 5 hours, as shown in Figure 3.54 blue line. Statistics was done with one-way analysis of variance (Bonferroni's multiple comparison test), $P < 0.05$.

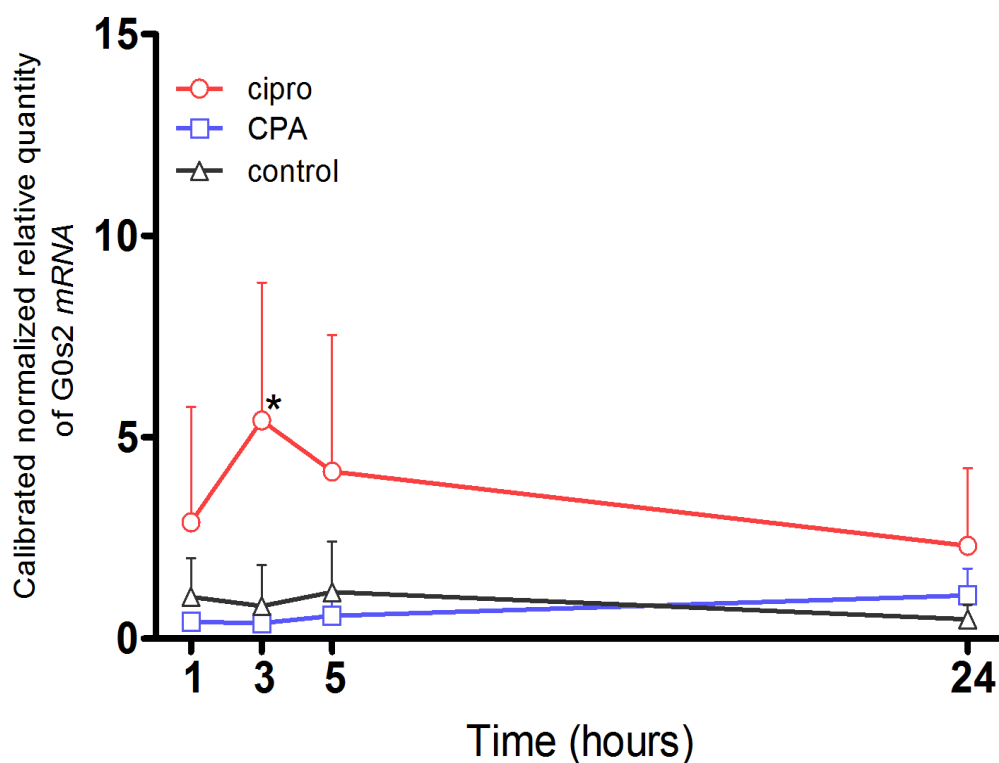


Figure 3.54 Effect of treatments on *G0s2* mRNA expression. The effects of treatments on *G0s2* mRNA expression, on cDNA from hepatic RNA of rats treated for 1, 3, 5 and 24 hours with 50 mg kg⁻¹ ciprofibrate (red circles), 100 mg kg⁻¹ CPA (blue squares) or corn oil (black triangles). Real-time PCR was performed as in Section 2.2.7.3 and analysed as in Section 2.2.7.4. Points are the mean value, and the error bars depict one standard deviation. Statistically significant difference from the control group is indicated by an asterisk. Statistics was done with one-way analysis of variance (Bonferroni's multiple comparison test), $P < 0.05$.

Section 3.5.2.4 Effects of ciprofibrate and CPA on *Ccnd1*

This part of the experiment was prepared to determine the effects of ciprofibrate and CPA on the induction levels of cyclin D1 (*Ccnd1*) (also involved in the cell cycle), with real- time PCR analysis.

The results for the samples treated with ciprofibrate, show a significant induction of *Ccnd1* at 3 hours (1.8 fold change) when compared with the control at the same time point (Figure 3.55/ red line). This result is comparable with the results from RNAseq at 3 hours where the *Ccnd1* gene was induced 2.9 fold (Section 3.4.6.2).

The induction of *Ccnd1* in Fisher rats treated with CPA is significantly higher than the control at 5 hours, while the 1, 3 and 24 hour samples treated with CPA have no significant difference when compared with the control as shown in (Figure 3.55).

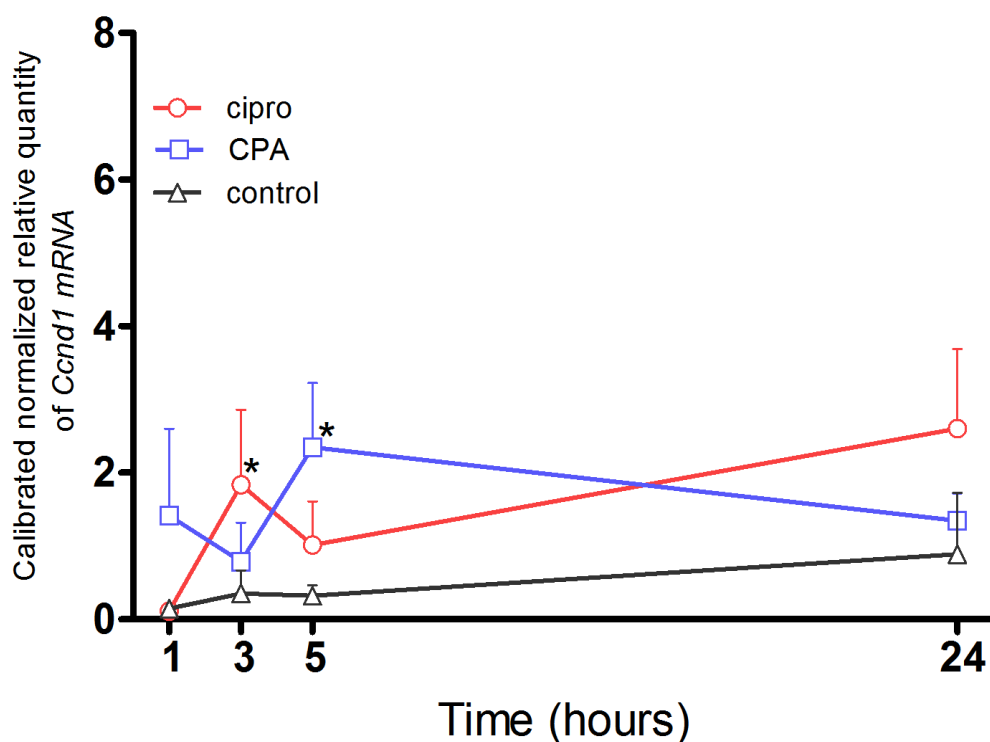


Figure 3.55 Effect of treatments on *Ccnd1* mRNA expression. The effects of treatments on *Ccnd1* mRNA expression, on cDNA from hepatic RNA of rats treated for 1, 3, 5 and 24 hours with 50 mg kg⁻¹ ciprofibrate (red circles), 100 mg kg⁻¹ CPA (blue squares) or corn oil (black triangles). Real-time PCR was performed as in Section 2.2.7.3 and analysed as in Section 2.2.7.4. Points are the mean value, and the error bars depict one standard deviation. Statistically significant difference from the control group is indicated by an asterisk. Statistics was done with one-way analysis of variance (Bonferroni's multiple comparison test), $P < 0.05$.

Section 3.5.2.5 Effects of ciprofibrate and CPA on *Scd1*

This experiment was done to test the induction levels of the *Scd1* gene expression, which encodes stearoyl-coenzyme A desaturase 1 (engaged in the PPAR α pathway) with real-time PCR. The results for the samples treated with ciprofibrate (Figure 3.56) showed a significant induction of *Scd1* at 1 hour, and at other time points *Scd1* expression in the ciprofibrate samples was slightly higher than in the control.

On the other hand the expression of *Scd1* in CPA- treated Fisher rats was lower than in the control albeit not significantly different (Figure 3.56).

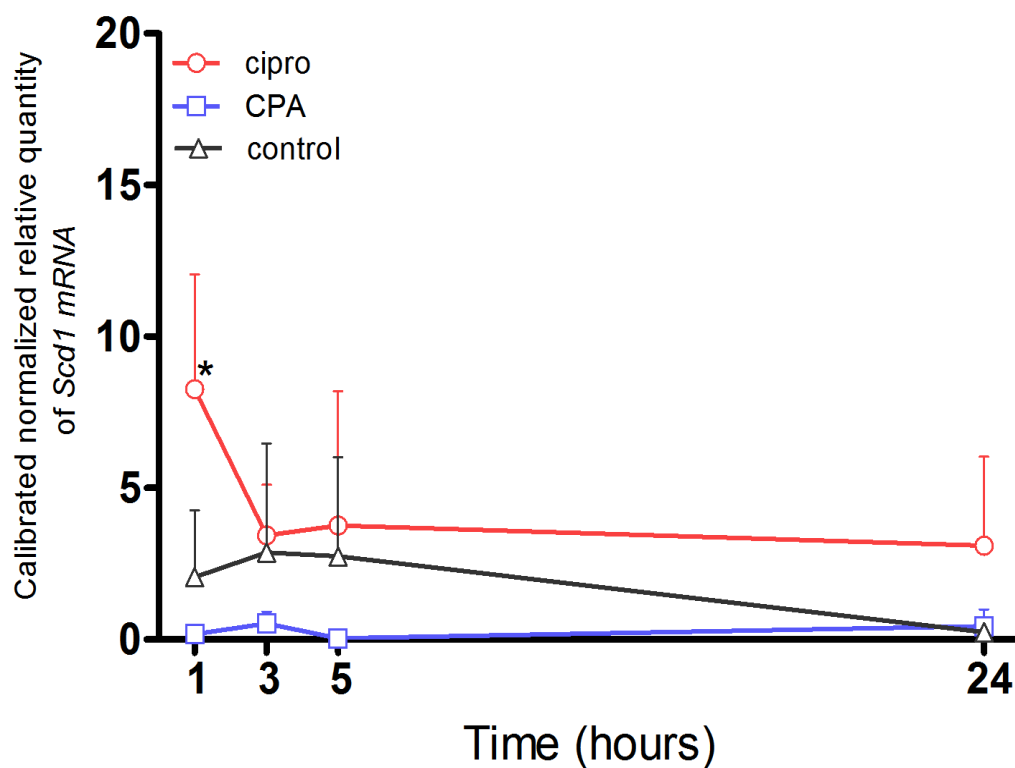


Figure 3.56 Effect of treatments on *Scd1* mRNA expression. The effects of treatments on *Scd1* mRNA expression, on cDNA from hepatic RNA of rats treated for 1, 3, 5 and 24 hours with 50 mg kg⁻¹ ciprofibrate (red circles), 100 mg kg⁻¹ CPA (blue squares) or corn oil (black triangles). Real-time PCR was performed as in Section 2.2.7.3 and analysed as in Section 2.2.7.4. Points are the mean value, and the error bars depict one standard deviation. Statistically significant difference from the control group is indicated by an asterisk. Statistics was done with one-way analysis of variance (Bonferroni's multiple comparison test), $P < 0.05$.

Chapter 4 Discussion

Section 4.1 Induction of hepatic DNA synthesis

The main goal of this thesis was to identify the genes induced by peroxisome proliferators (primarily ciprofibrate), which may be involved in the induction of hepatic DNA synthesis, and consequently, play an important role in hepato-carcinogenesis. First it was essential to demonstrate the effects of ciprofibrate on hepatic DNA synthesis.

Section 4.1.1 Effect of ciprofibrate on hepatic DNA synthesis in mice

To study the hepatic response to ciprofibrate in different strains of mice, two inbred strains of mice were chosen: C57BL/6JCrI which is a strain resistant to liver carcinogenesis and DBA/2JCrI which is fairly vulnerable to liver carcinogenesis (Diwan et al., 1986).

This was done with an immunohistochemical protocol, as has been used by many in the detection of proliferating cells *in vivo* with BrdU (deFazio et al., 1987)(Beyer et al., 2008)(Menegazzi et al., 1997)(Peters et al., 1997a). The validity of the immunohistochemical procedure in the detection of incorporated BrdU in the replicating hepatocytes has been broadly tested (e.g., (Eldridge et al., 1990) (Ledda-Columbano et al., 2003)).

The toxicity of BrdU to the DBA/2JCrI mouse strain was assessed by testing the gross effect on the animals' body weight, which showed there was no negative effect from the BrdU on the mice for the duration of the experiment (Section 3.1.1.2.1). This was consistent with results of (Al Kholafi, 2008).

Also, the results showed there was no toxic effect from the ciprofibrate indicated by the body weight study, and that the induction of hepatic DNA synthesis did not start until after 4 days of

exposure to ciprofibrate in DBA/2JCrI mice (Figure 3.5). This was substantively the same as previous results established by (Al Kholaiifi et al., 2008) with MCP in 129S4/SvJae mice, where he found that induction of hepatic DNA synthesis occurs 3- 4 days after dosing.

On the contrary, (Styles et al., 1990) showed that when dosing C57BL/6J mice with MCP, the induction of hepatic DNA synthesis started at 24 hours. This discrepancy could be the result of strain differences as this could affect liver function (Akiyama et al., 2001), although no previous evidence indicates that strain differences affect the response to peroxisome proliferators (Budroe et al., 1992).

Because of the above issues, the same experiment was repeated in C57BL/6JCrI mouse strain (Section 3.1.1.3), and this established that there were no toxic effects from the ciprofibrate or BrdU, and the induction of hepatic DNA synthesis did not start until after 4 days of exposure to ciprofibrate. The C57BL/6JCrI mice are inbred mice, so genetic variability is very small. Consequently, the difference between this study and the Styles study is less likely to relate to genetic differences. The present result was consistent with the findings for DBA/2JCrI mice (Section 3.1.1.2.2) AP mice and 129S4/SvJae mice (Al Kholaiifi et al., 2008).

The difference in the kinetics of induction of DNA synthesis between this and the Styles study could be due to the fact that this and the Al Kholaiifi studies (Al Kholaiifi et al., 2008) used an immunohistochemical detection of incorporated BrdU protocol, whereas Styles detected incorporated BrdU by isolation of hepatocytes and flow cytometry (Styles et al., 1987).

Miller (Miller et al., 1996) also tried to reproduce the same results that were from Styles but with the same flow cytometry procedure, and failed to produce the same results indicating that the results produced with the flow cytometry procedure are not fully reproducible.

As a conclusion, the induction of hepatic DNA synthesis was delayed until after 4 days dosing of ciprofibrate in the C57BL/6JCrI and the DBA/2JCrI mouse strain. This result demonstrates a confirmation of earlier studies on other strains with different peroxisome proliferators (Ledda-Columbano et al., 2003)(Al Kholaiifi et al., 2008).

Section 4.1.2 Effect of ciprofibrate on hepatic DNA synthesis in rats

The large time window of 4 days between the dosing of the ciprofibrate and the hepatic DNA synthesis makes it difficult to indicate when exactly the regulation of the genes responsible for the induction of DNA synthesis occurred. Thus, it was desirable to try an alternative system, and the rat liver was chosen.

It was known from previous *in vivo* and *in vitro* studies in Wistar and Fisher344 rats dosed with peroxisome proliferators (nafenopin and Wyeth-14,643) that the hepatic induction of DNA synthesis starts as early as 24 hours (Miller et al., 1996)(Menegazzi et al., 1997)(Al Kholaiifi, 2008) (Al Kholaiifi et al., 2008)(Bell and Elcombe, 1991a)(Bell et al., 1991).

In the current study a time course of the hepatic DNA synthesis revealed similar results. Hepatic DNA synthesis in male F-344/NHsd rats was induced by ciprofibrate within 10 to 30 hours, which peaks at 24 hours and again at 48 hours. This was consistent with (Plant et al., 1998) who demonstrated that PPAR α ligands cause rapid induction of DNA synthesis in rat hepatocytes *in vitro* (~24 hours). The second peak at 48 hours was probably related to the fact that the dose was given each day, meaning that the first peak would be a response to the first dose and the second peak would be a response to the second dose.

These results were similar to (Aboshofa's data) who also found two peaks of hepatic DNA synthesis at 24 and at 48 hours, using two labelling methods (BrdU and EdU). Aboshofa's data also

found that the induced hepatocytes at 24 hours are different from the hepatocytes induced at 48 hours (not the exact same hepatocyte). This was achieved by dosing the rats with ciprofibrate at 0 and 24 hours and dosing the animals with BrdU at 22 hours to label the proliferating cells at 22- 26 hours (as a response to the first ciprofibrate dose), and then giving the animals a dose of EdU at 46 hours to label the cells dividing at 46- 50 hours (as a response to the second ciprofibrate dose) (Aboshofa). After examining the hepatocytes with immunohistochemistry (to detect the BrdU labelled cells) and with fluorescent microscopy (to indicate the cells labelled with EdU), the images were merged together. Aboshofa found that the cells labelled with BrdU were different from the cells labelled with EdU, this suggests that the hepatocytes that divide at 24 hours are different from the hepatocytes that divide at 48 hours (Aboshofa personal communications).

Al Kholaiifi's (Al Kholaiifi et al., 2008) results with F-344/NHsd rats showed that hepatic DNA synthesis started to rise at 24 hours and continued after 48- 96 hours. This was consistent with the results found in this study (Figure 3.9). However, Al Kholaiifi used the chronic dosing system for BrdU labelling which gives the cumulative number of BrdU-labelled hepatocytes, whereas this study used the acute dosing system, showing what happened at each time point within a 2- 4 hour window.

It was desirable to determine the optimal dose of ciprofibrate in F-344/NHsd rats that gave the highest hepatic replicative DNA synthesis. Al Kholaiifi (Al Kholaiifi, 2008) used $50 \text{ mg kg}^{-1} \text{ day}^{-1}$ but did not test the effects of higher doses. This information was not available in the literature, so this study demonstrated (with a dose response experiment of 0, 50, 100, 200, 300 mg kg^{-1} body weight for 24 hours), that the liver labelling index increased significantly at 50, 100 and 200 mg kg^{-1} ciprofibrate, but the labelling index was low at 300 mg kg^{-1} . The reduced response at 300 mg kg^{-1} suggests a possible toxic reaction suppressing the liver growth response (Section

3.1.2.1.1). The concentration of 50 mg kg^{-1} was selected for the assay as an optimal dose of ciprofibrate for F-344NHsd rats, because it was deemed sufficiently low to avoid toxicity but gave a reasonably good labelling index.

(Woods et al., 2007) suggested that PPAR α ligands mediate chronic oxidative DNA damage, therefore, it was of importance to test the serum ALT concentrations as an indicator of liver cell damage. A single dose of $50\text{-}300 \text{ mg kg}^{-1}$ ciprofibrate did not increase serum ALT levels (Figure 3.11), indicating that there is no gross hepatotoxicity, and consequently no potential for regenerative DNA synthesis. Also the body weight of the animals was tested on a daily basis and this showed that there was no growth inhibition, indicating no toxic effects from the ciprofibrate. In addition the examination of histological sections revealed no necrosis in the liver tissue.

The immunohistochemical protocol yielded satisfactory staining of intestinal nuclei with BrdU as a positive control for labelling. Ciprofibrate-induced hepatocyte DNA synthesis was at its peak for rats at 24 hours after dosing, in contrast to the mouse strains where there was no significant induction until 4 days after treatment. These results demonstrate a species difference between mouse and rat in the kinetics of induction of hepatocyte DNA synthesis by PPAR α ligands.

The early induction of hepatic DNA synthesis in rat by PPAR α ligands unlocks doors of opportunity to study the mechanism of induction of the liver growth response, by linking early measurements of altered gene regulation to subsequent hepatic DNA synthesis.

Section 4.1.3 Effects of CPA on hepatic DNA synthesis in rats

The anti-androgen cyproterone acetate (CPA) is known to cause liver tumours in rats, and it

strongly enhances hepatic DNA synthesis of male and female Wistar rats (Schulte-Hermann et al., 1980), with it being more strongly inducing in female F-344 rats (Aboshofa). This drug was used to compare the genes that were induced with the CPA (PXR) with those that were induced by ciprofibrate (PPAR- α).

Although female rats gave higher induction than males from the preliminary tests, and from the literature (Topinka et al., 2004a) (Topinka et al., 2004b), it was reasonable to use male rats, as male rats had been used for the ciprofibrate dosing in this study. It might be expected that similar genes are induced in the male and the female rats, just more strongly in the latter (it would be interesting to follow up on this).

The doses used for the CPA (or PCN used in the preliminary experiment) were chosen from the literature (Topinka et al., 2004a) (Topinka et al., 2004b) and (Guzelian et al., 2006), at 100 mg kg⁻¹. The time was fixed to 24 hours so it would be comparable to the ciprofibrate experiment. The results showed a significantly high labelling index in the treated over the control group and over the groups treated with ciprofibrate at 24 hours (Figure 3.13B) and this was similar to results from previous studies (Topinka et al., 2004a).

Section 4.2 Zonal distribution of DNA synthesis in liver

The fact that the zonal distribution of DNA synthesis in the rat liver is known (Barrass et al., 1993) and that Barrass has established a reliable method to study this, gave reason to make similar studies on mouse liver, as there are no directly comparable data in mice (Burkhardt et al., 2004).

The main objective was to study the hepatic zonal distribution of cell proliferation in the liver of mice treated with ciprofibrate. Rats liver (used as a positive control) treated with ciprofibrate or CPA or corn oil (control) showed a significant zonal distribution of the labelled nuclei, with ~20 fold more labelled cells in the periportal region than in the perivenous region. This was in agreement with the literature (Barrass et al., 1993). Equally, the preferential periportal distribution of hepatic labelling in ciprofibrate treated rats was consistent over a dose range of 50-200 mg kg⁻¹ body weight (because of the low number of labelled cells in the 300 mg kg⁻¹ it was not possible to study the zonation in these slides.).

The zonal distribution of hepatic labelling index in male mouse showed no statistically significant difference between the periportal or perivenous zones in most of the mouse strains studied (1294S/SvJae wild type, PPAR α null male mice, C57BL/6JCrI and DBA/2JCrI), with the different doses of PPAR agonist, or choice of drug PPAR α (ciprofibrate, methylclofenapate) or CAR (TCPOBOP) (Section 3.2.2).

Exceptions were female 129S4/SvJae mice when treated with TCPOBOP as they showed a significantly perivenous preference (Figure 3.20), but the magnitude of the effect is relatively small, with the number of labelled cells in the perivenous region less than two times greater than those in the periportal region. Also AP mice when treated with MCP showed a similar zonal reaction to the rats, as the control and the group treated for 4 days had a periportal preference,

but for the group treated for 3 days the difference was not significant.

Of the total liver volume, 20% is made up of non-parenchymal cells, which are stellate cells, Kupffer cells, pit cells and endothelial cells (Oinonen and Lindros, 1998). These types of cells display modest quantitative zonal differentiation, in general being more abundant in the periportal area (Sasse, 1986). This could be related to the periportal/perivenous zonal behaviour noticed in the hepatic DNA induction in the liver population. It is observed furthermore that the periportal zonation of the hepatic labelling index in rat is different from the centrilobular induction of peroxisomal enzymes, cytochrome P450 and acyl CoA oxidase induced by PPAR α ligands, suggesting that the factors that mediate the induction mechanisms are located to specific zones of the liver lobule (Bell and Elcombe, 1991b)(Bars et al., 1993).

However, the induction of both enzymes and DNA synthesis are dependent on the PPAR α , this suggests that the zonation of the peroxisomal enzymes is via a PPAR α independent means (Al Kholaifi et al., 2008). Candidates that may intervene with these PPAR α -independent zonal effects are the zonal distribution of the PPAR α -associated microRNAs, or co-activators that are known to be required for the induction of hepatic DNA synthesis (Matsumoto et al., 2007) (Shah et al., 2007a).

Whereas it is acknowledged in mice that the *Apc* gene is the liver zonation keeper (Benhamouche et al., 2006), it is unknown if similar pathways apply for rat hepatocytes. Generally, however, the zonation of the hepatic induction of DNA synthesis is less studied, especially in mice.

These results illustrate species differences in zonation and kinetics of liver growth between rat and mouse, and predict species differences in zonation and timing of the regulatory factors responsible for the liver growth. These differences are similar in extent to those seen in extensive

zation studies on the induction of enzymes by various xenobiotics in isolated rat cells (Oinonen et al., 1994).

Bars (Bars et al., 1992) reported that the induction and expression of the cytochrome P4502B1/2 iso enzyme has a hepatic zonal pattern in the intact liver, whereas different reaction to PPAR α occurred *in vitro*, where they found that P4502B1/2 hepatic immunostaining was stronger in cells isolated from the perivenous liver region than in hepatocytes isolated from the periportal region. They also suggested that periportal and perivenous hepatocytes are assigned differentially to maintain regio- specific factors.

The diversity in inter species and inter strain response to the peroxisome proliferators could be most likely as a result of a difference in PPAR and/or PPRE structure, but this needs to be studied further and more data is required (Aldridge et al., 1995).

Section 4.3 Gene expression analysis

It has been acknowledged that the hepatic DNA synthesis induced by ciprofibrate in rat occurs after 24 hours (Section 3.1.2), and it is known that cells need ~19 hours to undergo hepatic DNA synthesis (Alberts et al., 2002). This implies that the RNA signal is increased within the first five hours after dosing. To determine the immediate early genes that may be induced the first 1-5 hours post induction were examined, as these genes could be responsible for the hepatic DNA synthesis at 24 hours.

This was done with cDNA microarray, RNA transcriptome sequencing and quantitative real-time PCR. Studies have shown that two or more different methodologies used to measure gene expression change have a tendency to agree when the magnitude of change in gene expression is large (Draghici et al., 2006).

cDNA microarrays can concurrently measure the expression intensity of thousands of genes within a specific mRNA sample (Schena et al., 1995)(Schena et al., 1998). The microarray analysis was applied to measure the expression levels of genes induced by 50 mg kg⁻¹ ciprofibrate in livers of F-344NHsd rats after 1, 3, 5 and 24 hours in comparison with the control.

The microarray analysis was carefully controlled with usage of: (1) vehicle/ corn oil as a negative control, (2) rats dosed with PXR ligands (CPA) as positive controls, to compare genes that are involved in hepatic DNA synthesis, but which might not be specific for the PPAR ligands (ciprofibrate), (3) the 24 hour samples were used to exclude the genes that are generally regulated by ciprofibrate but might not be specifically related to the early-immediate reaction that might be involved in the hepatic DNA synthesis, (4) a common reference/control (a sample containing an aliquot from each of the corn oil, CPA and ciprofibrate samples) was used which is the most commonly used design of microarray experiments, for a complete comparison of sam-

ples and to remove any bias from the 555 and 647 Alexa dyes (Cherkaoui-Malki et al., 2001).

The cDNA microarray analysis was optimized and each step controlled and assessed, beginning with the RNA quality and quantity measurements (Section 3.3.1). The incorporation of Alexa dyes 555 and 647 in the cDNA were also made with a negative control (sample with no RNA) and a positive control (RNA supplied in the kit).

Section 4.3.1 Optimization of the microarray technique

Optimization of the microarray technique was essential to ensure the results are reliable and repeatable. So, the following issues were addressed: (1) reproducibility of the scanning, (2) reproducibility of hybridization, and (3) the testing of different concentrations of cDNA. The reproducibility of scanning (60- 90%) was good (Section 3.3.3.1). However, initial experiments on the reproducibility of hybridization were not as successful, since the same cDNA when hybridized on three different sets of slides did not give repeatable results. This was a problem, and results from these hybridizations were not reliable. Therefore, it was essential to address this issue before being able to consider using the microarray technique for further analysis (Section 3.3.3.2). Subsequently, by considering all the microarray procedures systematically, it was revealed that the gene names were misplaced on the gene spots of the slides (human error) when placing the features from the GAL file on the slides. This issue was resolved by using slides with markers indicating where the genes were situated.

It was also essential that all the hybridizations were replicated at least twice with data giving a minimum of 60% similarity to be considered in the analysis. A 60% similarity in the reproducibility of the technical replicates was proposed acceptable in other studies, and it has been reported that the correlation coefficient can range between 0.5 and 0.95 (this corresponds to 50 - 95%) (Bammler et al., 2005)(Jarvinen et al., 2004)(Jenssen et al., 2002)(Carter et al, 2005).

This variability issue with microarrays could be related to the fact that they do not measure with great accuracy the genes expressed at low levels (Tarca et al., 2006). Also fluorescence from the extremely highly induced genes may surpass the saturation limit of the detector. These factors may have contributed to the differences evident in this study between data from the microarray analysis and the transcriptome sequencing, as the latter would detect all the genes.

From this study the measures used to improve the quality of microarray results were to use four biological replicates and two technical replicates with the technical replicates being accepted only where $r^2 > 0.6$.

Section 4.3.2 Microarray analysis of rats dosed with ciprofibrate

The 24 hour samples were initially tested to measure the expression levels of genes associated with induction of hepatic DNA synthesis, because this time point has been thoroughly investigated in the literature and a large number of genes have been detected to be highly induced. This was to ensure that the analysis and processing of the data were acceptable and reliable, and that specific genes used as markers were discovered.

Microarray analysis of rats dosed with ciprofibrate after 24 hours revealed 1597 significantly changed genes. In the samples treated with ciprofibrate for 24 hours 39 genes were up-regulated by double or more and 23 genes down-regulated by half or less. Up-regulated genes included *Cyp4a1*, *Cte1*, *Rab19*, *Slc9a2*, *18A*, *Ldh3*, *Narg1* and *Polr2a*. Genes recently reported to be up-regulated after 14 days of orally repeated dosing of WY-14,643 were *Apex1*, *Xrcc5*, *Gadd45* and *Aih1* (Suzuki et al., 2010). Down-regulated genes included *Mup2*, *Zfp598*, *Fgg*, *Cyp2c29* and *Sepina3a* (a list of all the genes are found in a CD at the end of the Thesis).

Cyp4A1 is a known marker for PPAR α agonists (Morris and Davila, 1996)(Hardwick et al.,

1987)(Bell and Elcombe, 1991a)(Bell and Elcombe, 1991b)(Orton and Parker, 1982)(Baker et al., 2004). The induction of *CYP4A1* via the PPAR α receptor has been reported in hepatocytes both *in vitro* (Yaacob et al., 1997) and *in vivo* (Correia, 1995)(Simpson, 1997). Furthermore, clofibrate exposure induced *CYP4A1* with both RNA microarrays and quantitative real-time PCR (Baker et al., 2004). The induction of *CYP4A1* with ciprofibrate here, while not proof, supported the validity of the present microarray analyses.

The microarray analysis of ciprofibrate dosed F-344NHsd fisher rats after 1, 3, and 5 hours showed 1377 significant genes analysed for each of the time points. Some of the up-regulated genes were *Fxyd2*, *Igf1*, *cpa5*, *Cd68*, *Abcc2*, *Mup1*, *Cmklr1*, *mmu* and *Ndrgr1* for the 1 hour samples, while *Psemb9*, *Inpp11*, *Saa3*, *Serpina1b*, *Ins1* and *Gsta1* were up-regulated for the 3 hour samples. The genes *Gc*, *Apob*, *Dusp14* *Mug1*, *Serpina 1b*, *Kcnh1*, *Apcs* and *Gcap15* were up-regulated in the 5 hour samples. These are all novelty findings for the immediate early genes. Later genes have been reported to be induced by clofibrate after 3 days of treatment: *Hcd*, *Cyp4a1*, *3kcta*, *3kctb*, *ACox*, *Cyp17* and *Adh1*(Baker et al., 2004).

Down-regulated genes can be just as informative as up-regulated genes when studying the mechanism of action of liver growth induced by ciprofibrate. After 1 hour treatment with ciprofibrate a number of the down-regulated genes were *Fgf21*, *Rgs12*, *Usp28*, *Dars*, *Vnn3* and *Fzd7*, some of the down-regulated genes after 3 hours were *Pfkfb2*, *Wdr10*, *Clecsf9*, *Kcnq2*, *Mylc2pl* and *Rgs12*, while genes down-regulated at 5 hours were *Nedd4*, *Rem1*, *Adcy8*, *Rab40c*, *Acadsb* and *Ccb3*. Other genes that have been reported to be down-regulated after 24 hours of dosing with ciprofibrate are *Apo-A1*, *CYP1A2*, *CPT1* and tumour necrosis factor receptor 1 (Baker et al., 2004).

Section 4.3.3 Transcriptome sequencing of rats dosed with ciprofibrate for 3 hours

Transcriptome sequencing analysis uses high- efficiency high- throughput sequencing to measure gene expression (Marioni et al., 2008). Transcriptome sequencing analysis was used to help find differently expressed genes in hepatic cells of F344/NHsd Fisher rats dosed with 50 mg kg⁻¹ ciprofibrate for 3 hours. A key advantage of the transcriptome sequencing analysis procedure is the number of genes that could be detected in this study, at just under 25000 genes. This was because the transcriptome analysis was sincere and was used against a complete rat genome data base. The microarray analysis had only 5826 genes because a large number of genes were missed when spreading the cDNA on the slides, or the cDNA does not hybridize well enough to be able to be scanned, or just washes off at the washing stages, also the microarray slides were specified for mice, so only the rat genes that match the mouse genome are included.

The different outcomes from the microarray results and the transcriptome sequencing results and real-time PCR measurements are expected, and could reflect the fact that the microarray analysis excluded the extremely high induced genes and the very low expressed genes (Tarca et al., 2006) (Gerhold et al., 2001). While the transcriptome sequencing data were more inclusive, and real- time PCR detects smaller changes in gene expressions, that are undetectable with microarrays (Baker et al., 2004). Hence the recommendation that microarrays are used to screen thousands of genes, whereas quantitative real-time PCR could be used to validate observations with chosen genes of interest (Baker et al., 2004).

The transcriptional sequencing was carefully monitored, with the samples confirmed to be assigned to their original groups and the hierarchical and k-means clustering showing that the control and treatment groups were clustered together.

Up-regulated genes included *CYP4A1*, *G0s2* (*G0/G1 switch 2*), *Aadacl1*, *Acot5+6* (*Acyl-CoA thioesterase 5+6*) *Acpp*, *Apol9a*, *Ccnd1*, *Epha2*, *Prrg4*, *Myc11*, *Etv6* and *Scd1*, and the down-regulated known/ named genes were *Igfbp2*, *ppargcla*, *Cyp1a2*, *AhR*, *Acsm2*, *Cldn9 +4 +6*, *Hps4* and *Rps14*. These were similar to results found by Peters who revealed that the *Aco*, and *c-myc* alongside other genes were up-regulated when mice were dosed with Wy-14,643 for 5 weeks and 11 months (Peters et al., 1997a). He also concluded that the peroxisome proliferators contribute to the hepato-carcinogenicity effects.

Section 4.3.4 Ciprofibrate pathways of over-represented genes

The *acute phase response* pathway was over-represented among the functions of up-regulated genes in both the 3 and 5 hour groups treated with ciprofibrate and analysed with microarray analysis. At 3 hours 4 up-regulated genes out of 23 (of all genes in the genome) that are involved in the *acute phase response* (17.4%) and 8 out of 23 genes (34.8%) in the 5 hour group. The *acute inflammatory response* and the *wounding response* were over-represented at 3 and 5 hours post-treatment. These results were not unexpected, as these responses, and especially the *acute phase response*, are a complex chain of reactions that are known to be the initial response to infections, or malignancy, which prevent further damage to the tissue (Baumann and Gauldie, 1994).

Hierarchical clustering was also used in these experiments to clarify different pathways that were over-represented, among up-regulated genes after ciprofibrate treatment for 24 hours. These functional categories included the *fatty acid metabolic process*, the *monocarboxylic acid metabolic process*, the *long chain fatty acid metabolic process* (100%), *fatty acid β oxidation* and *acyl-CoA metabolic process*. Among the changes documented to occur during liver growth after treatment with peroxisome proliferators in rats are β -oxidation of fatty acids and very long-

chain fatty acid β -oxidation (Latruffe et al., 2001)(Hamadeh et al., 2002)(Schoonjans et al., 1996). Therefore, similar results were revealed in this study.

Genes known to be involved in the response to peroxisome proliferators were also up- regulated, including genes responsible for liver fatty acid binding protein and acyl-CoA and growth regulatory genes (*c-myc*, *c-Ha-ras*, *fos*, *jun*, and *egr-1*) (Corton et al., 2000).

Over-represented pathways among up- regulated genes from the transcriptome sequencing data included those with relationships to DNA synthesis, cell cycle or PPAR signalling: The *PPAR signalling pathway*, the *fatty acid metabolic pathway* and *cell cycle* regulatory genes. Other smaller groups of genes (5-6 genes) were involved in *sodium ion transport*, *oxidation reduction* and *palmitoyl-CoA hydrolase activity*. Other pathways were the *metabolic process* and *bioactivation pathway via cytochrome P450*.

Others reported that in rodents treated acutely with Wy-14,643 a number of the significantly induced genes were *Il1 β* , *Il1r1*, *Hnf4* and *Stat3*, and when they were treated chronically the genes that had increased expression were *Il1 β* , *Il1r1*, *Il6* and *Ppar γ* (Anderson et al., 2001). With nullizygous mice for TNF-receptor I, II or both, they also demonstrated that the hepatic carcinogenesis caused by peroxisome proliferators is not mediated through TNF α .

A number of the genes identified here were followed up with the quantitative real- time PCR. Chosen pathways were tested in a time course for the ciprofibrate and CPA treatments. These pathways were the *DNA synthesis*, the *cell cycle pathway* and *PPAR signalling pathway* genes as they are related to the hepatic DNA synthesis and so to the liver growth mechanism. The genes were G0/G1 switch 2 (*G0s2*), cyclin D1 (*Ccnd1*), stearoyl-coenzyme A desaturase 1 (*Scd1*), Cytochrome P450, family 4, subfamily a, polypeptide 1 member *CYP4A1* and Cytochrome P450, family 3, subfamily a, polypeptide 1 member *CYP3A1*.

Section 4.3.5 Microarray analysis of rats dosed with CPA

CPA is an artificial progesterone analogue with progestogenic and antiandrogenic activities, used in the treatment of prostate cancer to resolve androgenisation symptoms in females and to reduce the male hormone testosterone to suppress the sexual drive in males (Neumann, 1994). In rat liver, CPA has shown evidence of genotoxic activity, promoting specific DNA adducts and mutagenesis (Krebs et al., 1998).

Microarray analysis of dosed rats with CPA after 24 hours resulted in 1597 significantly changed genes (up and down- regulated), the 24 hour group was initially studied to specify genes that are known to be induced at this time with PXR ligands, to act as a control for the analysis and processing of the data, and to be able to relate this to the experimental system. This was done by finding specific genes that were used as markers, *CYP3A1* (Slatter et al., 2006) or *Gstm3 (GstYb4)* (Krebs et al., 1998).

Results from rats treated with 100 mg kg⁻¹ CPA for 24 hours resulted in 21 up- regulated genes and 10 down- regulated genes with Student's t- test. Some of these up- regulated genes included *Gsta1*, *Hspa8*, *Gstm3 (GstYb4)*, *Ugt1a5*, *Apoa1*, *Mup3*, *Ef1A1* and *Rps6*, and down- regulated genes included are *Mup2*, *Rnase4*, *Trf* and *CYP2d26* (a list of all the genes are found in a CD at the end of the Thesis).

Although *CYP3A1* was not found in the initial 24 hour analysis, it is a well known marker for PXR agonists in rats (Slatter et al., 2006), this could be related to the fact that the *CYP3A1* is highly induced at 24 hours and the microarray analysis (as reported earlier) can lose the ability to detect the extremely high induced genes and the very low expressed genes (Tarca et al., 2006)(Gerhold et al., 2001). On the other hand, *Gstm3 (GstYb4)* was found, and this gene was also widely reported in the literature to be induced after 24 hours of dosing with CPA, and could

also be used as a marker (Slatter et al, 2006)(Krebs et al., 1998).

The microarray analysis of CPA dosed rats at the immediate early hours at 1, 3, and 5 hours, with 1377 significantly up/ down- regulated genes. Some of the up- regulated genes at 1 hour were *Sstr4*, *Cd97*, *4930477O03Rik*, *E2f3*, *Sirt2*, *Nrxn3*, *Trpc4ap* and *Rnf141*. A number of the 3 hour up- regulated genes were *Hpxn*, *Serpina3m*, *Cops4*, *E130101E03Rik*, *Cyp2c29* and *A1481316*. While the genes up- regulated in the 5 hour group included *mt*, *Mug1*, *Fabp1*, *Scd1*, *Adamts1*, *Gc*, *Eef1a1* *Fzd7* and *Cyp2c29*, other reported genes in rats and mice induced after 3 daily doses of PXR are *Gstm3*, *Abcd3*, *Abcc2*, *Amtl*, *CYP51*, *Foxa1*, *Fmo2*, *Gsta4*, *Gstt2*, *Nr2f2*, *Slc16a6* (Slatter et al., 2006). Shah used qPCR to find that a number of genes were up-regulated in guts of mice treated with PXR, *Gsta1*, *Gstm1*, *Gstt1*, and *Mdr1a* (Shah et al., 2007b). From this it is noticeable that the *Gst*'s and *Gstm*'s are all up- regulated in this study and in a number of previous studies.

While the down- regulated genes in the 1 hour group included *Pla2g2c*, *Card10*, *Apom*, *Sfrs16*, *Rem1*, *Rac1* and *Bmp7*, the down- regulated genes in the 3 hour group were *Axot*, *Alas1*, *Ugt1a13*, *Edem1* and *Sif1*, and the down- regulated genes in the group treated for 5 hours included *Acadsb*, *Dbil5*, *Sfrs16*, *Inpp1l*, *Cental* and *Gpx1*. Also Shah found that PCN (a PXR) inhibited TNF α activated NF kB luciferase reporter in the colon (Shah et al., 2007b) and in rat liver (Menegazzi et al., 1997).

Section 4.3.6 CPA pathways of over-represented genes

The *glutathione transferase activity pathway* was over-represented among the functions of up-regulated genes when the rat livers were treated with CPA. These results were similar to the findings of Slatter (Slatter et al., 2006). The process *interphase of mitotic cell cycle* was also over-represented among up-regulated genes along with the *phospholipid metabolic process* and

the *steroid metabolic process*, as also found by (Slatter et al., 2006).

According to a heatmap from the 3 hour samples treated with CPA, *immune responses*, *a response to chemical stimulus*, and *a response to organic substance* and other pathways were over-represented. Unfortunately, there was not enough data from the analysis to perform a heatmap analysis for the 1 and 5 hour time points. Previous studies have illustrated that *citrate cycle*, *pyruvate metabolism*, *fatty acid metabolism* and *bile acid biosynthesis* were over-represented among the data for PXR-treated cells suggesting that the PXR regulates bile and lipid acid metabolism (Slatter et al., 2006).

Section 4.3.7 Quantifying genes with real-time PCR

To confirm and quantify genes of interest found by the microarray and the RNAseq analysis, real-time PCR was applied. The choice of the genes analysed was based largely on the RNAseq results: *CYP4A1* was used as a positive control for ciprofibrate because it is known to be induced by PPAR α agonists (Bell et al., 1991)(Bars et al., 1993)(Morris and Davila, 1996). *CYP3A1* was used as a positive control for CPA as this is known to be highly induced by PXR agonists (Lehmann et al., 1998)(Hosoe et al., 2005), and *G0s2*, *Ccnd1*, and *Scd1* were also detected as genes of interest, to check when their expression is increased and if the ciprofibrate induces similar responses in these genes in comparison to the CPA samples.

Validation of the normalization genes (*AhR* and β -*actin* genes) was essential to normalise the measurements of mRNA to correct for loading differences, and was performed in each of the experiments. This was done to exclude any sample differences attributable to RNA quality or quantity. *AhR* and β -*actin* genes were found to have a minimal change according to the literature and they are well known housekeeping genes regularly used for this purpose (Bazzi et al., 2009) (Lee et al., 1995). The normalization / housekeeping gene expression is not always stable across

the treatments (Pohjanvirta et al., 2006), and it is optimal practice to use more than one such normalization/ reference genes (Vandesompele et al., 2002). Thus, two reference genes were used in this study.

Section 4.3.7.1 Effects of ciprofibrate and CPA on *CYP4A1*

The rat liver *CYP4A1* is induced in rodent livers treated with peroxisome proliferators (Hardwick et al., 1987). This rapid induction is usually followed by liver hyperplasia and hypertrophy (Bell et al., 1991), as liver enlargement of the treated rats was also found in this study.

The induction levels of *CYP4A1* mRNA at different time points on ciprofibrate or CPA treated samples in comparison with controls were measured. This generated a statistically significant high induction of *CYP4A1* after 3, 5 and 24 hours of ciprofibrate treatment with the 24 hour treatment giving an induction of ~30- fold. The 3 hour ciprofibrate samples gave results with real- time PCR consistent with the RNAseq analysis, ~2.4 fold induction and ~3.2 fold- induction, respectively. As for the microarray data it appears to be less sensitive and unreliable (in particular at early hours), as the *CYP4A1* did not correlate with the RT-PCR or the RNAseq, and the genes were not consistent between the time points as expected, also no positive controls were found in the early time points for the treatments. The literature reports *CYP4A1* to be highly induced with PPAR α agonists *in vivo* and *in vitro* (Bell et al., 1991)(Bars et al., 1993)(Morris and Davila, 1996).

The high induction of *CYP4A1* that has been seen here and the pathways that have been highlighted earlier in Section 4.3.4 correlate with results that state that induction of *CYP4A1* happens at the same time as other genes which are involved in the peroxisomal β oxidation pathway (Furuta et al., 1982). It has been reported that the *CYP4A1* is involved in the metabolism of a range of xenobiotics and in the hydroxylation of fatty acids (Aoyama et al., 1990)(Gonzalez and

Nebert, 1990). The induction of *CYP4A1* expression in the fisher rats treated with CPA was not affected in the 1, 3 and 24 hour samples while it was significantly lower than the control at 5 hours.

Section 4.3.7.2 Effects of ciprofibrate and CPA on *CYP3A1*

The quantitative real- time PCR results for the CPA treated samples showed a statistically significant > 30- fold induction of *CYP3A1* after 24 hours. These results were consistent with findings in the literature that found that *CYP3A1* was highly induced by PXR agonists (Lehmann et al., 1998)(Hosoe et al., 2005).

It has been reported that PXR mediates the genomic effects of a number of steroid hormones and xenobiotics via the *CYP3A* gene family (Masuyama et al., 2001), and the difference in the response to PXR between the males and female rodents could not be overlooked, with the latter having a higher induction of *CYP3A1* (Eliasson et al., 1994)(Larsen and Jefcoate, 1995). PXR activates *CYP3As* specifically in hepatic cells because of the need for a hepatic nuclear factor-4 α (HNF4 α) which apparently is involved in the transcriptional activation of the *CYP3As* (Tirrona et al., 2003). As for the induction of *CYP3A1* in ciprofibrate treated fisher rats, there was no difference from the control at any of the time points.

Section 4.3.7.3 Effects of ciprofibrate and CPA on *Ccnd1*

Cyclin D1 is involved in the cell cycle and was known to be induced after partial hepatectomy (Boylan and Gruppuso, 2005)(Michalopoulos, 2007). Cyclin D1 is considered an important intracellular mediator for the mitogenic signals that are responsible for hepatocyte proliferation in the regenerating liver (Nelsen et al., 2001). It is also a delayed target of the c-Jun-N-terminal kinase pathway during liver regeneration (Schwabe et al., 2003). Other studies demonstrate that

the over expression of the cyclin D1 gene results in fast growth of a subset of hepatocellular carcinomas (Nishida et al., 1994).

Real- time PCR analysis of *Ccnd1* showed an up- regulation with ciprofibrate treated samples at 3 hours (1.8 fold change) and this was similar to the corresponding RNAseq analysis (2.9 fold change). These results were similar to previous studies that showed a high induction of cyclin D1 mRNA in Wy14643 treated mice (Peters et al., 1998).

Ccnd1 was induced at 5 hours in CPA treated Fisher rats but not at 1, 3 or 24 hour samples. This demonstrates that the *Ccnd1* could be affected by both the ciprofibrate and the CPA at the immediate early time points.

These results demonstrate that *Ccnd1* might be a common target for the hepatic DNA synthesis, and is not specific for the peroxisome proliferators nor the PXR. This was also suggested by Pibiri, who found the *Ccnd1* was highly induced at the early hours after a single dose with the thyroid hormone (T3), and also suggested that *Ccnd1* could be responsible for the mitogenic activity of nuclear receptor ligands (Pibiri et al., 2001).

Therefore, conclusions could be drawn in relation to the effects of the *Ccnd1* on the hepatic DNA synthesis, as it is induced after 3 hours presumably stimulating the DNA to enter the cell cycle and so cause the hepatic DNA synthesis that is observed after 24 hours. However, this phenomenon is noticed additionally with the CPA signifying that the *Ccnd1* is not specific to the PPAR α and that the PXR have the same effects on the *Ccnd1*.

Section 4.3.7.4 Effects of ciprofibrate and CPA on *Scd1*

It has been shown previously that poly-unsaturated fatty-acids like linoleic acid repress the expression of *Scd* gene encoding (stearoyl-coenzyme A desaturase 1) in adipocytes (Sessler et al.,

1996). When mice were fed a high-carbohydrate fat free diet, *Scd1* was induced around 50-fold, which could have been caused by carbohydrate or insulin (Waters and Ntambi, 1994)(Ntambi, 1995).

In the present study ciprofibrate significantly induced *Scd1* at 1 hour (~9 fold change according to real-timePCR, ~13- fold change according to RNAseq analysis) and at the other time points *Scd1* expression in ciprofibrate samples were slightly higher than the control samples, while *Scd1* expression in CPA treated rats was slightly lower than the control at all the time points, though this difference was not significant. Research done by Miller showed that peroxisome proliferators induced *Scd1* in mouse liver (Miller and Ntambi, 1996).

This is interesting as it suggests that the signals from the *Scd1*, occurring as soon as 1 hour after dosing with the ciprofibrate, stimulates the hepatocytes to divide. The hepatic DNA synthesis could be seen after 24 hours with the immunohistochemical protocol. Apparently the induction just stops within two hours, and this is reflected on the hepatic DNA synthesis as the cells slow down their dividing within a couple of hours after the 24 hours peak. Furthermore, a noteworthy reality is that the CPA is not involved in this pathway as it is specific for the PPAR α , which is as confirmed by the fact that *Scd1* expression is not induced by CPA.

Section 4.3.7.5 Effects of ciprofibrate and CPA on *G0s2*

The *G0s2* gene was identified by chance during a screen to find differentially expressed genes with links to lectin treatments of lymphocytes, and is involved with cell cycle development from the G0 to the G1 stage (Cristillo et al., 1997). It was also considered that *G0s2* is a target gene of the all-trans-retinoic acid (this is an oxidized form of vitamin A used in the treatment of acute leukemia) (Kitareewan et al., 2008). It has been reported that the PPARs agonist WY14,643 induces *G0s2* expression *in vivo* after 6 hours and *in vitro* after 5 days (Zandbergen et al., 2005).

The results for the ciprofibrate samples show that the induction of *G0s2* at 3 hours was significantly higher than the control (5.4 fold induction). This result is similar to the findings from the RNAseq analysis at 3 hours as the *G0s2* gene expression was found to be 5.8 fold above control samples (Section 3.4.6.2). The induction then subsided gradually at 5 hours, and then more so at 24 hours. This is very interesting as the timing correlates with the hepatic DNA synthesis results (Section 3.1.2.1.1) which showed the hepatic DNA synthesis peaks at 24 hours before subsequently declining.

The data are consistent with a possible relationship between *G0s2* expression and hepatic DNA synthesis, as a *G0s2* provoked G0 to G1 transition at ~ 3 hours, would be expected to give induction of DNA synthesis during the following cell cycle at ~ 20 hours later, indicating a close relation between the *G0s2* and the hepatic DNA synthesis.

The expression of *G0s2* in fisher rats treated with CPA show no significant difference from the control in all the samples treated at 1, 3, 5 and 24 hours, showing that the CPA promotes a different response from the ciprofibrate, and the *G0s2* is specific to the PPAR α .

Section 4.4 Concluding remarks

In this thesis the effects of 50 mg/kg ciprofibrate treatment for 1, 3, 5 and 24 hours on immediate-early liver gene expression in rats was studied. This was done by using cDNA microarrays, transcriptome sequencing and quantitative real-time PCR.

The hepatic DNA synthesis induced by ciprofibrate was initially characterized. This was done with BrdU immunohistochemical detection protocol, and was found to be significantly increased at 24 hours in rat liver, while it was not significantly induced until after 4 days in mice. For rats treated with ciprofibrate, the zonation of the hepatocytes found in S-phase were significantly different between centrilobular and periportal liver regions, being more dominant in the periportal region by ~20 fold. The zonal distribution of S-phase hepatocytes in mice was not different between the periportal and the perivenous zones, but had a random distribution across the liver tissue. The difference in the period and localization of the hepatic DNA synthesis between mouse and rat liver suggests there is a species difference in the kinetics between the mouse and rat's response to the ciprofibrate.

This study identifies many genes that appear to be up-regulated by ciprofibrate including previously known PPAR α agonist-responsive genes involved in processes such as PPAR signaling pathways, fatty acid metabolic pathway, cell cycle, palmitoyl-coa hydrolase activity, lipid metabolism, inflammatory responses, and stress responses. In addition, the immediate-early gene response shows novel candidate genes, consistent with a link between the genes expression at 1-3 hours and the hepatic DNA synthesis at 24 hours. The gene induction was up-regulated at 1-3 hours, then the induction came to control levels after 5 hours dosing, incorporating the hepatic DNA synthesis that was significantly high at 24 hours and then gradually decreased to control levels at 36 hours. This suggests that these genes were responsible for the hepatic DNA

synthesis that occurred after 24 hours of dosing with ciprofibrate. This is similar to previous results obtained after partial hepatectomy with the gene expression for cell cycle regulated genes, that peaks after 2-3 hours after partial hepatectomy (Taub, 2004).

Five genes were followed up in ciprofibrate and CPA treated rats with quantitative real-time PCR, these genes were *CYP4A1*, *CYP3A1*, *Ccnd1*, *Scd1* and *G0s2*. *CYP4A1* and *CYP3A1* were used as positive controls for ciprofibrate and CPA dosed animals, respectively. The *Ccnd1* and *G0s2* genes both had a significantly high induction at 3 hours with ciprofibrate treated animals, but only *Ccnd1* expression was additionally significantly high at 5 hours with the CPA dosed animals. *Scd1* expression was induced significantly with the ciprofibrate at 1 hour and not with the CPA. These results suggest that *Scd1* and *G0s2* are affected at the immediate-early hours by ciprofibrate, while they are not affected by the CPA, so they would be expected to be in a different pathway than the CPA. Also, the *Ccnd1* is a common gene that is expressed by both the ciprofibrate and the CPA at the early time points after dosing and apparently it is induced by other DNA inducers e.g. T3.

Finally, this study was able to shed some light on the mechanism of action of hepatic growth induced by ciprofibrate by studying the hepatic DNA synthesis in mouse and rat liver tissue, and by characterizing the immediate early induced genes that are provoked by peroxisome proliferators, and can cause hepatic proliferation, which might be helpful in cancer studies.

Section 4.5 Prospective work

A large number of the genes identified here could be interesting to follow up, as the microarray and the transcriptome sequencing analysis both provide a large data base for the immediate early time points and the acute response to ciprofibrate or CPA after a range of time points.

Chosen pathways or particular genes could be tested with real-time PCR at different time points, and in female rats treated with CPA as it is known that female rats give a hepatic DNA synthesis higher than the males that were used in this study, so it would be expected to have much more induction of the same genes, in particular, the DNA synthesis, the cell cycle pathway or PPAR signalling pathway genes, such as G0/G1 switch 2 (G0s2), cyclin D1 (Ccmd1), stearyl-coenzyme A desaturase 1 (Scd1), carnitine palmitoyltransferase 1a (Cpt1a), cyclin T1 (Ccnt1), Myeloblastosis oncogene-like (Mybl1), Ets variant 3+6 (Etv31 + Etv6), Kelch repeat and BTB (POZ) domain containing 11 (Kbtbd11), Myc myelocytomatosis viral oncogene homolog 1 (Myc11 v), Protein phosphatase 1K (Ppm1k) or T-box 3 (Tbx3).

Additionally, the proteins induced by the PPAR could be studied by western blotting. These genes could be tested in other species in particular human tissues for comparison and to predict the level of danger, if any, from the peroxisome proliferators in concern with the hepatocarcinogenesis.

Also, immunohistochemical protocols could be used against specific antigens to indicate, where in the liver zones do particular genes begin their induction (by following the pattern at the different time points) if periportal or perivenous, and this could help to tie groups of genes together and introduce a small view of their mechanism of action in response to both the ciprofibrate and the CPA.

Furthermore, to prove that the genes identified in this study are actually involved in or required for the hepatic DNA synthesis induction, it would be exciting to follow up with knock-out mice or rats of these genes and test if the induction is affected. It might also be worth checking if the rodents that do not react to peroxisome proliferators have these genes, and to examine if these genes are induced after dosing.

Appendix I of RNA, cDNA and dye yields of the microarray samples

Sample ID	RNA yield $\mu\text{g}/\mu\text{l}$	cDNA yield $\text{ng}/\mu\text{l}$ for microarray	Alexa flour 555 dye attachment to cDNA $\text{pmol}/\mu\text{l}$
CIPROFIBRATE			
Treated cipro 1h-1	10.08	79.6	4.7
Treated cipro 1h-2	15.66	92.7	2.5
Treated cipro 1h-3	9.4	112.9	10.5
Treated cipro 1h-4	8.38	159.4	12
Treated cipro 3h-1	13.6	350.6	13.6
Treated cipro 3h-2	11.7	144	11.7
Treated cipro 3h-3	13.36	43.97	6.3
Treated cipro 3h-4	11.06	35.2	5.2
Treated cipro 5h-1	9.14	218.1	50.1
Treated cipro 5h-2	10.26	152.2	33.5
Treated cipro 5h-3	12.82	71.32	19.7
Treated cipro 5h-4	13.8	129.6	15.9
Treated cipro 24h-2	4.88	19.8	3.6
Treated cipro 24h-3	3.59	13	1.3
Treated cipro 24h-5	3.06	17.3	1.7
Treated cipro 24h-6	5.33	51	8.8

CYROTERONE ACETATE			
Treated CPA 1h-1	3.95	99.6	11.3
Treated CPA 1h-2	5.2	60.3	5.7
Treated CPA 1h-3	7.8	77.1	16.8
Treated CPA 1h-4	9.75	69.1	7.5
Treated CPA 3h-1	8.43	12	4.9
Treated CPA 3h-2	2.93	7.9	2.3
Treated CPA 3h-3	4.67	39.9	3
Treated CPA 3h-4	7.32	19.3	6.7
Treated CPA 5h-1	7.01	19.9	10.8
Treated CPA 5h-2	7.3	57.5	13
Treated CPA 5h-3	4.75	34.1	11.5
Treated CPA 5h-4	3.69	5.9	4.4
Treated CPA 24h-1	8.34	84.4	11.4
Treated CPA 24h-2	8.24	140.5	10.8
Treated CPA 24h-3	12.34	62.8	9.6
Treated CPA 24h-4	4.02	140.6	44.9
CORN OIL (CONTROL)			
control 1h-1	15.54	153.7	14.6
control 1h-2	8.08	114.4	7.6
control 1h-3	14.04	127.8	13
control 1h-4	7.46	136.5	16.7

control 3h-1	8.14	36	4.4
control 3h-2	7.56	10.7	1.5
control 3h-3	5.31	16.8	3.8
control 3h-4	4.88	12.7	3.2
control 5h-1	11.2	9.8	2.3
control 5h-2	6.72	24	4.6
control 5h-3	8.14	8	2.4
control 5h-4	7.33	18	6.5
control 24h-1	3.65	142.5	26.9
control 24h-2	6.24	373	46.2
control 24h-3	3.86	45.4	6.7
control 24h-6	5.32	71.4	8.4

Record of Training Courses

Mrs. Abeer H. A. Amer

student ID number 4047465

Record of Training courses	credits
Institute of genetics courses	
Radiation training course	
Safety training for postgraduate research student	
Biological safety training	
Training of Personnel under the Animals (Scientific Procedures) Act 1986 Home office license course (model 1)	2
Home office license course (model 2)	2
Home office license course (model 3)	2
RT-PCR course	2
Seminars	
Graduate School courses	
Library induction and use of software	1
Exploiting the power of MS word a: for individual chapters and academic papers	1
Exploiting the power of MS word b: combining chapters into the thesis	1
MS Excel functionality a: entering, formatting and storing data	1
MS Power Point: developing a professional presentation	1
Computer and molecular biology	4
Further presentation skills	1
Getting started with research design and statistics	3
Planning research and time management	1

Introduction to image and photo editing	2
Nature of the PhD and the supervision process	1
Guidelines for planning and formatting theses	1
Getting going on your thesis	2
Intensive learning and teaching program	8
Beginning writing	1
Academic writing/ synthesis sources	
Introductory events to teaching for PG research students and researchers	
Training for reading to tape for disabled students	
Training for note-taking	
Training for invigilation	
Training for information technology (IT) support	
Presentations at the School Post-Graduate Symposia	2
First platform presentation	
Second year poster presentation	
Third year platform presentation	
Demonstrating	2
Molecular and development neuroscience (ligand binding practical class)	
The manipulating genes and genomes	
The genes and cellular control	
The C. Elegans practical	
TOTAL CREDITS	41

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