



Published in final edited form as:

J Clin Gastroenterol. 2016 January ; 50(1): 80–84. doi:10.1097/MCG.0000000000000400.

Insulin resistance is associated with significant liver fibrosis in chronic hepatitis C patients: A systemic review and meta-analysis

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Abstract

Background—The role of insulin resistance (IR) on fibrosis progression in HCV patients has not been systematically evaluated. Therefore, this systemic review aimed to summarize the available epidemiologic evidence to evaluate the strength of association between IR and advanced liver fibrosis in these patients.

Methods—We performed a systemic literature search in PubMed, OvidSP and MEDLINE from January 1990 to April 2015 without language restriction using the following search terms: insulin resistance, liver fibrosis, cirrhosis, diabetes mellitus and chronic hepatitis C. Publication bias was assessed using the Begg and Egger's tests and with a visual inspection of funnel plot. All analyses were performed using Comprehensive Meta-Analysis version 2 software.

Results—A total of 3,659 participants with HCV infection from 14 studies were included in the analysis. After adjusting for publication bias, the RR for significant hepatic fibrosis among HCV subjects with IR was 1.63 (95% CI 1.34-2.01). Subgroup analysis by genotypes showed RR of 2.16 (95% CI 1.52-3.06) for genotype 1; however, the association was no longer significant when we analyzed the data for HCV genotype 3; RR 1.40 (95% CI 0.8-2.45).

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Conflict of interest: All authors reported no conflicts of interest to the content of this manuscript

Authors contribution:

SP and SL – study design

SP and RP – literature search, review and collection of data

SP and RJ – analysis and interpretation of data

SP, RJ and RP – drafted manuscript

SL – made critical revisions to manuscript

Conclusion—Our study showed significant association between IR and significant hepatic fibrosis in patients with HCV genotype 1 infection.

Keywords

insulin resistance; liver fibrosis; chronic hepatitis C

INTRODUCTION

Hepatitis C virus (HCV) infection is a major global public health problem and it is estimated that ~2%-3% of the world population are infected with the virus¹. It can now be successfully eradicated with the new therapeutic regimens; which are safe and highly efficacious². However, the cost of these new drugs prohibits their use in many countries with limited resources around the world¹. Undoubtedly, the morbidity and mortality from HCV infection continue to increase¹.

Most patients who acquire HCV develop chronic HCV infection. However, the rate of disease progression from the time of infection till the development of advanced fibrosis or cirrhosis is variable³. In fact, the natural history of chronic HCV infection is difficult to determine because of the difficulty in estimating the time and duration of infection and other factors that can affect disease course. Though the mechanism is poorly understood, it is likely that both host and viral factors play an important role in the disease progression. Several factors have been reported to influence the fibrosis progression, including age⁴, ethnic background⁵, gender⁶, and alcohol use⁷.

Recent epidemiological studies suggested that HCV infection is an independent predictor for the development of type 2 diabetes mellitus (DM), and that type 2 DM is more prevalent among patients with chronic HCV infection than in those with other causes of liver diseases⁸⁻¹⁰. It is likely that the HCV itself or the inflammatory response to HCV infection contributes to the development of insulin resistance (IR) and thus increasing the risk for type 2 DM¹¹. The presence of IR and type 2 DM are independent predictors of severe fibrosis in patients with non-alcoholic fatty liver disease^{12, 13}. The role of IR and advanced fibrosis in HCV patients has not been systematically evaluated. Therefore, this systemic review aimed to summarize the available epidemiologic evidence to evaluate the strength of association between IR and liver fibrosis in these patients.

METHODS

Study selection/search Strategy

We performed a systemic literature search in PubMed, OvidSP and MEDLINE from January 1990 to April 2015 without language restriction using the following search terms: insulin resistance, liver fibrosis, cirrhosis, diabetes mellitus and chronic hepatitis C. The reference list of each included study was comprehensively searched to further identify relevant studies. The process of systematic review was conducted in adherence to standards of quality for reporting meta-analyses¹⁴.

Inclusion and exclusion criteria

Relevant studies were included if they met the following criteria: (1) the studies were either case-control or cohort study designs; (2) the study participants were ≥ 18 years old; and (3) the relative risk (RR) estimate or odds ratio was reported for significant hepatic fibrosis in those with HCV infection.

Definition of hepatic fibrosis/cirrhosis

The following pathological classification for hepatic fibrosis was used; METAVIR, Scheuer, Ishak and histological activity index (HAI)^{4, 15-17}. METAVIR scoring system was specifically designed for patients with HCV infection⁴. The fibrosis score is assigned a number from F0-F4 where F3-F4 representing advanced fibrosis. Scheuer classification defines stages of fibrosis from 0 to 4 where stages 2-4 indicate significant¹⁷. Ishak scale assesses liver fibrosis in 7 categories, ranging from normal (stage 0) to cirrhosis (stage 6)¹⁵. For this scoring system, stages 4-6 indicate significant fibrosis. Lastly, HAI classifies fibrosis from stage A to D and gives a score to each stage where score 3 or more indicates advanced fibrosis¹⁶.

Definition of insulin resistance

Insulin resistance was measured using the HOMA (Homeostasis Model Assessment). The cut-off for the HOMA to define the presence of IR in each study is shown in Table 1.

Data extraction

The following information were extracted from each study: publication data (such as first author's last name and first name initials, year of publication and country of origin), sample size, participants' demographic data, types of study design (case-control/cohort), number of cases and controls (for case-control studies), number of exposed and unexposed (for cohort studies), criteria used to define significant hepatic fibrosis stratified by different pathological classification, the levels of HOMA scores to define IR, risk estimates with their corresponding confidence intervals (CIs), and the covariates (if any) which were used in the multivariate modeling. We carefully reviewed the potential confounders; that might be associated with the risk of liver fibrosis in the studied population. In this study, odds ratios (ORs) from case-control studies were considered as estimate of relative risk. This is based on the assumption that the prevalence of HCV infection is $<10\%$, and in this case the odds ratio and relative risk will be approximately the same. Two independent reviewers (SP and RJ) reviewed the studies and any discrepancies regarding inclusion/exclusion or risk estimates were resolved through the discussion by authors. We used Cohen's kappa coefficient to assess the agreement among reviewers for inclusion/exclusion of specific studies¹⁸.

Assessment of methodological quality

To assess methodological quality of all the publications that were included in the final analysis, the Newcastle-Ottawa Scale was used¹⁹. The scale allocates stars, maximum of nine, for the following categories: quality of selection, comparability, exposure and outcome of study participants. Any studies with the scale < 5 were excluded.

Statistical analysis

Summaries of relative risk (RR) estimates were evaluated using both fixed- and random-effects methods. Initial analysis was performed to look for association between IR and significant liver fibrosis. We used Cochran's Q -test and I^2 -statistic to determine the heterogeneity of the publications. Publication bias was assessed by (i) construction and visual inspection of funnel plot and (ii) employing the Egger's and Begg and Mazumdar tests. Duval and Tweedie's trim and fill method was utilized to obtain RR after adjustments of the publication bias. The p value of < 0.05 indicated statistical significance. All analyses were performed using Comprehensive Meta-analysis Version 2 (Biostat, Englewood, New Jersey)²⁰.

RESULTS

Literature Search and Study Characteristics

The schematic diagram of the detailed literature selection is shown in Figure 1. We identified 380 studies from different databases, either in full publications or in abstract forms. After title appraisal and extensive review, 56 publications were considered to be potentially relevant. Of these, we excluded 8 review articles, 5 animal studies, 2 letters to the editors, 8 studies which described HIV/HCV co-infections, 2 studies with post liver transplant HCV patients, 1 clinical trial, and 12 studies which did not provide RR estimate. Four additional studies were excluded as they were cross sectional studies. Fourteen studies (12 cohort and 2 case-control studies) were considered for full article assessment and included in the final analysis.

Publication quality and bias

The Newcastle-Ottawa Scale (range, 1-9 stars) to assess the publication quality revealed average 5.6 stars for the twelve cohort studies and 6.5 for the two case-control studies. The detail for the scale for each study is shown in Table 1. Due to the presence of publication bias (funnel plot, Supplemental Fig 1, Begg and Mazumdar test: $p = 0.04$; Egger's test: $p = 0.006$), adjusted RR was used to report the results (Figure 2).

Association between insulin resistance and liver fibrosis

A total of 3,659 participants with HCV infection were included in the analysis. Of these, 12 were cohort studies consisting of 3,259 subjects. Due to evidence of heterogeneity ($Q=29.83$, p value for heterogeneity 0.005, $I^2= 56.42\%$), we used random-effect model to report the pooled RR. The pooled RR for significant hepatic fibrosis among HCV subjects with IR was 1.89 (95% CI 1.54 -2.33) (Figure 2 and Fig S1). After adjusting for publication bias, the association remained significant with the adjusted RR 1.63 (95% CI 1.34-2.01).

We also performed subgroup analysis. Analysis of the 12 cohort studies showed RR of 2.02 (95% CI 1.6-2.55, $p<0.001$). Subgroup analysis by genotypes showed RR of 2.16 (95% CI 1.52-3.06) for Genotype 1 (Figures 3, 3 studies²¹⁻²³); however, the association was no longer significant when we analyzed the data for HCV genotype 3 (3 studies^{21, 24, 25}); RR 1.40 (95% CI 0.8-2.45)(Figure 3 and Fig S2). We also analyzed the strength of association between IR and significant fibrosis, stratified by the different pathological classification. We

found that the RR was 1.80 (95%CI 1.28-2.52), when considered the studies using METAVIR method (Figure 4 and Fig S3, 6 studies²⁶⁻³¹). The association was still significant when other classifications (Figure 5, 8 studies^{21-25, 30, 32, 33}) were used, RR 1.73 (95% CI 1.34-2.23).

Subgroup analysis was also performed based on the geographical location. There were no geographical differences in the association between IR and hepatic fibrosis; RR 1.69 (95% CI 1.32-2.17) for the studies from Europe and Australia (Figure S4, 10 studies^{21-25, 27-29, 33, 34}) and RR 1.90 (95% CI 1.27-2.83) for those from Asia (Figure S5, 4 studies^{26, 30, 32}).

DISCUSSION

The major findings of our study are the followings: 1) the presence of IR is a significant risk factor for advanced hepatic fibrosis in HCV patients and 2) whereas no association is observed for those infected with HCV genotype 3, the risk for significant fibrosis is increased for those infected with HCV genotype 1.

Association between hepatitis C infection and IR

The causal relationship of HCV infection and IR development has been demonstrated by the increased prevalence of IR in chronic HCV infection. The prevalence of IR in those infected with HCV is significantly higher than that in the general population^{35, 36}. The mechanism of HCV-induced IR is complex. Following inflammatory response in the liver to HCV infection, a profound impairment of insulin signaling occurs at the level of insulin receptor substrate (IRS) tyrosine phosphorylation and phosphoinositide 3-kinase activation³⁷. The increase in the levels of tumor necrotic factor- α by HCV core protein may also lead to proteasomal degradation of IRS1 and IRS2, resulting in the alteration of insulin function and the development of IR³⁷.

In addition to the direct effect of HCV on insulin signaling, the development of IR can also be mediated through hepatic steatosis. This can coexist with HCV, regardless of genotype, in patients with risk factors such as obesity and hyperlipidemia. Hepatic steatosis can also be related to the direct hepatopathic effect of genotype 3 viral infection³⁸. In this scenario, the relationship between IR and HCV infection is bidirectional³⁷; HCV induces steatosis and the latter could also cause IR³⁹. HCV-associated hepatic steatosis is mainly virus-induced in genotype 3 infected patients due to the impairment in very low-density lipoprotein (VLDL) secretion³⁸. However, in non-genotype 3, the development of IR is likely to play a major role in steatosis^{37, 38}.

Insulin resistance and hepatic fibrosis

Once developed, IR plays an important role in promoting hepatic fibrosis. Hyperinsulinemic state associated with IR directly activates stellate cells⁴⁷⁻⁵⁰. Furthermore, IR-induced hepatic lipid accumulation and generation of ROS can also indirectly activate stellate cells and initiate the cellular signaling cascades triggering hepatic fibrosis. Our findings that the progression of fibrosis in patients with IR is genotype-specific deserve further comments. Although the interference with the insulin sensitivity shows some HCV genotype-

specificity, IR has been reported to occur in all HCV genotypes, but to a different extent^{37, 40}. Patients infected with the genotype 3 virus have a lower prevalence of IR when compared with those infected with the other viral genotypes, even after adjustment for the effects of body mass index (BMI) and other confounders^{41, 42}. However, genotype 1 infection was found to be a significant determinant of severe IR, even in patients without underlying diabetes mellitus³⁷. The effect of different genotypes of HCV on the severity of IR is likely explained our findings.

Limitation

Our systemic review has some limitations. First, there are factors which were not taken into consideration for the adjusted RR analysis such as age, gender, body mass index, duration of HCV infection and family history, primarily due to unavailability of these data in the original studies. Second, information on the history of alcohol intake was not uniformly provided. In the studies that mentioned alcohol intake, there were discrepancies in the amount as well as the cut-off levels for hazardous alcohol use. One study excluded any amount of alcohol users²³ while others^{26, 28, 33, 34} used different levels of alcohol intake as the cut-off and one study³¹ did not consider alcohol consumption in exclusion criteria.

Summary and Clinical implications

This study elucidates the important relation for a genotype-specific association between IR and significant fibrosis in patients with HCV infection. Improvement in IR either by weight loss, life style change or insulin sensitizer can significantly improve SVR rates and treatment outcomes⁴³. At present, it is unclear how insulin resistance will impact the response to treatment with the newly effective anti-viral agents for HCV. However, given the limited access to these new medications in other parts of the world, strategies to improve insulin sensitivity should be explored as they might mitigate against the progression of fibrosis.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgments

Financial support/Sources of funding: This study is supported by K08 AA016570 from the NIH/NIAAA, 1I01CX000361-01 from the Veterans Affairs Research and Administration, and W81XWH-12-1-0497 from United States Department of Defense (All to S.L)

Abbreviations

HCV	Hepatitis C virus
HOMA	Homeostasis model assessment
IR	Insulin Resistance
OR	Odds Ratio
RR	Relative Risk

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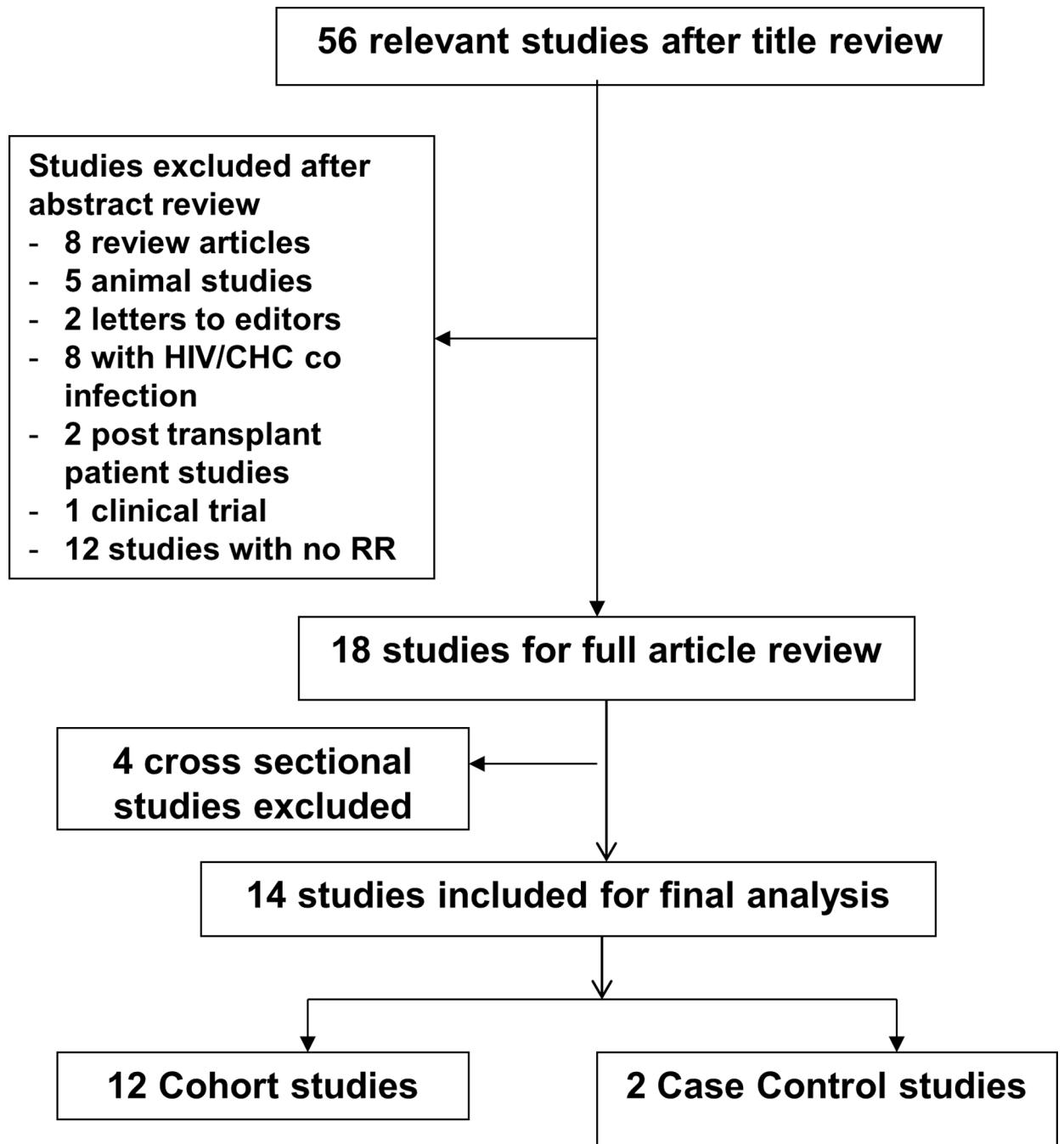


Figure 1.
Schematic diagram for study selection of the relevant articles

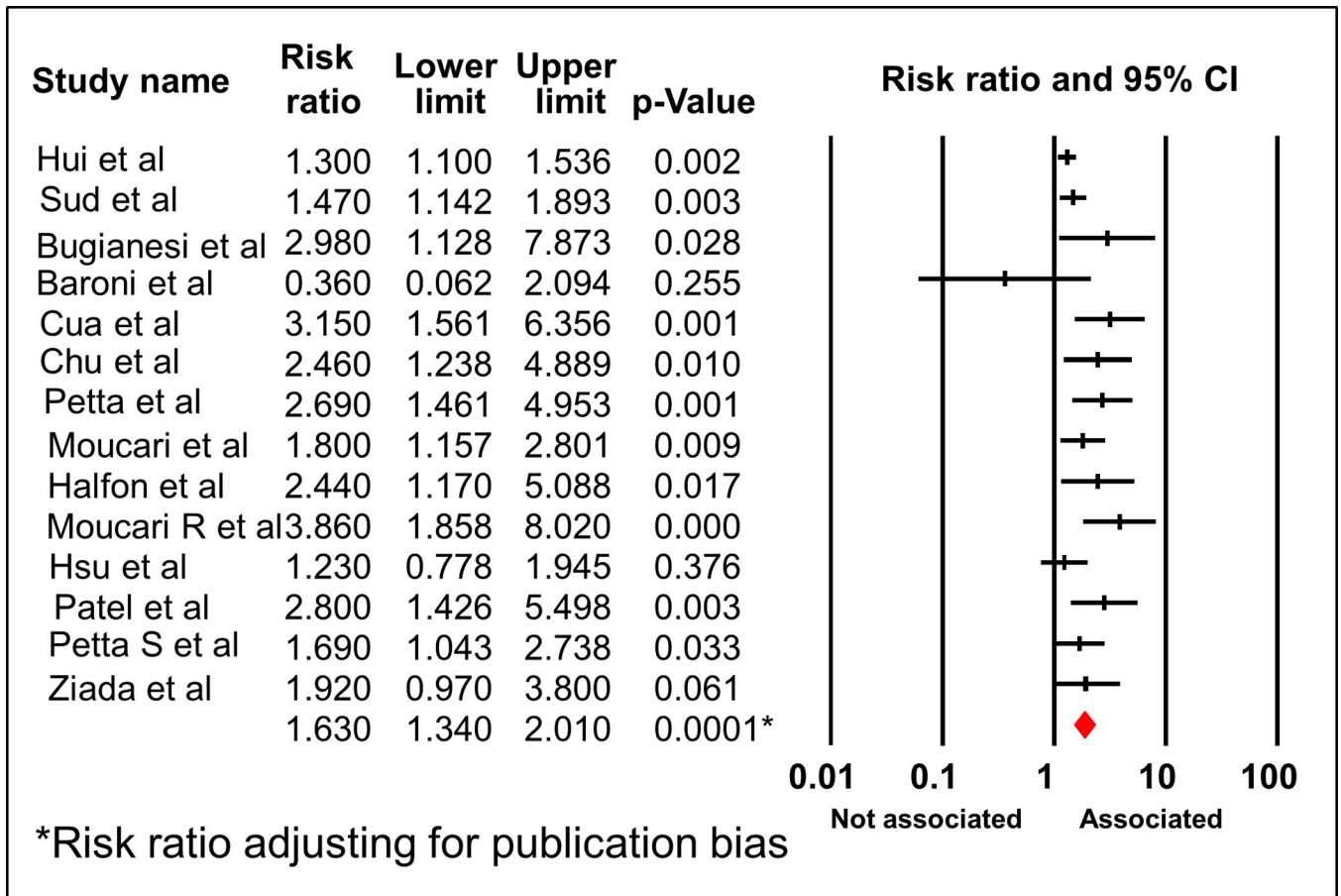
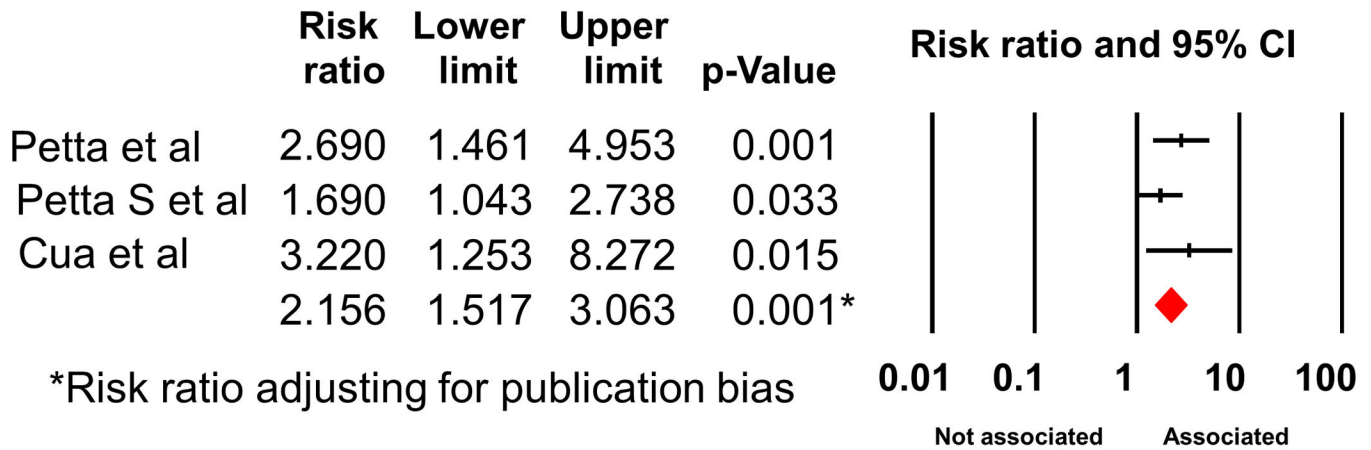


Figure 2. Forest plot of meta-analyses demonstrating the association between IR and significant fibrosis of all 14 studies

Subgroup analysis for Genotype 1



Subgroup analysis for Genotype 3

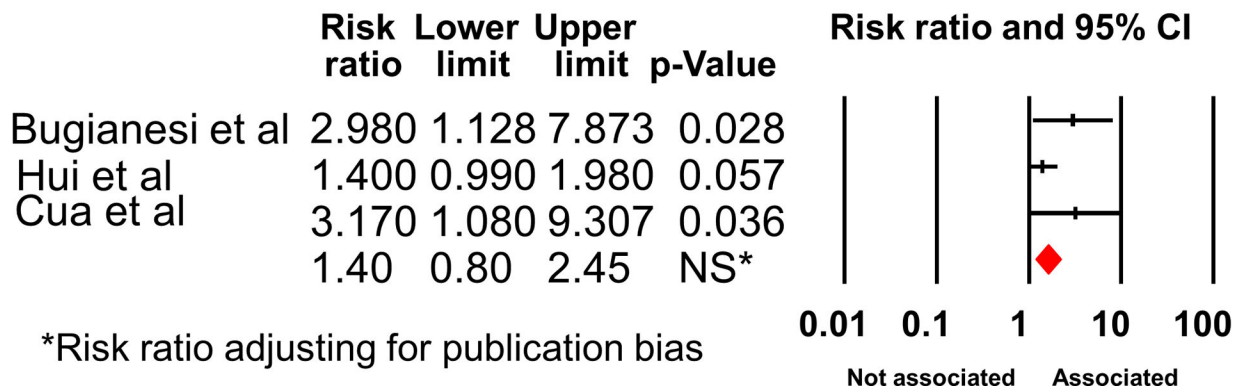


Figure 3. Forest plot demonstrating association between IR and significant fibrosis, subgroup analysis stratified by genotypes

Subgroup analysis - Liver Fibrosis Staging using METAVIR

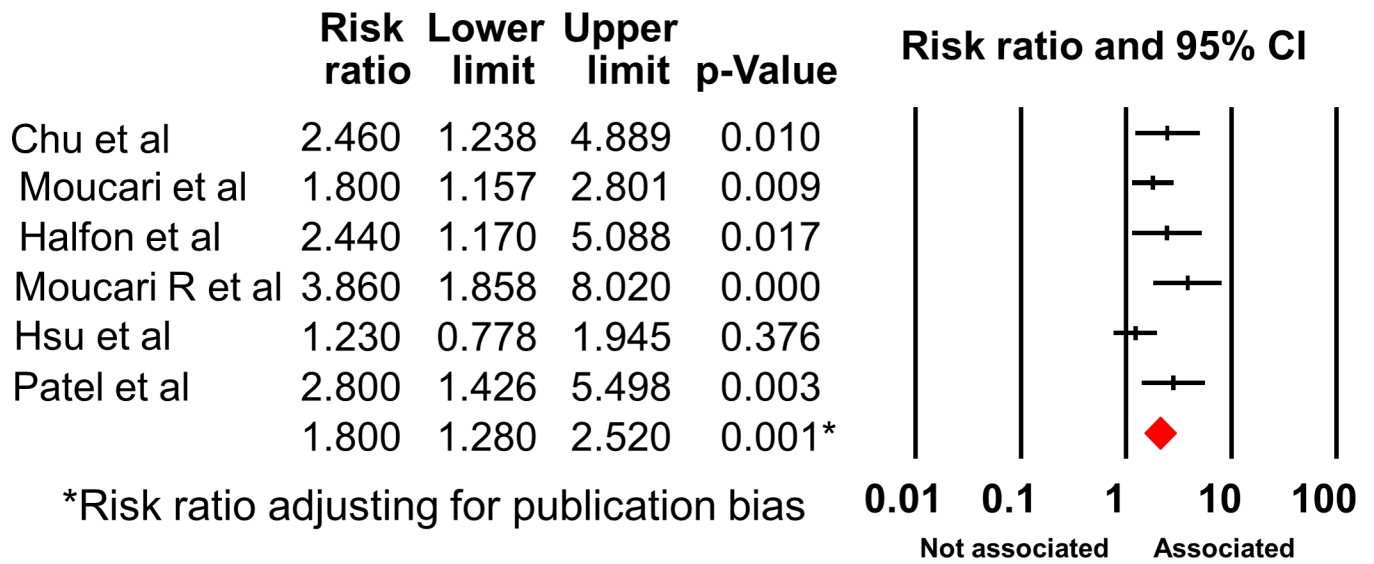
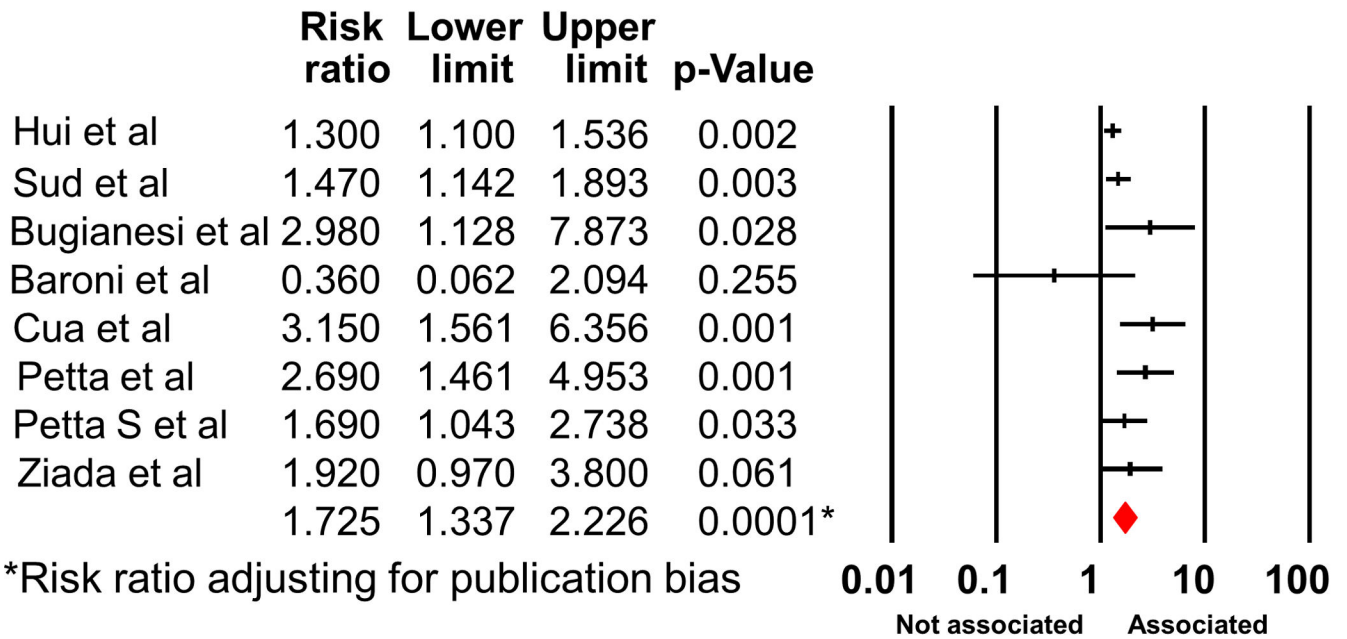


Figure 4. Forest plot demonstrating association between IR and significant fibrosis, subgroup analysis for the studies using METAVIR as the pathological classification for fibrosis

Subgroup analysis - Liver Fibrosis Staging by other methods[§]



[§] Fibrosis classification by Scheuer, Ishak and histological activity index (HAI)

Figure 5.

Forest plot demonstrating association between IR and significant fibrosis, subgroup analysis for the studies using non-METAVIR methods as the pathological classification for fibrosis

Table 1

Study characteristics and RR with CI for individual studies

No	Study Name	Year of publication	Country	Type	Male: Female	Single center Vs Multi Center	N-O scale*	Cases of CHC	Genotype	Fibrosis definition	IR definition	Total RR (95% CI)
1	Hui et al ²⁴	2003	AUS	Case-control	175:85	Single	7	260	3 and Non 3	Scheuer	HOMA	1.3 (1.1– 1.4)
2	Sud et al ³⁴	2004	AUS	Cohort	115:61	Multi	6	176	**	Scheuer	HOMA	1.47 (1.14– 1.89)
3	Bugi anesi et al ²⁵	2006	AUS	Cohort	101:31	Single	7	132	3	Scheuer	HOMA	2.98 (1.13– 7.89)
4	Baroni et al ³³	2007	Italy	Cohort	62:28	Multi	7	90	3 and Non 3	Ishak	HOMA >2.7	0.36 (0.06– 2.05)
5	Cua et al ²¹	2008	AUS	Cohort	235:111	Single	5	346	1 and 3	Scheuer	HOMA>2	3.15 (1.56– 6.35)
6	Chu et al ²⁶	2008	China	Cohort	**	Single	5	192	**	Metavir	HOMA >2.5	2.46 (1.24– 4.9)
7	Petta et al ²²	2008	Italy	Cohort	105:96	Single	6	201	1	Scheuer	HOMA >2.7	2.69 (1.46–4.95)
8	Moucari et al ²⁷	2008	France	Cohort	254:200	Single	6	454	1 and 4	Metavir	HOMA >3	1.8 (1.16– 2.82)
9	Halfon et al ²⁹	2009	France	Cohort	121:49	Single	5	170	**	Metavir	HOMA >2	2.44 (1.15– 5)
10	Moucari et al ²⁸	2009	France	Cohort	168:58	Single	6	226	4	Metavir	HOMA>3	3.86 (1.86– 8.03)
11	Hsu et al ³⁰	2010	Taiwan	Cohort	288:246	Single	5	528	1 and 2	Metavir	HOMA >2.4	1.23 (0.78– 1.95)
12	Patel et al ³¹	2011	Asia	Cohort	141:122	Multi	5	263	2 and 3	Metavir	HOMA>2	2.80 (1.4– 5.4)
13	Petta et al ²³	2012	Italy	Cohort	239:242	Single	5	481	1	Scheuer	HOMA>3	1.69 (1.04– 2.73)
14	Ziada et al ³²	2013	Egypt	Case-control	75:65	Single	6	140	**	HAI	HOMA 2	1.92 (0.97– 3.4)

* Newcastle-Ottawa Scale