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**Identification of SNPs in genes potentially affected by  
domestication to efficiently detect hybridization  
between wild and domestic cats**

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## Abstract

The European wildcat (*F. s. silvestris*) is a medium-sized carnivorous mammal that has suffered significant population decline and range contraction and fragmentation during the last century. The species is also currently threatened by human-mediated hybridization with domestic cats. Their hybrid offspring is fertile and can backcross with pure wildcats, leading to the introgression of domestic cat genes into the wildcat gene pool. This poses a danger to the genetic integrity of the European wildcat, and may contribute to its extinction. Hence, methods are urgently needed that are able to distinguish between pure wildcats and hybrids. However, the accurate detection of hybridization based on available morphological characters is particularly difficult, and even the panels of molecular markers that have been reported either are not highly reliable in detecting hybrids or are too cumbersome to be of practical value for routine use.

Here, I aimed to identify single nucleotide polymorphisms (SNPs) located in genes that may have been affected during the domestication process of the domestic cat (*F. s. catus*). A total of 51 candidate genes were selected for sequencing of exonic and flanking intronic regions in samples of wildcats and domestic cats. Analysis of the resulting sequences revealed 13 SNPs with elevated genetic differentiation ( $F_{st} > 0.8$ ) between the two species. Subsequently, genotypes of first-generation hybrids (F1), second-generation hybrids (F2), and backcrosses with wildcats (B x FSI) and with domestic cats (B x FCA), were simulated in order to evaluate the diagnostic power of the 13 SNP set. Admixture analysis of the simulated genotypes showed that the marker panel could identify and assign each genotype to the correct category, with an accuracy of about 90%.

Given the current need for efficient and reliable tools to detect and discriminate samples from wildcats, domestic cats, and their hybrids, this small panel of 13 SNPs with apparently high diagnostic power offers the promise of a convenient and effective assay for surveying and monitoring the distribution and hybridization status of European wildcats.

**Keywords:** *Felis silvestris*, hybridization, SNP, candidate genes, diagnostic markers

## Resumo

A crescente desflorestação e expansão das áreas afectadas pela presença e actividade humana no planeta têm tido inúmeras consequências directas e indirectas na abundância e sobrevivência de espécies sensíveis aos impactos antropogénicos. Entre estas espécies em Portugal destaca-se o gato-bravo europeu (*Felis silvestris silvestris*). A fragmentação e a degradação do habitat e a perseguição e a caça pelo Homem obrigaram este carnívoro de médio porte a refugiar-se em áreas geográficas restritas, isoladas e dispersas da Europa.

Uma ameaça adicional, críptica e ainda pouco estudada, e potencialmente uma grande ameaça à conservação do gato-bravo europeu, é o seu contacto e interacção com o gato doméstico (*Felis catus* ou *Felis silvestris catus*). O gato doméstico é um animal de estimação popular, com uma população mundial estimada em 600 milhões de indivíduos. Estudos anteriores concluíram que o gato doméstico é o resultado de gerações de seleção artificial iniciada em gatos-bravos na pré-história humana, existindo evidências arqueológicas de relações entre gatos e homens desde há 9500 anos. Análises genéticas sugeriram que a área de origem do gato doméstico terá sido o Cresceste Fértil no Médio Oriente, nos primórdios da agricultura e da pecuária, onde a função dos gatos seria principalmente de controlo de roedores. Desde essa altura até tempos recentes a seleção artificial no gato doméstico terá tido uma intensidade relativamente moderada, mas a partir do século XIX esta selecção intensificou-se e, por razões de preferências estéticas, gerou uma abundância de raças com diferentes cores e padrões de pelagem, ainda que mantendo em geral a morfologia anatómica típica da espécie ancestral.

Além de outros potenciais impactos, incluindo ecológicos, comportamentais, parasitológicos e epidemiológicos, um maior contacto entre gatos domésticos e gatos-bravos, tendo em conta a raridade da segunda espécie, aumenta o risco de hibridação entre as duas espécies. Observações de campo e capturas de indivíduos têm demonstrado a existência de híbridos. Sendo estes férteis e capazes de hibridar com indivíduos parentais, a hibridação entre as duas espécies, especialmente se for comum ou frequente, pode assim constituir uma grave ameaça para o gato-bravo, uma vez que os retrocruzamentos entre híbridos e gatos-bravos puros podem levar à introgressão de genes de gato doméstico no genoma do gato-bravo. Este processo pode conduzir à quebra da integridade genética do gato-bravo e, em última análise, à extinção da espécie. Como o gato-bravo é um predador de topo em muitos ecossistemas, a sua extinção pode ter consequências graves nas cadeias tróficas e redes ecológicas. Por conseguinte, é importante implementar medidas de conservação para o gato-bravo. A fim de monitorizar, controlar e reduzir a hibridação, são necessários métodos que sejam capazes de distinguir gatos-bravos puros de híbridos. Os primeiros métodos propostos com este objectivo eram baseados em caracteres anatómicos ou morfológicos, especialmente medidas cranianas e padrões da pelagem. Contudo, vários estudos demonstraram a dificuldade em utilizar estes métodos, bem como a sua ineficiência em certos casos, devido à grande semelhança morfológica frequentemente observada entre gatos-bravos puros e híbridos. Consequentemente, tem crescido o interesse no desenvolvimento e aplicação de abordagens baseadas em marcadores moleculares.

Um dos marcadores genéticos com resultados promissores neste contexto são os microssatélites, que são unidades de pequenas repetições da mesma sequência de nucleótidos (por exemplo, AGAGAGAG) e que têm elevadas taxas de mutação em comparação com outras regiões do genoma. Através do uso de microssatélites, foi possível avaliar o grau de hibridação em várias regiões europeias, desde países como a Itália e Alemanha em que as populações de gatos-bravos e gatos domésticos ainda se encontram bastante diferenciadas, até aos casos da Escócia e Hungria, onde os níveis de introgressão são elevados. No entanto, devido à sua elevada taxa mutacional,

os microssatélites tendem a sofrer de homoplasia, o que pode levar à falsa detecção de híbridos e a dificuldades em identificar e distinguir híbridos de gerações diferentes e retrocruzamentos. Assim, as atenções têm-se voltado para outro tipo de marcador molecular potencialmente útil em estudos de hibridação e que, devido à sua baixa taxa de mutação têm menos problemas de homoplasia do que os microssatélites: os “single nucleotide polymorphisms” (SNPs).

Os SNPs consistem em variações na sequência de ADN num só nucleótido, o que faz com que sejam os polimorfismos mais abundantes no genoma. Tendo os SNPs frequentemente apenas duas variações alélicas, se estas tiverem frequências muito diferentes entre duas espécies ou populações, e considerando a geralmente limitada homoplasia nos SNPs, nesses casos os SNPs podem revelar-se altamente diagnósticos. Estudos anteriores têm apresentado painéis de SNPs com capacidade de identificar e distinguir gatos-bravos puros, gatos domésticos puros, e híbridos até à segunda geração. Porém, esses painéis contêm um elevado número de marcadores, não sendo portanto testes práticos e eficientes. O facto de o poder estatístico desses painéis depender aparentemente do uso de um número elevado de SNPs é possivelmente consequência da maioria deles não estar fixada para alelos diferentes nas duas espécies. É expectável que esses SNPs fixados e altamente diagnósticos estejam em regiões do genoma onde exista elevada diferenciação entre gatos-bravos e gatos domésticos. Recentemente, um estudo procurou identificar as regiões que terão sido mais afectadas pela selecção e influentes na domesticação do gato doméstico. Esse estudo baseou-se na sequenciação do genoma de seis gatos-bravos e 22 gatos domésticos pertencentes a raças com origens geográficas distantes. Regiões do genoma exibindo elevados níveis de diferenciação genética entre gatos-bravos e gatos domésticos, identificados através de valores elevados do índice de fixação ( $F_{st}$ ), e simultaneamente baixos níveis de variação intraespecífica, indicados por valores reduzidos de heterozigosidade ( $H_p$ ), foram consideradas como potencialmente impactadas por selecção positiva. Assim, identificaram-se cinco regiões possivelmente influenciadas por selecção positiva. Estas regiões contêm vários genes envolvidos em diversos processos neurais, sendo muitos desses genes em particular determinantes do comportamento. Notavelmente, vários desses genes codificam proteínas que controlam a sobrevivência e migração das células da crista neural. Esta observação está de acordo e dá suporte à hipótese da síndrome da domesticação, uma teoria que postula que os caracteres morfológicos e comportamentais modificados em mamíferos domesticados têm origem em alterações na migração das células da crista neural durante o desenvolvimento embrionário. Com base nas descobertas desse estudo e de outros semelhantes em outros mamíferos domésticos, é possível focar em genes candidatos a pesquisa de marcadores potencialmente discriminantes entre mamíferos domésticos e os seus ancestrais selvagens.

O presente estudo teve como objetivo descobrir, através da análise de genes candidatos, e testar um painel de SNPs com elevado poder diagnóstico para distinguir com fiabilidade diferentes classes de gatos puros e híbridos. Os genes candidatos estudados pertencem a três categorias funcionais diferentes: i) desenvolvimento neuronal, ii) cor e estrutura da pelagem, e iii) sistema sensorial. Para cada gene selecionado, um ou dois pares de “primers” foram desenhados para amplificar fragmentos de 700-900 pares de bases em amostras de gatos-bravos de Portugal, França e Roménia, e gatos domésticos portugueses. Um conjunto de 13 genes revelou SNPs com valores elevados de diferenciação ( $F_{st} > 0.8$ ) entre gatos-bravos e gatos domésticos. Posteriormente, genótipos de híbridos de primeira geração (F1), segunda geração (F2), e retrocruzamentos com gatos-bravos (B x FSI) e com gatos domésticos (B x FCA) foram computacionalmente simulados com base nos genótipos das classes parentais amostrados. Usando os 13 SNPs, cerca de 90% dos indivíduos analisados foram corretamente identificados e atribuídos à sua categoria. Com base nestes resultados, constata-se preliminarmente que os SNPs identificados neste estudo possuem um valor diagnóstico potencialmente comparável ao de painéis com mais SNPs reportados em estudos anteriores. O painel aqui desenvolvido, carecendo obviamente de análises adicionais que confirmem os resultados obtidos, pode futuramente

constituir um método eficaz e fiável para caracterizar padrões de hibridação e introgressão em populações de gato-bravo, facilitando o trabalho de indivíduos e instituições envolvidas na conservação da espécie na Europa.

Palavras-Chave: Hibridação, gato doméstico, gato-bravo, marcadores diagnósticos, SNP.

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## 1. Introduction

### 1.1 Hybridization and the species concept

The process of hybridization is often defined as the reproduction between individuals belonging to genetically distinguishable populations (Barton and Hewitt 1985). In most cases involving different species, these matings result in sterile offspring (Todesco et al. 2016). However, in other instances, such crosses can produce fertile hybrids that are capable of reproducing with individuals of both parental species. This backcrossing can induce the movement of genes from one of the parental species to the other, which is called introgressive hybridization (Randi et al. 2008).

The phenomenon of hybridisation is closely tied to the discussion around the definition of “Species” as a biological unit. One of the most relevant species concepts brought forth was the one by Ernst Mayr in 1942. In his book, *Systematics and the Origins of Species*, Mayr defined his “Biological Species Concept” (BSC) in which species are groups of interbreeding populations that are unable to exchange genes with other such groups living in the same area. From this definition, one of the intrinsic properties required for a population to achieve species status is Reproductive Isolation (RI). The mechanisms of reproductive isolation can be categorized as preventing mating and fertilization (prezygotic) or contributing to the low fertility and survival of hybrid offspring (postzygotic) (Servedio and Sætre et al. 2003). Due to various criticisms directed to the BSC, Wu (2001) updated the concept by shifting the focus from RI at the genomic level to interspecific incompatibility at the genic level, specifically in genes that affect differential adaptation. Nevertheless, the BSC is not strictly followed today, as species are no longer viewed as discrete single units, but as more continuous standing points in the hierarchy of biodiversity, partially due to an ever increasing perception of the pervasiveness of hybridization and its importance in evolutionary biology (Mallet 2005).

### 1.2 Hybridization and domestication

Many authors argue that hybridization has a large beneficial impact on the evolution and diversification of taxa. One reason for this is that hybridization can act as a source of adaptive variation for diverging species, allowing the movement of alleles between genomes, with selection maintaining the differences in genes involved in reproductive isolation (Barton 2013). Hybridization can also lead to the formation of new species if the hybrid populations have improved traits over the parental types, a phenomenon known as heterosis, facilitating their successful establishment and adaptation to new environments (Rieseberg et al. 1999). New hybrid species may carry higher numbers of homologous chromosome pairs (allopolyploidy) than their diploid parents (Abbott et al. 2013). In plant evolution, polyploid hybridization is of major significance, with around 31% of ferns and 15% of flowering plant species having polyploidy origins (Mallet et al. 2007). As for animal species, allopolyploid hybrid speciation is more difficult, since most animals do not reproduce asexually like plants (Seehausen 2004), with fewer exceptions (e.g. Kobel and Pasquier 1986; Chenuil et al. 1999; Ereskovsky et al. 2007). There have also been instances of new hybrid animal species originating without changes in ploidy (homoploid), including butterflies (Beltrán 2007), ants (Schwander et al. 2007), crustaceans (Taylor et al. 2005), and the controversial case of the red wolf (vonHoldt et al. 2011).

Although hybridization has been associated with increased biodiversity and speciation, other researchers have also underlined that the formation of hybrids can have negative consequences for the survival of one or both parental species (Rhymer and Simberloff 1986, Wolf et al. 2001).

In general, hybridization can contribute to species extinction in two different ways in function of the fitness of the hybrids. The first is termed demographic swamping, and occurs when hybridization is common and hybrids have lower fitness than either parental species. In this scenario, hybrids may not pose a competitive and genetic threat to the parental species, but the hybridization represents wasted reproductive effort for individuals of the parental species and this can result in a reproductive output below the required replacement levels (Todesco et al. 2016). This process has been reported in pied flycatchers (*Ficedula hypoleuca*) on the Swedish island of Öland, which has been colonized by another species, the collared flycatcher (*F. albicollis*). (Wiley et al. 2009; Vallin et al. 2010). An alternative mechanism by which hybridization can be a threat to species survival is called genetic swamping. In this case, hybrids do not have a significantly lower fitness than the parental types, and hybridization may thus lead to the loss of one or both parental types (Todesco et al. 2016). Genetic swamping has been documented in several animal taxa (e.g. Nogueira et al. 2011; Derr et al. 2012; Mucci et al. 2012) and appears to be more common than demographic swamping (Ellstrand and Rieseberg 2016).

One of the main scenarios in which hybridization often has negative consequences is the arrival of invasive species that reproduce with native species and decrease their fitness. If this fitness reduction is extreme, it may threaten their survival (Mooney and Cleland 2001). This has been shown to be a common phenomenon and is most likely due to weak prezygotic barriers, as native and introduced species did not evolve together (Todesco et al. 2016). The number of alien species has been steadily increasing during the last 200 years (Seebens et al. 2017). A large part of this increase is due to human activities and interference, from commercial translocations of species to simplifying habitat structure, removing natural barriers to migration (Mooney and Cleland 2001). As a consequence of habitat homogeneity, previously isolated taxa may come into contact and hybridize (Seehausen et al. 2008).

In a survey of 143 empirical studies focused on hybridization, ranging from 1975 to 2015, around 48% of them highlighted hybridization as an extinction threat. For the studies reporting human involvement, 72% indicated the occurrence of at least one extinction (Todesco et al. 2016). Among the anthropogenic extinction threats is the occurrence of interbreeding, especially if frequent, between domestic and wild species (Crispo et al. 2011). Domestication is defined as the selective breeding of individuals originated from a wild ancestral species, in order to exploit or harvest a resource of interest (Terrel et al. 2003). Selective pressure for traits of interest to humans caused the domestic populations to become increasingly genetically distinct from their wild ancestors (Diamond 2002). Therefore, introgressive hybridization with domestic taxa can have some serious detrimental effects. Due to the in general extremely low genetic diversity in domestic populations, their interbreeding with wild species may reduce genetic variation in the latter. Perhaps even more importantly, traits artificially selected in domesticates may be maladaptive in natural environments and, thus, their introgression into wild populations can lead to outbreeding depression and decreased local adaptation (Crispo et al. 2011). Although in some instances, introgressive hybridization appears to not be yet a very serious threat, such as in the Italian population of grey wolves (*Canis lupus*), in which a study using microsatellites only detected 11 (5%) out of 220 genotyped individuals as admixed (Verardi et al. 2006), in other cases, like the hybridization between the common quail (*Coturnix c. coturnix*) and domestic Japanese quail (*C. c. japonica*) in Spain and Italy, it has had effects on phenotypic traits of the

former species, with migratory behaviour in particular having been observed to decrease in first-generation hybrids (Barilani et al. 2005).

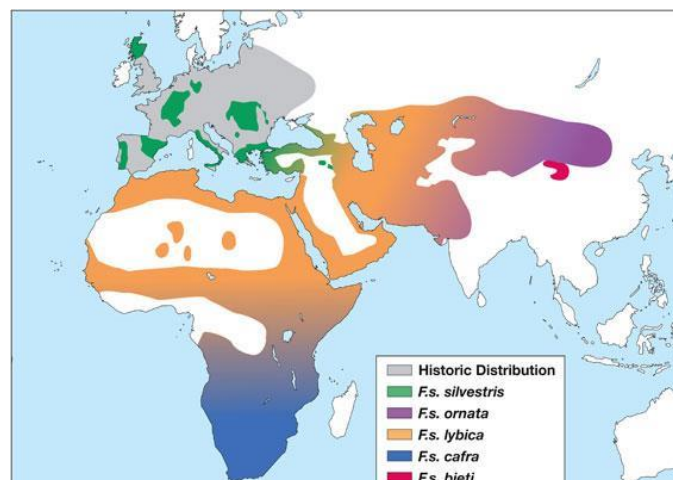
### 1.3. The wildcat

The wildcat (*Felis silvestris*) is a small carnivore belonging to the family Felidae, which comprises around 38 species distributed across the world, inhabiting diverse ecological niches that reflect different evolutionary adaptations (Sunquist and Sunquist 2002) (Fig.1). It is considered a polytypic species, with five recognized subspecies: *F. s. silvestris* in Europe, *F. s. cafra* in Southern Africa, *F. s. lybica* in North Africa and Middle East, and *F. s. ornata* and *F. s. bieti* in Central Asia (Fig. 2) (Driscoll et al. 2007). The domestic cat is sometimes also considered a subspecies, *F. s. catus*, since it is derived by domestication of wildcat (Driscoll et al. 2007). A global genetic survey and population structure analysis of wildcats and domestic cats revealed that the latter should have been domesticated in the Near East, being a descendant of domesticated *F. s. lybica* wildcats (Driscoll et al. 2007). The same study suggested that the domestication process started more than 9000 years ago, contemporary with the development of agriculture and farming in the Fertile Crescent (Driscoll et al. 2007). This is supported by archaeological findings from the site of Shillourokambos in Cyprus, in which a cat skeleton alongside a human skeleton were dated to about 9500 years ago (Vigne et al. 2004). Since then, selective pressures may have been minimal, given that cats were mostly used for rodent control and that most of the 30 to 40 cat breeds have been formed, for essentially aesthetic traits, within the last 150 years (Montague et al. 2014). This left the domestic cat with the same basic morphology as its wild ancestral species, with the main phenotypic difference being coat colour variability, which is mostly controlled by a relatively small number of genes (Pierpaoli et al. 2003).



**Fig. 1.** The European wildcat (*F. s. silvestris*) (photo from <http://www.wild-scotland.org.uk/>)

In order to reveal the genomic signatures underlying the domestication of cats, a study by Montague et al. (2014) used a whole-genome analysis of different domestic cat breeds and wildcats to find regions putatively affected by selection. They developed a new higher-quality reference assembly of the domestic cat genome, named FelCat5, which contained 2.35 gigabases (Gb) assigned to all 18 autosomal and X chromosomes. Identifying regions under positive selection can be complicated by the effect of random allele fixation during the formation of new breeds. This effect of genetic drift was mitigated by sequencing 22 domestic cats from six phylogenetically distant and geographically dispersed breeds. These sequences from the 22 domestic cats were then combined to reconstruct an “ancestral” domestic cat genome. Including also a pool of wildcat sequences from the European (*F. s. silvestris*) and Near East (*F. s. lybica*) subspecies, the sequences of domestic cat and wildcat were aligned separately to the FelCat5 reference genome for variant detection. The goal was to find regions with low levels of diversity, measured by pooled heterozygosity (Hp), and high levels of domestic cat - wildcat divergence, estimated by the fixation index (Fst). A total of five chromosomal regions fulfilled both these criteria, with 13 identified genes within these locations. Each of these genes plays important roles in neural processes, mainly in synaptic circuitry that influences behaviours like stimulus-reward learning, and neural crest cell migration (Montague et al. 2014). The results seem to be in concordance with the predictions of the domestication syndrome hypothesis, which posits that morphological and physiological traits are altered during domestication as consequences of mild neural crest cell deficits during embryonic development (Wilkins et al. 2014). Moreover, this study reinforces the notion that differences between wild and domesticated cats mostly concern aspects in behaviour like docility and tameness, which do not necessarily represent major prezygotic barriers to reproduction if individuals of the two species meet.



**Fig 2.** Geographic distribution of the five wildcat subspecies (Ramos 2014).

### 1.3.1. The degree of hybridization in European wildcat populations

The European wildcat represents one of the most well-known examples of anthropogenic hybridization. The subspecies suffered a drastic population decline and fragmentation in the 19<sup>th</sup> century (Enserink and Vogel 2006) and is, in general, currently restricted to relatively small isolated areas, usually densely forested but in proximity to human settlements (McOrist and Kitchener 1994). This has facilitated hybridization between wildcats and their domestic congeners. Being the offspring of such crosses fertile, this makes possible the introgression of domestic cat genes into the wildcat genome, potentially leading to genetic dilution and outbreeding depression (Rhymer and Simberloff 1996). Although the wildcat is currently classified as “Least Concern” by the IUCN Red List of Threatened Species (Yamaguchi et al. 2015), the same assessment underlines that hybridization and introgression in the European subspecies are of major concern and may have resulted in cryptic extirpations of some populations, and recommends further research on those topics.

Recently, researchers have attempted to assess the degree of hybridization in several European regions, and the incidence of this threat is variable between them (Virgós and Moleón 2014). In some areas there have been reports showing high levels of hybridization, as in Scotland (Kitchener et al. 2005) and Hungary (Pierpaoli et al. 2003). In contrast, in countries like Italy (Pierpaoli et al. 2003), France (O’Brien et al. 2009), and Portugal (Oliveira et al. 2008a), wildcat populations appear to be much less introgressed. Importantly, all these studies emphasized the need to continuously monitor hybridization rates and to develop simple but powerful methods that are able to distinguish wildcats from domestic cats and their hybrids.

### 1.4. Identifying pure and hybrid individuals: from morphology to genetics

Classical methods to assess the pure versus hybrid status, and determine the level of hybridization, of a specimen or population were based on the analysis of their morphology. For instance, Kitchener et al. (2005) analysed the pelage markings and skull measurements of 135 wildcats, domestic cats and hybrid specimens kept in Natural History Museums. They classified the specimens through the use of 20 pelage characters and 37 skull measures, each scored with an integer from 1 (domestic cat) to 3 (wildcat). Skull parameters were not able to separate nominal wildcats from hybrids. Also, no single pelage character was able to accurately distinguish any of the three groups of cats. The most reliable method to distinguish wildcats from hybrids and domestic cats was to use the top seven pelage markings (7PS) that exhibited less intra-group variation across groups, but the authors warned that using this system may not be practical in the field. Natural variation in pelage traits seems to be larger than previously thought, with cats classified as hybrids upon crude examination in the field, subsequently ascertained as wildcats following additional morphological and genetic analyses (Virgós et al. 2014). Accordingly, the development of genetic tests using molecular markers, coupled with data analysis using sophisticated Bayesian clustering algorithms, has vastly improved our ability to screen the genetic makeup of wildcat populations (Randi et al. 2001, Oliveira et al. 2015). However, there is still room for improved methods to reliably distinguish pure wildcats from feral domestic cats and their hybrids, and specially needed are assays that are accurate, cost-effective, and convenient to use (Kitchener et al. 2005, Virgós et al. 2014).

#### 1.4.1. Microsatellites

Microsatellites, also known as short tandem repeats (STRs), are molecular markers consisting of short runs of tandemly repeated sequence motifs (e.g. TCTCTCTCTC) abundantly and evenly distributed throughout eukaryotic genomes (Ellegren 2004). Microsatellite polymorphisms derive mainly from variability in length rather than in the primary sequence. With the advent of the Polymerase Chain Reaction (PCR) technique in the 1980s, the genotyping of microsatellite polymorphisms became straightforward and popular (Ellegren 2004). The increasing availability of large panels of highly polymorphic microsatellite markers, together with the use of model-based Bayesian methods for data analysis, provided improved detection of hybridization and hybrid zones (Vähä and Primmer 2006).

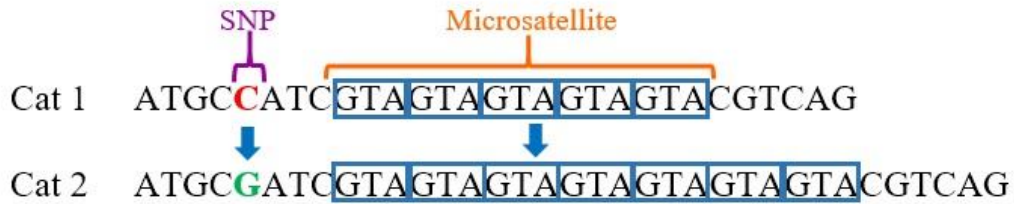
Their widespread use was also evident in the case of the wildcat. In a study conducted by Pierpaoli et al. (2003), a total of 211 wild and domestic cats, and 25 known or presumptive hybrids, were obtained from various localities in Portugal, Belgium, Switzerland, Italy, Germany, United Kingdom, Bulgaria, Slovenia and Hungary. Using 12 *F. catus* microsatellites and multivariate and Bayesian analysis, the study showed that wildcats are still genetically distinct from their domestic counterparts in most central and southwest European locations, but are extensively admixed in the UK and Hungary. A similar study focused on the hybridization levels in the Iberian Peninsula, with a sample size of 75 Portuguese and Spanish wildcats, and 109 feral or purebred domestic cats, estimated that at least 11.4% of the Portuguese wildcats showed signs of introgressive hybridization (Oliveira et al. 2008a; Oliveira et al. 2008b).

However, microsatellites have some issues such as null alleles and size homoplasy, due to their high mutation rate, that introduce ambiguity in data analysis (Morin et al. 2004), and thus optimal marker selection is critical (Vähä and Primmer 2006, Oliveira et al. 2008b).

#### 1.4.2. Single nucleotide polymorphisms

When comparing genomes from individuals of the same or very closely related species, they will be nearly identical, but one of the key differences between them are variations of a single nucleotide at a specific position in the genome. These are called “Single Nucleotide Polymorphisms” (SNPs), which are highly abundant in the genome, for example occurring about once every 1,000 bases in the human genome (Syvänen 2001). SNPs are mostly biallelic, and cannot be more than tetrallelic (Nussberger et al. 2013). As such, they have less variability than STRs. Nevertheless, and overall, SNPs have several advantages over STRs, namely: (1) higher density and more uniform distribution along the genome, (2) less homoplasy as a consequence of a much lower mutation rate, (3) better suited for high-throughput genotyping, (4) easier comparison of results between labs, with less extensive protocol standardization and genotype scoring calibration, (5) Highly applicable to samples with degraded DNA, such as non-invasive or historical samples (Oliveira et al. 2015). These make them highly desirable molecular markers for different types of studies, ranging from estimating population structure (Marchini et al. 2004) and genetic variation (Morin et al. 2004) to phylogeography (Emerson et al. 2010). The use of SNPs in studies of admixture in wild populations is on the rise but still relatively limited (Oliveira et al. 2015). Notably, a study using SNPs to investigate the hybridization between two carp

species estimated hybridization rates that were four times higher than those identified using microsatellites, which was interpreted as suggesting a greater diagnostic power for SNPs (Lamer et al. 2015).



**Fig. 3.** Examples of mutation in a single nucleotide polymorphisms (SNP) and in a microsatellite locus.

The study by Nussberger et al. (2013) was the first that attempted to obtain a set of diagnostic nuclear SNP markers to identify European wildcats, domestic cats, and their hybrids. The authors created a reduced representation library (RRL) by sequencing a small random part of the genome of six wildcats and three domestic cats, and identified 48 SNPs showing high levels of differentiation between the species ( $F_{st} > 0.8$ ). An additional of 42 wildcats and domestic cats were genotyped for these 48 SNPs, and used as reference parental genotypes for simulating hybrids. Around 97.3% of all simulated individuals were assigned to their correct hybrid category. However, 9% of parental domestic cats and 18% of parental wildcats were misclassified as third generation hybrids (Nussberger et al. 2013). The main limitation of the study is that it used a geographically restricted set of wildcat samples (Oliveira et al. 2015), all originating from a local population in Switzerland (Nussberger et al. 2013). This is an issue because the European wildcat population has dwindled significantly and become highly fragmented, and the current patterns of genetic diversity and structure are still poorly known. Analysis of parental individuals sampled from a limited geographical area is likely to under-represent variability of the parental species and overestimate their differentiation (Oliveira et al. 2015). Hence, ascertainment bias may affect the utility of the SNP panel in other European wildcat populations.

The more recent study by Oliveira et al. (2015) tried to address the concern for ascertainment bias, by sampling cats across the distribution of the European wildcat. A total of 107 putative wildcats, collected from nine locations far from each other, and 121 domestic cats were analysed. The SNPs used were a combination of markers discovered in previous studies (Kurushima et al. 2012, Lyons et al. 2005, Johnson et al. 2006, Esteves et al. 2007). The results of the admixture analysis showed that the 158 SNPs were successful in identifying F1, F2 and backcross hybrids. The authors also showed that reducing the marker set to the top 35 SNPs with the highest  $F_{st}$  values did not result in losing too much inference power. However, none of the 158 SNPs were fixed between the two species, with only 22 markers being monomorphic in the wildcat sample (Oliveira et al. 2015). Also, the average of the  $F_{st}$  values for the top 35 SNPs was 0.74, which is lower than that for the larger panel of 48 SNPs (average  $F_{st} = 0.89$ ) in the study of Nussberger et al. (2013).

Despite the major contributions represented by those two studies, the search is still on for a set of SNP markers that are fixed or extremely differentiated between European wildcats and domestic cats, so that it has power to identify pure wildcats and hybrids across Europe, but that at the same time is small enough to allow the development of a simple genotyping assay that is convenient, rapid and inexpensive.



## 1.5. Objectives

This thesis had the following objectives: i) survey previous literature to select candidate genes that were impacted by positive selection during the domestication of the cat and other mammals; ii) amplify these candidate genes in wildcat and domestic cat samples originating from Portugal, France and Romania; iii) identify possible diagnostic SNPs during sequence analysis; and iv) assess the diagnostic power of the SNPs through assignment tests of observed and simulated genotypes.

## 2. Materials and Methods

### 2.1. Cat reference samples

Analyses were conducted on 16 tissue samples from domestic cats and wildcats (Table 1). Domestic cat samples were collected in the municipal catteries of Lisbon (2) and Porto (3). Portuguese wildcat samples (4) were obtained from the Tissue Bank of Wild Vertebrates of the Portuguese Institute of Nature and Forest Conservation (BTVS-ICNF), and the purity of the four wildcat samples has been suggested by the results of the study by Oliveira et al. (2008b). Lastly, samples of wildcats from France (4) and Romania (3) were kindly provided, respectively, by Sébastien Devillard and Sandrine Ruetten and by Zsolt Hegyeli; the purity of the French samples is supported by the results of Devillard et al. (2013) and the purity of the Romanian samples is indicated by morphological analyses carried out by Zsolt Hegyeli using the diagnostic definitions proposed by Kitchener et al. (2005). Total genomic DNA was extracted from tissue using the EZNA Tissue DNA kit (Omega Bio-Tek) following the manufacturer's protocol, and then stored at -20 °C.

**Table 1.** Information about the wildcat and domestic cat samples used in this study.

Code number	Sample Label	Species	Country of Origin
1	99-0081	<i>Felis silvestris</i>	Portugal
2	02-0325	<i>Felis silvestris</i>	Portugal
3	GP9	<i>Felis catus</i>	Portugal
4	GL10	<i>Felis catus</i>	Portugal
5	02-0100	<i>Felis silvestris</i>	Portugal
6	03-0072	<i>Felis silvestris</i>	Portugal
7	GP5	<i>Felis catus</i>	Portugal
8	GL8	<i>Felis catus</i>	Portugal
9	Fs3915	<i>Felis silvestris</i>	France
10	Fs39H	<i>Felis silvestris</i>	France
11	ZH4	<i>Felis silvestris</i>	Romania
12	ZH8	<i>Felis silvestris</i>	Romania
13	ZH58	<i>Felis silvestris</i>	Romania
14	GP3	<i>Felis catus</i>	Portugal
15	Fs39O	<i>Felis silvestris</i>	France
16	Fs39E	<i>Felis silvestris</i>	France

## 2.2. Candidate genes selection

As stated above, the study by Montague et al. (2014) indicated that genes with roles in neuronal development were among the most involved in cat domestication. Signs of positive selection were also detected in genes associated with sensory perception (e.g. vision and hearing). On the basis that coat colour is one of the most notable phenotypic differences between wildcats and domestic cats, even though artificial selection for coat colour in domestic cats has mostly occurred in the recent past for aesthetic reasons (Lyons 2015, Montague et al. 2014), genes controlling coat colour were considered as possibly eligible for inclusion in the study. Thus, in the search for candidate genes potentially involved in cat domestication, the focus was on those with known associations in domestic cat and/or in other domestic mammals to: (i) neuronal development; (ii) sensory system, specifically olfaction, vision and hearing; and (iii) coat coloration and hair length. In practical terms, the survey concentrated on the literature reporting genes putatively linked to these biological processes and phenotypic traits in the cat and, to increase the number of candidate genes tested, other domestic mammals (e.g. dog *Canis familiaris*, horse *Equus caballus*, pig *Sus scrofa*, goat *Capra hircus*, and sheep *Ovis aries*).

The Ensembl genome browser (Yates et al. 2016) was used to (i) verify that, in the case of candidate genes selected from studies on other species, the genes have been sequenced and annotated in the cat, (ii) locate the genes in the genomes of the cat and other domestic mammals, and learn the exon-intron structures, and (iii) export the gene sequences. The software GENEIOUS (Kearse et al. 2012) was used to align the sequences of each gene and design primers complementary to highly conserved regions. All primers were designed with a length of 18 nucleotides, 40-60% G+C content, and each primer pair amplifying 700-1200 base pairs (bp). For each of the 51 genes, two pairs of primers were constructed, with each pair targeting a different gene region and thus different sets of exons and introns (Table 2). For convenience during the laboratory work, two sets of primers were defined (A and B), each containing a primer pair for each gene.

## 2.3. PCR amplification

In general, after testing and optimization, PCRs were carried out in volumes of 15  $\mu$ l with 1 x PCR buffer (NZYTech), 1.7 mM MgCl<sub>2</sub>, 0.1 mM of each dNTP (Bioline), 0.3  $\mu$ M of each primer, 1 U of NZYTaQ II DNA polymerase (NZYTech), and 3–5 ng of DNA template. Thermal cycling conditions consisted of an initial denaturation at 94 °C for 3 min, followed by 35 cycles of 30 s at 94 °C, 30 s at Ta, 30 s at 72 °C, and a final extension of 7 min at 72 °C. Ta is the locus-specific annealing temperature and ranged from 46 to 63 °C. Initially, the annealing temperature tested for all primer pairs was 55 °C, but this was inadequate for many of them and optimization was required to obtain good PCR results. In several cases it was necessary to use stepdown PCR, a variant of PCR that aims to reduce nonspecific background by incrementally lowering the annealing temperature as PCR cycling progresses. This procedure can increase primer specificity and maximise yield (Hecker and Roux 1996). The results of the PCR amplifications were visualized on 2% agarose gels to verify PCR quality, and the PCR products were purified with an Exo-SAP protocol (Hanke and Winke, 1994; Werle et al., 1994) and sequenced at Macrogen Inc.

## 2.4. Preliminary analyses of sequences and SNPs

Sequences were edited, assembled and aligned using BioEdit v. 7.0.5 (Ibis Therapeutics) and Sequencher v. 4.2 (Genes Codes Corporation). Identification of fixed and highly divergent SNPs between wildcats and domestic cats was done by eye.

Preliminary analysis of each gene fragment involved amplification and sequencing of two wildcats and two domestic cats from Portugal. Fragments containing SNPs possibly fixed between the two species were analysed in other samples of Portuguese wildcats and domestic cats. Finally, fragments in which SNPs continued to appear fixed between the two species after increasing their sample size for Portugal, were further analysed in wildcats from France and Romania to check for ascertainment bias.

If a fragment contained more than one fixed or highly divergent SNP, only the one with the highest  $F_{st}$  was included in the subsequent statistical analyses, and the others were removed from the dataset to avoid using very tightly linked markers. Estimates of  $F_{st}$  were calculated using GENEPOP v.4.7.0 (Rousset 2008). The information content of each SNP was also evaluated by computing the “informativeness for assignment” ( $I_n$ ) using INFOCALC 1.1. (Rosenberg et al. 2003; Rosenberg 2005).  $I_n$  provides a measure of potential for assignment of an allele to one population compared with the “average” population; for a given set of populations ( $K$ ), the minimal  $I_n$  of 0 occurs when all alleles have equal frequencies in all populations, and the maximal value,  $\ln K$ , occurs when no alleles are shared between populations.

## 2.5. Population structure and admixture analysis

The SNPs with the highest  $F_{st}$  were evaluated in their capacity to identify population structure and hybrid classes. These assessments were made through Bayesian clustering methods, namely STRUCTURE v. 2.3.4 (Pritchard et al. 2000) and NewHybrids 1.1 Beta3 (Anderson and Thompson 2002). In STRUCTURE, initial population structure analysis was run with  $K$  values ranging from 1 to 3, with 15 replicate runs for each  $K$ . For each run, the average proportion of membership ( $Q$ ) of the sampled populations and the distribution of individual membership proportions ( $q_i$ ) to the two inferred clusters, with their 90% credibility intervals (CIs), were estimated. Parameters were set with the admixture model and correlated allele frequencies, and a burn-in length period of 100,000 followed by 500,000 Markov Chain Monte Carlo (MCMC) iterations. The most likely  $K$  value was inferred using the Evanno method (Evanno et al. 2005) in STRUCTURE HARVESTER 0.6.94 (Earl and vonHoldt 2012). The respective 15 runs were averaged using the program CLUMPP v.1.1.2 (Jakobsson and Rosenberg 2007), and the averages were graphically displayed with DISTRUCT v. 1.1 (Rosenberg 2004).

Subsequently, using the domestic cat and wildcat samples as reference genotypes, hybrids were simulated with HYBRIDLAB 1.0 (Nielsen et al. 2006). The program creates multilocus hybrid genotypes by randomly selecting one allele from each of the two parental populations, according to their frequency distribution (Nielsen et al. 2016). The diagnostic power of the markers to correctly identify the parental and hybrid status of the genotypes was then examined in NewHybrids (Anderson and Thompson 2002). NewHybrids estimates the posterior probability of each genotype belonging to one of the following six classes: domestic cats (D), wildcats (W), first generation hybrids (F1), second generation hybrids (F2), backcross with domestic cat (BD), backcross with wildcat (BW).

### 3. Results

#### 3.1 Candidates Genes

From the literature search for candidate genes potentially affected by cat domestication, a total of 51 genes were selected for this study. Of these, about 72% have roles in neuronal development, 16% in hair structure and colour, and 12% in the sensory system (Table 2).

**Table 2.** List of the 51 candidate genes analysed in this study. In the columns from left to right are the gene name (the letter in superscript indicates the consulted source, and these are listed below the table), biological role (N, neuronal development; C, coat colour; S, sensory perception), primer set (A or B), forward and reverse primers (5' to 3'), samples analysed (according to the sample numbers in Table 1), and the status of the analysis (FA: failed amplification; NE: not enough samples analysed to determine the presence or not of SNPs; NS: enough samples analysed to conclude that the fragment does not contain fixed or highly divergent SNPs between wildcats and domestic cats; PS: enough samples analysed to conclude that the fragment contains one or more SNPs potentially fixed or highly divergent between the species).

Gene Name	Role	Set	Primer Sequences (5' - 3')	Samples	Status
ACMSD <sup>d</sup>	N	A	F: GGTAACCTCTGAGTGCA R: TGTGTAGCATGTCATCAG	2,3,4	NS
		B	F: GCTGATAGGTTGCCTCTG R: CCTCCCTGTGAAGTGCA	1,3,4	NS
ADAMTS20 <sup>i</sup>	C	A	F: CCTGGTGTATCTGGACCT R: CACTAGCTCAACATCTGA	2	NE
		B	F: TCTGACTCTACTTGACTC R: TTAGTGCACCTTTCACGT	3,4	NE
ARID1B <sup>d</sup>	N	A	F: CCTGTAAAAGCACATCAG R: CTGAGATTAACAGGAGCT	2	NE
		B	F: CCCGTGTGAGGTTATGAC R: TCAGCACCAAGCCTGGAT	2,6	NE
ARID3B <sup>a</sup>	S	A	F: AGGTCAGAACACAGAAGT R: GACTGACCTGCTTGAGCT	1,2,3,4	NS
		B	F: TGGATTATTCTCCTCGCT R: ACAGAGAACAGGGCACTG	None	FA
ASIP <sup>b</sup>	C	A	F: CAGAGTGGATGCCATCAC R: GCAACCTAAGTAGCTATG	None	FA
B3GALTL <sup>f</sup>	N	A	F: GCTATGTCTGATTCTGCT R: ATGTGGTCACAGCTCGCA	2	NE
CDH5 <sup>l</sup>	N	A	F: CCAGTGAGGCTAAGCTAG R: CGTAAACACTCACAGTACA	1,2,3,4	NS
CLK3 <sup>a</sup>	N	A	F: ACCTCGGGACACAGACTT R: GCCCAAATCATACTGGA	None	FA
		B	F: GCCTAGATGCATCACTGT R: GAGGTGAGGCCTTTAAGC	2,3,4,5,6,7,8	PS
CNTRF <sup>g</sup>	N	A	F: GCTCAGAAAGTGTCTCAG R: CCTACTTGACTTACCAG	2	NE
COL9A3 <sup>a</sup>	S	A	F: AAGGAGACACAGGCCTGC R: CAGATGTTCTCACGGCAT	None	FA
		B	F: ACTCGAGAACCCATTTCAG R: TCACTTGCTTCAGTTGCG	8	NE
CYFIP1 <sup>d</sup>	N	A	F: AGCCTGCCATGGCTCATA R: AGTTGGTGTACACCTTATG	7	NE
CYP1A1 <sup>a</sup>	N	A	F: TCTCAGCAGCCATCTTGA R: CAGCCTGTACCTTCTCAA	2	NE
		B	F: CCTCATGTACCTGGTGAC R: TAAGCTGCAGGGCTCTCA	1,2,4,5,6,14	PS
CYP1A2 <sup>a</sup>	N	A	F: TGAATGGTCCATCTTGA R: ACCTCGTCAAAGTCCTGG	3,4	NE
		B	F: ACCATTTGCAAGGCCAAT R: CTGAAGCTGGTTGGTCAG	1,2,3,4	NS

<b>DCC<sup>a</sup></b>	N	A	F: GGATGCTTTCCAAAGACC R: ATTTGGAAGCACAACCTGC	1,2,3,4,5,8,12, 13,15,16	PS
<b>DDC<sup>e</sup></b>	N	A	F: CATCCTGTTTTGCTAACTG R: GGCTCAGCATGTTTGCAC	1,2	NE
<b>EDC3<sup>a</sup></b>	N	A	F: AGTTCATCCGTGGTGTAC R: TCTCCTAAAGGGGTAGAC	1,2,3,4,5,7,8	NS
<b>FAM114A2<sup>a</sup></b>	N	A	F: GTTGCATGTGTTTCCTAG R: GCTTGGAGACCAACCATT	1,2,3,4	NS
		B	F: CATGATGCTAACGTGTTC R: CTTGAACCAATGTGCATC	1,2,5,6,7,14	PS
<b>GALR1<sup>d</sup></b>	N	A	F: GTGCTGTTCGGCCTGATC R: CCGACTGCAAGTAACTT	None	FA
		B	F: TGTTGAGGCAGCCGTTCA R: AGTTGCAGGCTTGCTAAC	1,3,5,14	PS
<b>GNAQ<sup>i</sup></b>	C	A	F: ATCCAGAAGCTAGGTAGC R: ACACATTCACACAGAGTC	1,3,4,8	NE
<b>GRIA1<sup>a</sup></b>	N	A	F: AACTCAGATATGGCCTAG R: GTGAGTTGGAAATTAGCC	1,2,3,4,5,6,7,8	NS
<b>GRIA2<sup>a</sup></b>	N	A	F: TCTGTGTGATTGTATGCC R: GGTGTCGGAAGACTTATG	1,2,3,4,5,6,7,1 2,13,15,16	PS
<b>GRIK3<sup>d</sup></b>	N	A	F: CCAGATGAGGAGACAGGT R: CAGCTAGTGAGCGTCAGA	1,2,4	NE
<b>GRM8<sup>c</sup></b>	N	A	F: AGGTGGAAGACATGCAGT R: CCAGAGAAGGGTTGGCAT	1,2,4	NS
<b>HELLS<sup>i</sup></b>	C	A	F: CTGATAGAGGGTAGTAGC R: TGCAACTCTTGATCTCAG	4	NE
<b>HIPK2<sup>d</sup></b>	N	A	F: ACACCTGCTGTCACCTCCA R: TGTGCCGCATCTTCAGTA	None	FA
		B	F: ACTCTTGAATGGAGATCG R: AGAGCCAAGGCGATGCTA	4	NE
<b>HTR3A<sup>i</sup></b>	N	A	F: GATGTGAGGTCTGTCTGG R: AGAGACTCTCCACCACGT	1,2,4	NS
<b>KRT71<sup>b</sup></b>	C	A	F: TCCTCACTGCAAGCTGCA R: TCCTCATAGCTAACCTGA	1,2,3,4,5,6,7,9 ,12,13,14, 15,16	PS
<b>LPAR6<sup>b</sup></b>	C	A	F: AACTGTAAACTGGACGT R: AGAGGTGCCAGTGAAGT	1,2,3	NS
<b>MATN2<sup>f</sup></b>	N	A	F: CGTGGAGATTGGAGGTCT R: CCTTGCCAACACAGTACC	2,3	NE
		B	F: AGAGTACGGTGTGGACCT R: TCTAGACTAACAGCACAG	6	NE
<b>MFAP3<sup>a</sup></b>	N	A	F: AGGTGGTCTCCACTCAC R: GACAATAGCTCCTTGTC	1,2,3,4,5,8,12, 13	PS
<b>MITF<sup>i</sup></b>	C	A	F: AGATGCAATCAAGCTGAC R: GTTGCAACAGGCACCTGT	2,3	NE
<b>MLPH<sup>b</sup></b>	C	A	F: TAGTTGAGCGCCATAGGA R: CTGCTCTCTCAGACACGT	1,2,3,4	NS
<b>MYO15A<sup>a</sup></b>	S	A	F: ACTCCGCTCACCTTTGCT R: AAGATCCCGAGGCTGTTC	2,3	NE
		B	F: TCTCCAAGTCCCAGAGCT R: TAATCGTCGTAGCCGTAGC	None	FA
<b>MYO7A<sup>a</sup></b>	S	A	F: ATAGAGAGGCTGGAGGCT R: TACCAGGGTGAAGTCAGC	1,2,3	PS
<b>NINJ1<sup>f</sup></b>	N	A	F: CGGTGCCTACACTTCACT R: AAGGACCACAGCTAGTC	None	FA
		B	F: TGAGCTTCCGTTCTTGGA R: CACATAGGCAGTGTTTCAG	None	FA
<b>NPFFR2<sup>i</sup></b>	N	A	F: ACAAGGCAGTGTCCCACT R: GAGGGCTTTCAGTACATA	1,2	NE
<b>OR10K1<sup>a</sup></b>	S	A	F: AAGTCTTGCTGCCTGCA R: GATTCTAGGCATCCCATT	1	NE
<b>OR2B11<sup>a</sup></b>	S	A	F: TTGACTCATCACAGGCAC R: CTCAGAGCCTAATCTGCA	1,2,3,4,5,6,7,8	NS
<b>PCDHB4<sup>a</sup></b>	N	A	F: ACATCCTCATGACAGAAG R: CACCACTTAAAAGTGCCT	None	FA

		B	F: CCTGAGCTTACCATATCT R: AGAGGTTGGGAAATATCG	None	FA
<b>PKP4<sup>g</sup></b>	N	A	F: TCTGTAATGACAGTGGAC R: CAGCCTGAAGTCAGCAGT	1,2,3,4	NS
<b>PLEKHH1<sup>a</sup></b>	N	A	F: AACCACAGCTCTTCAGAT R: GAGGATGAGAGCCTCATC	1,2,3,4,5,7,8,1 2, 13	PS
<b>PPFIA4<sup>g</sup></b>	N	A	F: AGGTTACAAGAGTCTGAG R: AAGGGGAGGTTCCAGCAT	2	NE
<b>PTPRQ<sup>a</sup></b>	S	A	F: ACCTACCATAGTACCCGA R: GATTCAGGCTTACCTGAC	1,2,3,4	NS
		B	F: TCACTAGTCTATGTCAG R: CTTTGGTTTCTCAAGGTC	2,3,4,5	PS
<b>RNF103<sup>d</sup></b>	N	A	F: GCTTCATGATTGAGGATC R: CTGGACAGCTCTCGAAGT	2	NE
		B	F: TTATTTCTGGCTACCTG R: GCAAGCTGTAAGATACTC	2,3,4,5	NS
<b>SH3GL2<sup>d</sup></b>	N	A	F: ACTCATCAATGGTTCCT R: GAAACCACAGGACAGTCA	1,2,3,4,5,6,7,8 ,12,13,16	PS
<b>SLC6A4<sup>c</sup></b>	N	A	F: AATCCGCTGCTGCTGTA R: ACGCACTAAGGAGGCTGA	1,3,4	NS
		B	F: TGCTCAGGCCGTTTCATG R: GGAAGCTTTCAGTTAGT	1,2,3,4,5,6,7,8	PS
<b>SPOCK1<sup>h</sup></b>	N	A	F: TGTTCAAACGGCTTGAT R: TCTTCTGCCTGATCTCCA	1,2,4,5,8,14	PS
<b>SYNJ2<sup>f</sup></b>	N	A	F: GCTGGCCTTTACTGACT R: AAGAGCTGTGGTCCATGC	None	FA
<b>TLX3<sup>d</sup></b>	N	A	F: CAGTGTCAACCTGAGCCT R: TAGCGGGTAGCACAGAAT	None	FA
<b>VEZT<sup>d</sup></b>	N	A	F: CACTTTGAGAACCACTGC R: GTGGATTATCTAGCCTTG	1,2,3,4,5,6,7,8 ,12,13,14, 15	PS
<b>YWHAH<sup>d</sup></b>	N	A	F: TGGATTCAGATCTCAGTG R: TGGCAAGGAAGAATCAGT	2,3,4	NS

a – Montague et al. 2014

b – Lyons 2015

c – Wang et al. 2013

d – Axelsson et al. 2013

e – Cagan et al. 2015

f – Schubert et al. 2014

g – Moon et al. 2015

h – The Bovine HapMap Consortium 2015

i – Dong et al. 2015

### 3.2 SNP discovery

The results of the PCR and sequencing analyses of the first set of primers (set A) yielded that the primer pairs for 28 genes failed to amplify, or did so inconsistently and results were obtained for only a few samples, in both cases preventing any search for SNPs. For another 14 genes, enough samples were sequenced so that a preliminary SNP exploration could be performed, but no potentially useful SNPs were found. For the remaining nine genes, analysis revealed potentially discriminating SNPs between the Portuguese samples of the two species. Of these nine gene fragments, seven were sequenced in Romanian samples, and five in both Romanian and French samples. Since there was a large number of unsuccessful primer pairs (28) and gene regions without interesting SNPs (14), a second primer pair was designed for some of these genes; in the case of the latter group of genes, this second primer pair targeted a different gene region than the first primer pair. This second set of primer pairs (set B) targeted fragments of 18 genes. Amplification was unsuccessful for nine, three did not contain relevant SNPs, and the remaining six harboured potentially useful SNPs (Table 2).

Hence, potentially discriminatory SNPs were found in 15 genes (CLK3, CYP1A1, DCC, FAM114A2, GARL1, GRIA2, KRT71, MFAP3, MYO7A, PLEKHH1, PTPRQ, SH3GL2, SLC6A4, SPOCK1 and VEZT). However, for MYO7A only one domestic cat sequence was produced (sample GP9), and tentative support for the SNP could only be obtained by the addition of the reference domestic cat genome sequence (FelCat5, Montague et al. 2014) in the alignment. Therefore, conservatively, this gene was not included in further analyses. Additionally, sample Fs3915, a wildcat from France, could be sequenced for only one gene, KRT71, during the timeframe of this thesis. For this reason, this sample was removed from the data set prior to subsequent analyses. It is worth noting that most SNPs were located in intronic regions, with only three SNPs located in exons (Table 3).

### 3.3 Genetic Variation

Genetic variability analysis showed that for most of the 14 genes, SNPs were fixed for different alleles in the two species (Table 4). The exceptions were KRT71 and CYP1A1, for both of which one wildcat was heterozygous (samples ZH8 and 02-0325, respectively), GRIA2, for which one wildcat (sample 03-0072) was homozygous for the “domestic allele”, and FAM114A2, for which one domestic cat (sample GP5) was homozygous for the “wild” allele. The  $F_{st}$  values indicated that the SNPs are potentially useful in discriminating wildcats and domestic cats, since for almost all of the SNPs the values were above 0.8. The only exception was for FAM114A2 ( $F_{st} = 0.58$ ). The  $I_n$  scores supported the indications from the  $F_{st}$  values, with most SNPs achieving the maximum possible score for  $K = 2$  (i.e.,  $\ln(2) = 0.693$ ), with the lowest value belonging to FAM114A2. Given these results, the FAM114A2 SNP was not included in the admixture analysis.

**Table 3.** Summary of characteristics of the discovered SNPs that were found to be highly differentiated between wildcats and domestic cats. In the columns from left to right are the genes where the SNPs were identified, genomic location of the SNPs (chromosome and nucleotide position in the *F. catus* genome from Ensembl, FelCat5), the annealing temperatures used in the PCR (for most genes a stepdown protocol was performed), whether the SNPs are exonic or intronic, the alternative nucleotides at each SNP, whether the exonic SNPs are synonymous or nonsynonymous, and the amino acids associated with the alternative allele at each exonic SNP.

Gene	Location	T <sub>a</sub> (°C)	Region	Locus	Mutation	Amino acid
VEZT	B4_118813134	55 - 53	Exon	G/T	Nonsynonymous	V - G
SH3GL2	D4_42617102	55 - 53	Intron	G/A	-	-
DCC	D3_71413722	62 - 60	Exon	G/A	Synonymous	A
PLEKHH1	B3_144899186	55 - 53	Exon	C/T	Synonymous	L
MFAP3	A1_193825899	55 - 53	Intron	G/A	-	-
SLC6A4	E1_16571212	57 - 55- 53	Intron	T/C	-	-
CLK3	B3_31690860	55 - 53	Intron	A/T	-	-
SPOCK1	A1_113638109	57 - 55- 53	Intron	G/T	-	-
GARL1	D3_91747200	55 - 52 - 49	Intron	A/G	-	-
PTPRQ	B4_103742634	55	Intron	T/C	-	-
KRT71	B4_78971988	62 - 60	Intron	C/G	-	-
CYP1A1	B3_31603213	58 - 53	Intron	A/G	-	-
GRIA2	B1_70880038	55 - 53	Intron	A/G	-	-
FAM114A2	A1_193305654	57 - 55- 53	Intron	G/A	-	-

V – valine; G – glycine; A – alanine; L - leucine

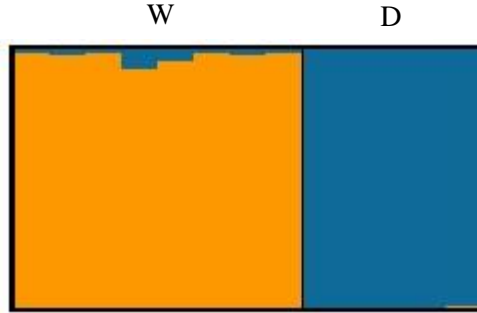
**Table 4.** Information about the diagnostic SNPs identified in this study. In the columns from left to right are the genes where the SNPs were identified, the “wildcat” allele (“p”), the “domestic cat” allele (“q”), the number of wildcats genotyped (“nW”), the frequency of the “wildcat” allele in the wildcat samples (“pW”), the frequency of the “domestic cat” allele in the wildcat samples (“qW”), the number of domestic cats genotyped (“nD”), the frequency of the “wildcat” allele in the domestic cat samples (“pD”), the frequency of the “domestic cat” allele in the domestic cat samples (“qD”), the  $F_{st}$  values between wildcats and domestic cats, and the  $I_n$  scores.

Gene	p	q	nW	pW	qW	nD	pD	qD	$F_{st}$	$I_n$
<b>VEZT</b>	G	T	8	1	0	4	0	1	1	0.693
<b>SH3GL2</b>	G	A	7	1	0	4	0	1	1	0.693
<b>DCC</b>	G	A	7	1	0	3	0	1	1	0.693
<b>PLEKHH1</b>	T	C	5	1	0	4	0	1	1	0.693
<b>MFAP3</b>	G	A	5	1	0	3	0	1	1	0.693
<b>SLC6A4</b>	T	C	4	1	0	4	0	1	1	0.693
<b>CLK3</b>	A	T	3	1	0	4	0	1	1	0.693
<b>SPOCK1</b>	G	T	3	1	0	3	0	1	1	0.693
<b>GARL1</b>	A	G	2	1	0	2	0	1	1	0.693
<b>PTPRQ</b>	T	C	2	1	0	2	0	1	1	0.693
<b>KRT71</b>	C	G	9	0.94	0.06	4	0	1	0.91	0.693
<b>CYP1A1</b>	A	G	4	0.88	0.12	2	0	1	0.81	0.537
<b>GRIA2</b>	A	G	8	0.87	0.13	3	0	1	0.80	0.497
<b>FAM114A2</b>	G	A	7	1	0	3	0.33	0.67	0.58	0.318

### 3.4 Performance of the SNP panel to distinguish wildcats from domestic cats

In order to quantify the performance of the selected panel of 13 SNPs to discriminate wildcats and domestic cats, Bayesian clustering analysis in STRUCTURE was conducted without prior population information. The optimal number of clusters was  $K = 2$  with wildcats and domestic cats being clearly separated into two distinct clusters, with all individuals correctly assigned to their putative species of origin (Fig. 4). When the assignment analysis was repeated assuming  $K = 2$ , all individuals had an estimated membership coefficient ( $Q$ )  $> 0.95$  to their cluster of origin, thus showing no signs of admixture, with the sole exception of a Portuguese wildcat (sample 03-0072) that had  $Q = 0.94$  and a lower limit of the 90% confidence interval below 0.75 (Table 5). Nonetheless, the  $Q$ -value of individual 03-0072 in its cluster of origin is still higher than those observed in individuals considered to be pure in previous studies of hybridization in wildcats using SNP markers (Oliveira et al. 2015). Overall, the final 13 individuals were considered as reliable reference wildcat and domestic cat samples for further analysis.





**Fig. 4.** Bar plot of STRUCTURE results, averaged across 15 replicate runs, for  $K = 2$  genetic groups. The analysis was based on 13 SNPs and included 13 individuals (eight wildcats, W, and five domestic cats, D). Each individual is depicted by a column that is partitioned into two ( $= K$ ) segments, which length is proportional to the ancestry probability of the individual to each group, coloured in orange (W) and blue (D).

**Table 5.** STRUCTURE's estimates and 90% confidence intervals (CI) for the membership proportions of each individual (Q) in, respectively, the wildcat (W) and domestic cat (D) clusters.

Sample code	STRUCTURE's Q			
	$Q_w$	90% CI on $Q_w$	$Q_D$	90% CI on $Q_D$
<b>99-0081</b>	0.997	(0.983,1.000)	0.003	(0.000,0.017)
<b>02-0325</b>	0.988	(0.931,1.000)	0.012	(0.000,0.069)
<b>02-0100</b>	0.998	(0.986,1.000)	0.002	(0.000,0.014)
<b>03-0072</b>	0.936	(0.737,1.000)	0.064	(0.000,0.263)
<b>ZH8</b>	0.965	(0.822,1.000)	0.035	(0.000,0.178)
<b>ZH58</b>	0.995	(0.973,1.000)	0.005	(0.000,0.027)
<b>Fs390</b>	0.992	(0.941,1.000)	0.008	(0.000,0.046)
<b>Fs39E</b>	0.994	(0.959,1.000)	0.006	(0.000,0.041)
<b>GP9</b>	0.003	(0.000,0.017)	0.997	(0.983,1.000)
<b>GL10</b>	0.003	(0.000,0.014)	0.997	(0.986,1.000)
<b>GP5</b>	0.005	(0.000,0.030)	0.995	(0.970,1.000)
<b>GL8</b>	0.004	(0.000,0.023)	0.996	(0.977,1.000)
<b>GP3</b>	0.008	(0.000,0.049)	0.992	(0.951,1.000)

### 3.5 Admixture Analysis

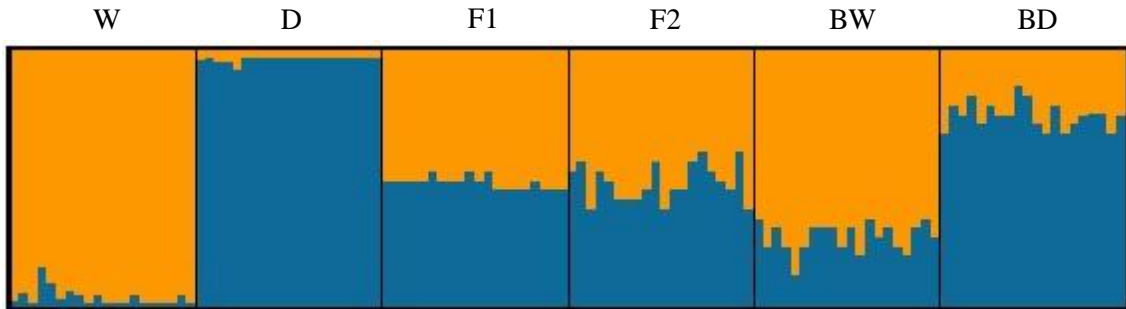
To further assess the power of the panel of 13 SNPs to identify pure and hybrid individuals, 13 hybrid F1 genotypes were simulated using HYBRIDLAB and the genotypes of the 13 reference wildcats and domestic cats. The 26 genotypes were analysed in NewHybrids to estimate their posterior probabilities of belonging to the parental and F1 classes. The results showed that all genotypes, both observed and simulated, were assigned to their correct class with a mean posterior probability  $> 0.99$  (Table 5). The performance of the panel was also investigated including a second generation of hybridization in a HYBRIDLAB simulation generating 20 cat genotypes of each of the six following categories: pure parental species, first-generation hybrids (F1), second-generation hybrids (F2), and backcrosses to one parental species or another. Similarly to the previous analysis, all genotypes were correctly assigned to their category with a mean posterior probability  $> 0.95$  (Table 5).

Admixture analysis was also conducted in STRUCTURE with the same 120 genotypes used in the second NewHybrids experiment, and enforcing  $K = 2$  (Fig. 5). STRUCTURE clustered the genotypes in a similar fashion to NewHybrids. The average membership coefficients were  $> 0.95$  for each parental category. All genotypes of the parental and F1 classes were correctly

assigned, and the second-generation hybrids and backcrosses also had high percentages of correctly assigned individuals, with 80% or more for each class (Table 7). Thus, in general, the panel had excellent diagnostic power to distinguish pure wildcats from domestic cats and F1 hybrids, and a remarkable ability to even discriminate different hybrid categories.

**Table 6.** NewHybrids results for the mean posterior probabilities of belonging to pure and hybrid categories. N indicates the number of individuals used for each category. Categories are: parental domestic cat (D), parental wildcat (W), first-generation hybrids (F1), second-generation hybrids (F2), backcrosses into domestic cat (BD), and backcrosses into wildcat (BW).

Experiment	Category	N	NewHybrids					
			D	W	F1	F2	BD	BW
Until first generation	D	5	<b>0.999</b>	0.000	0.001	-	-	-
	W	8	0.000	<b>0.999</b>	0.001	-	-	-
	F1	13	0.001	0.001	<b>0.998</b>	-	-	-
Until second generation	D	20	<b>0.995</b>	0.000	0.000	0.000	0.005	0.000
	W	20	0.000	<b>0.978</b>	0.001	0.003	0.000	0.018
	F1	20	0.000	0.000	<b>0.997</b>	0.001	0.001	0.001
	F2	20	0.000	0.000	0.001	<b>0.978</b>	0.009	0.012
	BD	20	0.005	0.000	0.001	0.017	<b>0.980</b>	0.000
	BW	20	0.000	0.003	0.000	0.039	0.000	<b>0.958</b>



**Fig. 5.** Bar plot of STRUCTURE results, assuming  $K=2$  and averaged across 15 replicate runs, for six genotypic classes (20 genotypes for each class). W - pure wildcats, D - pure domestic cats, F1 - first-generation hybrids, F2 - second-generation hybrids, BW - backcrosses with wildcat, and BD - backcrosses with domestic cat.

**Table 7.** Mean and (within brackets) range of STRUCTURE's estimates for the membership proportions of each individual belonging to six different genotypic categories in, respectively, the wildcat (W) and domestic cat (D) clusters. The categories, each with 20 individuals, are: parental domestic cats (D), parental wildcats (W), first-generation hybrids (F1), second-generation hybrids (F2), and backcrosses with domestic cats and wildcats (BD and BW, respectively). The last column shows the percentage of correctly assigned individuals (%N).

Categories	STRUCTURE		%N
	Q <sub>D</sub>	Q <sub>W</sub>	
<b>D</b> <b>Q &gt; 0.80</b>	0.975 (0.909, 1.000)	0.025 (0.000, 0.091)	100
<b>W</b> <b>Q &gt; 0.80</b>	0.043 (0.002, 0.128)	0.957 (0.872, 0.998)	100
<b>F1</b> <b>0.40 &lt; Q &lt; 0.60</b>	0.492 (0.335, 0.648)	0.508 (0.352, 0.665)	100
<b>F2</b> <b>0.40 &lt; Q &lt; 0.60</b>	0.491 (0.337, 0.646)	0.509 (0.354, 0.663)	80
<b>BD</b> <b>0.60 &lt; Q &lt; 0.80</b>	0.756 (0.614, 0.877)	0.244 (0.123, 0.386)	85
<b>BW</b> <b>0.60 &lt; Q &lt; 0.80</b>	0.278 (0.148, 0.425)	0.723 (0.575, 0.852)	95

#### 4. Discussion

The study of hybridization and its implications for evolution and conservation has increasingly become a major research area in biology. For instance, it is now widely recognized that hybridization poses a serious threat to species already under pressure from other factors (Todesco et al. 2016). This is due to the fact that often hybrids are fertile and can generate second-generation hybrids and backcross to parental species. A specific issue is the case of hybridization between feral individuals of domestic species and individuals of their wild ancestral species. Artificial selection alters phenotypic traits and likely makes them less suited to the natural environments inhabited by the wild ancestor. However, this artificial selection is still relatively recent, and in general has not led to the establishment of reproductive barriers between domestic and wild forms (Vilà et al. 2003). As such, the introgression of domestic traits into wild populations can hinder their survival due to outbreeding depression and genetic dilution (Leonard et al. 2014). This has been documented between dogs and wolves (Vilà and Wayne 1999, Verardi et al. 2006, vonHoldt et al. 2013). While initially wolf-dog hybrids were thought to be of little concern, with an estimated hybridization frequency of 1% in Scandinavian populations, based on mitochondrial DNA and autosomal markers (Vilà and Wayne 1999, Vilà et al. 2003), but higher levels of admixture have been detected in Italy, with 5% of identified hybrids (Verardi et al. 2006), and in the Iberian Peninsula with 4% of identified hybrids (Godinho et al. 2011). The authors of these studies have suggested that it is likely that the extent of hybridization has been underestimated and emphasized the need for improved markers, especially in terms of detecting backcrosses more effectively. Accordingly, vonHoldt et al. (2013) attempted to develop a panel of diagnostic SNPs able to detect mixed ancestry within up to four generations of wolf-dog hybridization. They performed a genome-wide scan that revealed 48,000 SNPs between 155 grey wolves and 912 domestic dogs. After filtering and ranking the SNPs through principal components analysis (PCA), a panel consisting of 100 unlinked SNP markers was created, which provided good resolution until F2 hybrids, with 82% of individuals assigned to the correct genotype class. For applications requiring more precise determination of wolf-dog ancestry, they

recommended the use of the full 48K SNP marker set. None of the SNPs were fixed between species, and the authors warned that finding such perfectly diagnostic markers could be an unrealistic goal.

The wildcat (*Felis silvestris*) represents another example of a wild species highly impacted by anthropogenic activities, including human-facilitated hybridization with feral domestic cats. This is illustrated by the case of the Scottish wildcat (*F. silvestris grampia*), which has been affected by extensive introgressive hybridization, with essentially all examined wildcats having at least some domestic cat ancestry (Beaumont et al. 2001). This also makes difficult to define a “pure wildcat” diagnosis based on morphological characters, due to a large overlap between species (Kitchener et al. 2005). In the study by Beaumont et al. (2001), using microsatellites, 28% of cats belonging to a non-domestic genetic group had domestic pelage characters. Yamaguchi et al. (2004) estimated that the proportion of Scottish wildcats failing to be identified as such on the basis of coat traits could be as high as 40%. Given this, recent discoveries of the underlying genetic makeup that distinguishes wildcats from their domestic congeners (Montague et al. 2014) provide a great opportunity to develop improved molecular assays for discriminating wildcats, domestic cats and their hybrid offspring.

#### 4.1 Candidate genes with diagnostic SNPs

Previous studies have attempted to pinpoint the mutations that explain the variation in phenotypic traits between the different cat breeds, especially those affecting coat colour and hair length. To date, around 26 genetic variants from at least 11 known genes have been documented to cause phenotypic variations. Some of these variants are caused by single nucleotide substitutions but others are due to the deletion or insertion of sequence segments (Lyons et al. 2015). However, the usefulness of these mutations for discriminating wildcats, domestic cats and their hybrids is uncertain. Many of these mutations, or at least their high frequencies, may be relatively new in domestic cats because artificial selection for aesthetic traits, particularly those related with pelage types, became widespread only recently. Over the last 140 years, since the first cat show in London in 1871, a plethora of different breeds have been created by man, with 57 breeds currently recognized by the International Cat Association (Kurushima et al. 2012). Most of this artificial selection has been imposed on purebred breeds, and therefore the selected mutations are likely to be rarer in random-bred cats. On the other hand, as mentioned above, a high frequency of genetically wildcat-like cats, as indicated by molecular markers, have domestic cat pelage characteristics (Yamaguchi et al. 2004), suggesting that mutations responsible for coat traits may be easily introgressed in hybridizing populations. In agreement, preliminary analyses of SNPs in coat trait genes did not show promising results in terms of distinguishing between wildcats and domestic cats (C Fernandes, pers. comm.). Nevertheless, Oliveira et al. (2015) used a few SNPs for such traits in their admixture analysis, and some of them were among the highest ranked in diagnostic power of the 158 SNPs in the panel.

Besides genes affecting coat traits, another intuitive set of potentially relevant candidate genes are those involved in neuronal processes. This stems from considering the combination of traits that are frequently modified during the domestication process, which are referred as the “domestication syndrome” (DS) (Wilkins et al. 2014). The list of traits that comprise the DS is varied and includes depigmentation, curly tails, short muzzles, delay in sexual maturity, and increased docility. This last trait, docility, is the only one in the DS that is common among all domestic mammals. This is most probably an effect of strong selection for tameness, a reduced

fear and aggression towards humans, in the initial stages of domestication (Wilkins et al. 2014). Tameness of domestic mammals has been associated to a reduction in the size and function of the adrenal glands, which regulate many physiological responses to stress and fear (Trut et al. 1999). The formation of the adrenal glands during embryonic development is linked to a group of cells called neural-crest cells (NCC), a vertebrate-specific group of embryonic cells that first appear at the junction between the neural and epidermal ectoderm, and then migrate from the dorsal region of the neural tube to give rise to diverse cell lineages (Hall 2008). As such, a recent hypothesis postulates that the initial selection for tameness led to a reduction in the proliferation or differentiation of neural crest cells, which in turn induced phenotypic changes in traits of the “domestication syndrome”. While this hypothesis is not accepted by all, evidence in its favour has been derived from studies on the domestic cat, since several genes implicated in neuronal cell migration (e.g. ARID3B, DCC, PLEKHH1) were inferred to be under positive selection when comparing the genomes of wildcats and domestic cats (Montague et al. 2014). Thus, several grounds support the idea that genes with a role in neuronal development may contain markers distinguishing the two species. In general, because selection on traits of the domestication syndrome seems to have occurred in most domestic mammals, it also makes sense to look for differentiation between wildcats and domestic cats in genes known to be associated with selected traits, other than coat characteristics and neuronal development, in the domestication of other mammalian species (e.g. dog, pig, horse).

Of the primer pairs designed to amplify and sequence fragments of the 51 selected candidate genes, those for 22 of the genes did not amplify at all or amplified inconsistently, even after several repeat attempts with different thermal cycling conditions, so that sequence data was not obtained from enough individuals to allow reliable searches for SNPs. Therefore, the question of whether these 22 genes may contain diagnostic SNPs between wildcats and domestic cats requires further examination. For the primers that worked well, this was sometimes achieved by reducing the final MgCl<sub>2</sub> concentration in the PCR reaction, since *Taq* DNA polymerase activity is sensitive to the concentration of magnesium ions, and very often by using a step down PCR protocol (Hacker and Roux 1996), which in general improved primer specificity and product yield.

In this study, of all the SNPs that were discovered, those found in 13 of the candidate genes (CLK3, CYP1A1, DCC, GARL1, GRIA2, KRT71, MFAP3, PLEKHH1, PTPRQ, SH3GL2, SLC6A4, SPOCK1, and VEZT) were highly differentiated ( $F_{st} > 0.8$ ) between wildcats and domestic cats, in a degree comparable to the SNP markers reported by Nussberger et al. (2013). Furthermore, the candidate gene approach in this study yielded 10 fixed SNPs ( $F_{st} = 1$ ) between wildcats and domestic cats, while seven such SNPs were identified by Nussberger et al. (2013) using random sequencing of about 2% of the genome. Thus, well-informed and carefully designed candidate gene studies can be useful to identify diagnostic SNPs to distinguish between hybridizing species. The candidate gene approach has been a successful strategy for identifying functional mutations causing phenotypic changes in non-model organisms (Aitken et al. 2004; Slate et al. 2009). Here, using candidate genes which previous research suggested as having undergone positive selection during domestication of mammals provided a targeted and effective means to identify diagnostic SNPs. The  $I_n$  scores from Infocalc were also indicative of the diagnostic power of the panel developed in this study, with most SNPs reaching the maximum possible value of  $I_n$ .

Several of the diagnostic SNPs found in this investigation are located in genes detected to be under positive selection in the study by Montague et al. (2014) on the genetic signatures

underlying cat domestication (e.g. DCC, PLEKHH1, MFAP3, CLK3, CYP1A1, GRIA2, PTPRQ). MFAP3, CLK3 and CYP1A1 have roles in neuron circuitry and metabolism, while DCC and PLEKHH1 have functions specifically related to neural crest cells migration, and PTPRQ and GRIA2 have functions related to the sensory system. Although the SNP identified in the gene FAM114A2 did not show a very high  $F_{st}$  ( $= 0.58$ ), this gene is also located in one of the regions identified by Montague et al. (2014) as under positive selection. The gene KRT71 was not identified as under positive selection in the same study (Montague et al. 2014), but previous studies in cat, dog, mouse, and rat, have identified mutations in KRT71 that cause effects on hair structure (Gandolfi et al. 2010). Namely, a variant localized in c.445-1 of the coding DNA sequence (c.445-1G>C) disrupts the conserved splicing site of intron 1, and seems to be linked to the curly hair coat phenotype of the Selkirk Rex cat breed (Gandolfi et al. 2013). Also, another splice variant polymorphism, a substitution of an adenine by a guanine near intron 4 (c.816+1G>A), was suggested to cause the hairless phenotype in the Sphynx breed (Gandolfi et al. 2010). The SNP identified here between wildcats and domestic cats is located in the intron between exons 1 and 2, near the mutation in the locus *Rexing* of the Selkirk Rex cats. However, because the curly hair mutation is specific to the Selkirk Rex breed, it is unlikely that there is a linkage between this mutation and the SNP identified here.

Interestingly, most other genes with diagnostic SNPs (GALR1, SH3GL2, SLC6A4, and VEZT) were candidate genes from studies of dog domestication. The SLC6A4 gene was found to be potentially under positive selection in genome comparisons between dogs of different breeds and grey wolves (Wang et al. 2013). Like most genes suggested being under positive selection in the domestic cat lineage by Montague et al. (2014), SLC6A4 has a role in neurological function, specifically coding for a serotonin transporter. Increased levels of serotonergic receptor have been associated with aggressiveness in dogs (Badino et al. 2004). Similarly, the genes GALR1, VEZT and SH3GL2, inferred to be under positive selection in the dog lineage in another genomic study comparing various dog breeds and wolves (Axelsson et al. 2014), have functions related to synaptic plasticity and modulation of action potentials (Axelsson et al. 2014).

A common feature of the 13 diagnostic SNPs was that most were intronic (Table 3), albeit closely flanked by one or two exons. Introns are less bounded by functional constraints than exons, and therefore tend to exhibit higher nucleotide diversity, making them preferred regions where to search for neutral markers (Slate et al. 2009). Even though these mutations may not have been targets of positive selection, their high levels of fixation between the two cat species may indicate that these intronic SNPs are closely linked to sites that were influenced by positive selection, and underwent a process known as genetic hitchhiking (Barton 2000). However, there were also SNPs found in exons, which are the parts of the gene that encode the mature messenger RNA (mRNA) and, as such, are generally highly conserved sequences (Ryynänen and Primmer 2006). Both SNPs in DCC and PLEKHH1 are synonymous mutations, meaning that they do not result in an amino acid change (Sauna and Kimchi-Sarfaty 2011). In contrast, the SNP in VEZT is a nonsynonymous missense mutation and, therefore, associated with an amino acid change. This type of mutation is most often linked to functional changes in proteins and the fitness of an organism (Kryukov et al. 2007). Recently, a study compared the performance of intronic versus exonic SNPs in revealing population differentiation within a widespread bird species, and showed that functional exonic SNPs outperformed intronic markers in this task (Zhan et al. 2015).

## 4.2 Power of the SNP panel to detect hybridization

The panel of the 13 diagnostic SNPs discovered in this study was able to successfully identify hybrid genotypes simulated based on the allele frequencies of the reference samples of wildcat and domestic cat. When only adding first generation hybrids to the empirical data set, all individuals were assigned to their category of origin with  $> 0.99$  posterior probability. With the further inclusion of second-generation hybrids, the panel still maintained a high accuracy, as individuals were assigned to the correct category with posterior probability  $> 0.95$ . The only significant decrease was in the wildcat parental class, with two individuals (03-0072 and ZH8) assigned to their correct class with posterior probability  $< 0.90$ . However, both of these animals were only genotyped at seven of the 13 markers. Moreover, sample 03-0072, a Portuguese wildcat, possessed the domestic cat allele in the GRIA2 gene, and the Romanian sample ZH8 was heterozygous for the KRT71 SNP. Both the not wild-like genotypes at these genes and the missing data for the other six loci likely contribute to explain their lower probability of assignment to the correct category.

When comparing the results obtained with the panel developed here with those for the 48 SNP marker set reported by Nussberger et al. (2013), the latter assigned reference parental cats and simulated hybrids to their respective six genotype classes with a higher probability in all cases. In their study over 97% of individuals were assigned to their correct hybrid category with  $> 0.95$  posterior probability, whereas in the present study the percentage was 90%. However, while Nussberger et al. (2013) only analysed samples from Switzerland, here we examined wildcats from Portugal, France, and Romania. Moreover, it is remarkable that the SNP panel presented here is much smaller than that of Nussberger et al. (2013), and therefore more practical and convenient (being this one of the objectives of this work), but still provides an extremely high accuracy and power.

In comparison to the results of Oliveira et al. (2015), the panel of 13 SNPs was able to assign parental domestic cats ( $Q > 0.995$ ) and wildcats ( $Q > 0.978$ ) to their correct category with a posterior probability similar to their panel of 158 SNPs ( $Q > 0.999$  for domestic cats;  $Q > 0.994$  for wildcats) in NewHybrids. For the hybrid categories (F1, F2, BD, BW), the panel of this study possessed higher assignment values for all hybrid categories. In STRUCURE analysis, both panels performed similarly for the parental and F1 categories, with every individual of each category being correctly assigned. However, for the remaining categories (F2, BD, and BW) the average assignment probabilities were lower in this study than in Oliveira et al. (2015). In the latter, using only the top 35 diagnostic SNPs identified by the authors, showed 0%, 8%, and 4% of misassignments in the simulated F2, BD, and BW, respectively; the corresponding percentages in the present study were 20%, 15%, and 5%. Using all 158 SNPs, they had no misassignments across all six categories. All of the 158 SNPs investigated by Oliveira et al. (2015) were polymorphic in the domestic cat samples, and only 22 SNPs were monomorphic in the wildcat. By comparison, in this study the 13 SNPs were monomorphic in the domestic cat, and only three SNPs (KRT71, GRIA2, and CYP1A1) were polymorphic in the wildcat. Having more reference parental samples polymorphic at more SNPs may result in less distinct simulated genotypes among the different genotype classes, since for instance HYBRIDLAB generates hybrid genotypes by randomly drawing one allele at each locus, as a function of the respective estimated allele frequency distributions (Nielsen et al. 2006). Also, it is important to note that the SNPs discovered in the present study were genotyped for different numbers of individuals, ranging from four (GARL1, PTPRQ) to more than 10. This may, for example, create a bias because a

monomorphic marker will cause estimated allele frequencies to appear fixed at opposite extremes, but the observed monomorphism may be an artefact due to the small number of samples genotyped. Clearly, comparison of results between this study and those of Nussberger et al. (2013) and Oliveira et al. (2015) must be done with caution, especially because here the number of analysed individuals was much smaller.

### 4.3 Future perspectives

The purpose of this study was to be a pilot effort towards developing an accurate, cost-effective, and convenient set of diagnostic SNPs for discriminating pure wildcats, domestic cats and their hybrids. Examining candidate genes suggested by previous studies to be involved in the process of domestication of the cat, dog, and other mammalian domesticates, permitted to discover an initial set of 13 markers showing high differentiation between the two cat species and that was found to have high power to detect hybrids and distinguish different hybrid classes (F1, F2 and backcrosses).

However, this preliminary work needs to be continued and expanded. For example, given that the designed primers failed to amplify, or did so inconsistently, almost half of the candidate genes selected for this study, new primers need to be designed and tested, following suggestions from the literature (e.g. Dieffenbach et al. 1993; Cha et al. 2014), for the same or different regions of these genes. Also, in general for all the gene regions containing potentially diagnostic SNPs, and especially for those that were sequenced in very few individuals (GALR1, PTPRQ, MYO7A, and FAM114A2), it is necessary in future research to analyse more specimens to confirm the extremely high differentiation observed between wildcats and domestic cats at these SNPs, and to allow more confident analyses of admixture using larger sample sizes. It is worthwhile to note that the reason why a few of the gene fragments with highly differentiated SNPs were only sequenced for a very few individuals was that those SNPs were identified late in the period allocated to laboratory work, and this precluded analysis of more samples in the time frame of this thesis. To further ascertain that the set of diagnostic SNPs is useful for identifying pure and hybrid wildcats throughout the range of the subspecies *silvestris*, it would be important to test wildcat and domestic cat samples from throughout the extant range of the subspecies. As there has been no or little research on hybridization between the two species outside Europe, an interesting future extension of this work would also be to determine whether the SNP panel is able to distinguish between wildcats and domestic cats across the rest of the distribution area of the former species, particularly in the Middle East, the area of origin of the domestic cat (Driscoll et al. 2007). Finally, for the missense SNP in the VEZT gene, if its fixation or near fixation between the two cat species is verified by further studies, it would be relevant to investigate the potential effect of the amino acid change in the structure and function of the encoded protein, through bioinformatics tools, like PROVEAN (Choi et al. 2012) that predicts the functional impacts of protein sequence variation, and experimental approaches such as CRISPR-Cas9 (Dow et al. 2015) and ectopic expression (Liu et al. 2017).



#### 4.4 Concluding remarks

The European wildcat *F. s. silvestris* is a mammalian mesocarnivore with important ecological roles, for instance being the top predator of the food chain in some of the ecosystems in which it occurs (Roemer et al. 2009). The continued habitat destruction and population decline have confined the species to small and disconnected fragments of its former range. The increasing human impacts on its habitats have also increased the probability of it encountering its domestic counterpart and, consequently, of hybridization and introgression, eroding its genetic integrity and fitness (O'Brien et al. 2009). Already there are signs of extensive admixture in central and northwestern Europe (Pierpaoli et al. 2003, Kitchener et al. 2005). To reliably assess and monitor this issue, methods that are able to accurately differentiate between pure and hybrid wildcats are pressingly needed. The increased use of molecular tools, such as microsatellites and SNPs, has improved hybrid detection, but the development of efficient and cost-effective assays remain a priority because they would facilitate the efforts of individuals and agencies involved in the conservation of the wildcat.

In this study, using a candidate gene approach, SNPs were searched in a selection of genes suggested by previous studies to be involved in the process of domestication of the cat and other mammalian domesticates, and a set of 13 highly divergent SNPs ( $F_{st} > 0.8$ ) between wildcats and domestic cats was identified. This panel allows the accurate discrimination between the two species and the detection of their hybrids, including the distinction between different hybrid classes. Pending the above-mentioned extensive research that is necessary to validate the panel, this new marker set may provide a convenient and reliable assay to accurately characterize hybridization and introgression patterns in different wildcat populations across Europe.

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