

Population Genetics of the European Roma—A Review

Giacomo Francesco Ena¹, Julen Aizpurua-Iraola¹, Neus Font-Porterias², Francesc Calafell¹, and David Comas^{1,*}

- ¹ Department of Medicine and Life Sciences (MELIS), Institute of Evolutionary Biology (CSIC-UPF), Universitat Pompeu Fabra, 08002 Barcelona, Spain
- ² Department of Biomedical Informatics, Anschutz Medical Campus, University of Colorado, Aurora, CO 80309, USA
- * Correspondence: david.comas@upf.edu

Abstract: The Roma are a group of populations with a common origin that share the Romani identity and cultural heritage. Their genetic history has been inferred through multiple studies based on uniparental and autosomal markers, and current genomic data have provided novel insights into their genetic background. This review was prompted by two factors: (i) new developments to estimate the genetic structure of the Roma at a fine-scale resolution have precisely identified the ancestral components and traced migrations that were previously documented only in historical sources, clarifying and solving debates on the origins and the diaspora of the Roma; (ii) while there has been an effort to review the health determinants of the Roma, the increasing literature on their population genetics has not been subjected to a dedicated review in the last two decades. We believe that a summary on the state of the art will benefit both the public and scholars that are approaching the subject.

Keywords: population genetics; genomics; Romani; genetic structure; bottleneck; founder event



Citation: Ena, G.F.; Aizpurua-Iraola, J.; Font-Porterias, N.; Calafell, F.; Comas, D. Population Genetics of the European Roma—A Review. *Genes* 2022, 13, 2068. https://doi.org/ 10.3390/genes13112068

Academic Editor: Luisa Azevedo

Received: 6 October 2022 Accepted: 3 November 2022 Published: 8 November 2022

Publisher's Note: MDPI stays neutral with regard to jurisdictional claims in published maps and institutional affiliations.



Copyright: © 2022 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (https:// creativecommons.org/licenses/by/ 4.0/).

1. Introduction

The Romani population constitute the largest transnational ethnic minority across the European continent, and while they are commonly known to be traditionally itinerant, most of them have been settled for centuries. Despite their long presence in numerous countries and the assimilation into part of the local culture to different extents, Romani groups share a common identity and cultural traditions [1,2]. For centuries, most of the information regarding Romani history originated from linguistics, written historical records from non-Romani populations and socio-anthropological comparisons, although recent insights have come from the characterization of their genetic background [3–7]. According to historical findings [2,8], this nomadic state of living, including cultural and social practices (for instance, endogamy) that are uncommon in the dominant European populations, has led to a cultural friction which very often resulted in social marginalization and persecution. Due to this, a significant number of the Roma might have decided not to register their ethnicity in official censuses [9]. In 2019, Romani people in Europe were estimated to number 10 million [10]; however, some Romani and international human rights organizations declare that they exceed 14 million, meaning that they constitute 1.5–1.8% of the total European population [10]. Most Romani populations are established in South-Eastern Europe, while other major groups are found in Central and Western European countries. Other Romani live outside of the European continent, in particular in the Middle East and in the Americas [2,8]. The Roma are a complex, structured population and considering just the national origin of its members usually fails to properly characterize this diversity. Understanding Romani structure requires the concerted efforts of many disciplines, as well as the involvement of the Romani public. For instance, the main linguistic units within

the Roma are based on seven main dialect groups: Balkan, Vlax, Central, North-Eastern, North-Western, British and Iberian [11,12].

Linguistic and historical research has provided evidence that the Roma originated in the northern regions of South Asia, in particular the Punjab and Kashmiri regions of northern India [5,11]. Anthropological records have shown resemblances between the cultures of different Indian groups and Romani people [13], who have a similar social structure to the jatis of India where groups are usually defined by profession and the endogamous group constitutes the primary unit [2,14]. The origins of the Roma have been a subject of debate for a long time and hypotheses on their Indian origins were already made in the 19th century [15–18]. However, it was as a result of studies based on autosomal [5,7] and uniparental markers [19–21] that their genetic South Asian origin was corroborated, which was also supported by previous evidence of Mendelian mutations that are shared by the Roma and individuals of South Asian ancestry [22,23]. Because their history of migrations, bottlenecks and endogamy practices have led to the presence of a number of founder mutations [23] and sharing of unique Mendelian disorders [3,22], as well as founder mitochondrial DNA (mtDNA) and Y-chromosome (Y-DNA) lineages [6], the Roma are described as a group of founder populations with a relatively complex demography and history [21,24].

Historical records show that the Roma were initially welcomed with mixed feelings in most of the European kingdoms and became known as "Egyptians" and locally derived terms, such as 'astigani', 'tsigane' and 'gitanos', leading to misconceptions regarding the origins of the population. The first descriptions of the Roma included a variety of differently skilled workers, such as farmworkers, blacksmiths, craftsmen and mercenary soldiers, as well as musicians and entertainers [2,8,25]. Due to changes in the political climate, between the 15th and 16th century, the Roma were engulfed by a wave of intolerance, often forced to settle or to leave the countries in which they were travelling [26–28]. When Romani individuals did not accept these new imposed laws, they were arrested, enslaved, deported or killed [29,30]. Since then, and until the second part of the 19th century, they lived in slavery in many parts of western, central and eastern Europe [8,31]; the persecutions culminated into the genocide that occurred during the Second World War, in which it is estimated that the Nazi regime and its allies exterminated around half of the Romani population of Europe [32–37].

Romani people nowadays are full-right citizens in the European Union, but despite improvements in their living conditions, there is still an anti-Romani sentiment which has proven to be a difficult prejudice to overcome [38]. The Roma still suffer many inequalities, with high rates of unemployment and poverty, low participation in education and a poor health status [39,40], and governments are still slow in taking action against their discrimination [38,40].

2. Origins of the Roma from Genetic Evidence

The first genetic evidence of the Indian origins of the Roma came from studies based on blood group genetic markers [41–43] and founder mutations in the γ -Sarcoglycan gene [44,45]. The presence of several disease-causing Mendelian mutations [22], which can be explained by founder events, such as the congenital myasthenia caused by the 1267delG mutation in the CHRNE gene, also supported the Indian origin of the Roma, as this form of inherited disorder was otherwise described only in patients of Indian and Pakistani genetic ancestry [23,46]. Other mutations, such as the W24X mutation present in the GJB2 (connexin 26) gene related to non-syndromic hearing loss or the mutations present on the LTBP2 gene, were found in the Roma [47–49] and are known to be present at a high frequency in a number of Indian populations [50].

However, most of the genetic evidence regarding the South Asian origin of the Roma has been provided by the analyses of uniparental lineages (mtDNA and Y-DNA), as the presence of South Asian lineages in Romani groups pointed to an Indian origin of the proto-Roma [3]. Regarding mtDNA, M haplogroups (such as M5a1, M18 and M35b),

which originated in South Asia, are present in Romani groups [21]. One of the most distinct maternal lineages found in the Roma is the M5a1b1a1 haplogroup that, as expected from a founder lineage, has been observed at a high frequency but low diversity [51], and whose diversity in the Roma has been dated around 1.5 kya [51]. Other strong evidence on the Indian origin comes from Y-DNA data due to the high frequency of the Y haplogroup H1a-M82 [19,52], which is originally from North-Western India [53]. While studies based on uniparental DNA agree on the estimating dates of the origins [51,53], some have contradicted each other about the exact location within the Indian subcontinent in which the proto-Roma originated [20,21]. A study suggesting that southern India could have also contributed to the gene pool of the proto-Roma, based on the exact matches of Y-DNA H haplotypes [20], contradicted previous reports based on mtDNA and sparked an intense debate. However, as some authors noted [4], the Indian sampling of the study in question did not include groups from the north-western regions, biasing the results. More recent findings have proven that Dravidian-speaking populations from South India are also involved in the South Asian source of the Romani individuals [7], appearing to solve the contradiction regarding the identification of uniparental Roma lineages with a North-Western Indian origin and the high level of Y-STR haplotype sharing among the Roma and South Indian populations.

Based on genome-wide SNP arrays and whole-genome sequences, it has been determined that the Romani people carry approximately 20–35% South Asian ancestry [4,7], and North-West India constitutes the major source of this component [4,7,54]. The South Asian genetic ancestry is complex and can be decoupled into two components: Ancestral North Indian (ANI) and Ancestral South Indian (ASI) [55]. The former is related to the admixture events that occurred in South Asia around 2–4 kya from West Eurasian migrations [56] and, as a result, the Roma may have already carried an Ancestral West Eurasian (AWE) component derived from the South Asian ANI prior to their arrival into Europe [4,7]. The general consensus on the origins was eventually reached with the results coming from analyses of uniparental markers and whole-genome sequences, which pinpointed the origins in north/northwestern India 1.5 kya [5,6,57], in agreement with the dates obtained from uniparental markers [21,52]. Additionally, these results proved that all the analyzed Romani people descended from a common single founder population [5–7], as previously suggested by Kalaydjieva et al. [24].

3. The Romani Diaspora out of India

Historical and genetic sources agree on the fact that after the proto-Roma left North-Western India, they migrated through the Middle East, mostly through Persia (in the Iranian Plateau) and the Caucasus, Armenia in particular [4,8,13,21]. Such evidence suggests the occurrence of a southern Black Sea migration through Anatolia into the Balkans [11,58], which might account for the near absence of Arabic and the existence of a few lexical items of Georgian derivation in the Romani language [11]. At the same time, a more northerly route through the Caucasus and Crimea into Eastern Europe has been deemed as unlikely by historical sources, due to the fact that the first evidence of the Roma in Russia appears rather late, already into the 16th century [13]. Nonetheless, there is no agreement on the amount of gene flow that the Roma received during their diaspora until reaching Europe. Some genome-wide studies have suggested a long stay of Romani groups in the Middle East with substantial gene flow from local populations [59], whereas others have proposed a moderate Middle East and Caucasus gene flow during the migration across these territories [5,7]. The migration through the Middle East and Caucasus is supported by the presence of uniparental haplogroups, which most likely have a Middle Eastern and Caucasus origin, having their highest frequency in these present-day regions, such as the mtDNA U3 haplogroup in Iran and Lebanon [60] or the Y-DNA J2 haplogroup in Ingushetia and Chechnya [61]. Despite this, Middle Eastern admixture is probably overestimated due to its genetic similarity with European-related ancestry, and the presence of high-frequency

uniparental haplogroups from this region in the Roma could be explained by the occurrence of high levels of genetic drift.

It has been historically suggested that the Roma reached the Anatolian peninsula between the 11th and 12th centuries [2,8,27] and the Balkans as early as the 13th century [8,31]; the dating of these events is also supported by genetic studies [4,5,7,57] (Figure 1). The Balkan genetic footprint in the Roma is well documented [4,57,59] and there is evidence of an ancestry gradient that correlates with the distance to the Balkans. In fact, the Balkan genetic component varies from 45% in Bulgarian, Greek and Serbian Roma to 25% in Baltic and Iberian Roma, which is further evidence that the dispersion into Europe took place via the Balkans [5,7]. After subsequent migrations and dispersal across Europe, Romani groups eventually reached Northern Europe and the Iberian Peninsula, the western-most part of the continent, in the 15th century [5,8]. We can speculate that one of the earlier migration waves of the Roma that started from the Balkans is the one that generated the Polish, Lithuanian and Iberian Roma [21], because only these three subpopulations show a high frequency of mtDNA U3 and Y-DNA J2 haplogroups, present in over a third of the individuals [3,51,62,63].



Figure 1. Early routes of the Romani diaspora. The figure is based on data from Kenrick 2007, Mendizabal et al., 2012, Moorjani et al., 2013, Font-Porterias et al., 2019 and Bianco et al., 2020, and reports the estimated date of arrival and settlement of the Romani populations in different regions of West Eurasia. The wheel symbols indicate the two main reported bottlenecks experienced by the Roma, the arrows indicate the main flow of the migrations and the dotted lines indicate the possible routes of the diaspora.

In general, Romani people carry approximately 65–80% West Eurasian (European, Middle Eastern and Caucasian) ancestry, estimated to have been acquired by extensive gene flow between the 13th and 16th centuries [4,7]. Among the West Eurasian sources, Eastern Europe constitutes the major source of European ancestry [4,6,7,54]. The different estimates of West Eurasian ancestry depend on the inclusion, or exclusion, of the South Asian AWE component in the Roma (derived from the ANI in Indian populations): estimates of the recent West Eurasian component increase from 65% [7] to 80% [4] when AWE is included. In addition, the influence of West Eurasian gene flow in the Roma is attested by the presence in the moderate to high frequencies of West Eurasian mtDNA (i.e., H, U, X, J, T) and Y-DNA lineages (i.e., J and R) [3,19,24,52,64].

4. The Genetic Differentiation and Heterogeneity among Different European Romani Subpopulations

Despite their common origin, Romani groups have experienced several population splits, founder effects, small population sizes and differential gene flow from non-Romani groups that have caused strong drift effects and rapid genetic differentiation and hetero-geneity among different Romani subpopulations. The effect of this genetic heterogeneity is shown in the diverse frequency patterns of uniparental haplogroups [65–68], as well as in the frequency of disease-causing mutations [22–24] and genome-wide structures [4,54,57,69,70]. This has resulted in a marked genetic substructure with different levels of complexity [57,71].

In addition to differential bottlenecks and the loss of genetic variation experienced by each Romani group, part of the heterogeneity might be explained by different exogamy patterns expressed as gene flow within non-Romani groups, which might have been different in sources and amount [7]. As part of the recent European admixture, every Romani exhibits a Balkan common element (with different percentages in each group) and some local gene flow that are specific to each group, i.e., non-Roma Spanish into the Spanish Roma or non-Roma Polish into the Polish Roma [4,7]. The Western European Roma generally differ by exhibiting higher amounts of non-Roma European ancestry and lower levels of South Asian ancestry [4,7,72] (Figure 2).



Figure 2. South Asian and West Eurasian ancestry components in Romani groups based on the results from Font-Porterias et al., 2019. The weighted means for the individuals obtained in the original analysis by genetic clusters are regrouped by country of origin. For sake of clarity, only West Eurasian ancestry percentages are shown.

Some differences between Romani groups can be spotted from the heterogeneous distribution of the mtDNA haplogroups and from the low frequency of M5 in the Polish, Iberian and Lithuanian Roma [3] to the presence of some exclusive branches, such as X2e, in the same Roma populations [73]. As other instances of differentiation, the Polish Roma frequently show haplotypes from W and K haplogroups that appear to be absent in other Romani populations [65], while the Romungro from Ukraine have the highest frequency of the J1 haplogroup and show a complete absence of the M5 haplogroup [6].

Differences in the time of admixture between the Roma and non-Romani groups have also been explored through the use of haplotype-based methods of dating, as these results show that the older dates of admixture for the Balkan Roma genetic clusters (a grouping method based on haplotype affinity) are around 800 ya, whereas the Iberian Roma genetic clusters are estimated to be around 600 ya [7]. The presence of South-Western European elements in the Iberian Roma also supports this hypothesis, as these elements are lacking in the Balkan groups. Nonetheless, Balkan genetic clusters, such as the Croatian Roma, show a higher level of more recent admixture, with an estimated date of admixture around 400 ya [7].

Local ancestry analyses have been used to infer more details about the time of these admixture events by measuring the length of ancestral chromosomal segments. As suggested by Mendizabal et al. [5], recent gene flow coming from non-Romani groups is supposed to generate individuals with long chromosomal segments of non-Roma European ancestry and individuals without traces of these segments, whereas cumulative recombination tends to shorten and spread the non-Roma European chromosomal tracts across the Roma population [74]. This is the case for several Romani groups from Central Europe and the Balkans, where a few individuals have shown very long ancestral chromosomal segments of non-Roma European origin; meanwhile, the populations overall have low mean values of admixture, which is possibly reflective of modern changes in social rules and endogamy practices, in particular [5]. The Roma from Lithuania and the Iberian Peninsula instead show higher values of non-Roma European admixture but in shorter segments, which might be proof of a more recent genetic isolation [5].

5. The Genetic Diversity of the Roma: Current Insights in the Bottleneck and Founder Events

The analysis of the demographic history of the Romani people presents two main levels of complexity. The first level of complexity comes from the gene flow with non-Romani groups that occurred all over their history, as shown in the previous sections. The second level is represented by the series of bottlenecks and splits experienced during their diaspora [5,57,75] and the different amounts of endogamy that have left traces of low intragroup diversity and high intergroup heterogeneity [6,52,63–65,76–78]. In this sense, two major bottlenecks have impacted the history of the Roma: the out-of-India and the out-of-Balkans migrations (Figure 2), events that are discernible through variations in the effective population size (*Ne*), which fluctuated moderately for generations and drastically reduced in these particular instances [5,57].

According to genetic evidence, initially, the *N*e of the Roma overlapped with the *N*e of Northern Indian individuals until ~125 generations ago (ga), in agreement with the Roma origins in this region of the subcontinent [5]. The proto-Roma started to differentiate soon after the out-of-India event, proven by the occurrence of a bottleneck in which the proto-Roma Ne became half of the parental Indian population [5,57]. It is thought that the proto-Roma generated in this circumstance left their place of origin in a single migration event [6,7]. In the period of time that goes from 125 ga to 50 ga, the Roma *N*e became lower than that of Northern Indians, and around 50 ga, it became even lower than the Dravidic groups of southern India [79,80].

The out-of-Balkans bottleneck is estimated to have happened around 900 ya, which coincides with the divergence between the Eastern and Western European Roma; the Western European Roma appear to have undergone an additional bottleneck, which reduced their *N*e to a third of that of the Eastern European Roma [5]. In the following period, both the Western and Eastern European Roma admixed with non-Roma European populations, and in a few generations, extensive gene flow from non-Roma Europeans increased the Roma *N*e, thus, compensating for the previous loss of genetic diversity [5,57]. Historically, the cessation of the *N*e reduction coincides with the settlement of the Roma in Europe and the beginning of more intense assimilation policies during the 17th century [2,57,81]. The increasing genetic distance from the Balkans and the decaying *N*e in the Western Roma suggest that cumulative drift events within Europe are one of the main catalysts of genetic differentiation within the subpopulations [5,82,83]. For this reason, Roma diversity decreased from Eastern towards Western and Northern groups due to the accumulation of drift effects during successive population splitting and migrations along the dispersion

within Europe [5]. The signature of the founder effects and subsequent bottlenecks, amplified by higher levels of isolation and rates of endogamy, is also evident in the presence of a high number and total length of runs of homozygosity (RoH), in comparison to other European and Northern Indian populations [57]. In addition, a lower number of RoH was found in the Roma from the Balkans [57], possibly explicable by the fact that this group did not experience the out-of-Balkans bottleneck.

6. Gene Flow Sex Bias in the Roma Population

The Roma, as many human societies [84], traditionally have patrilocal residence patterns where the sociocultural group affiliation is patrilinearly inherited [85,86].

The genetic evidence for sexual asymmetries in gene flow has been provided by the data coming from studies on maternally inherited mtDNA and paternally inherited Y-DNA, due to their lack of recombination and well-established phylogeography [87]. Besides uniparental lineages, comparing differences in ancestry proportions between the X chromosome and the autosomes also provides information about sex-specific population history as two-thirds of the X chromosomes in a population are carried by women; therefore, the influence of female ancestry is higher than that of male ancestry [88]. In the case of the Roma, sex-biased patterns have been found in their genetic variation. The South Asian ancestral component in the Romani groups has been found in significantly different proportions in male and female lineages [3,6,89]. In this sense, South Asian lineages (mainly represented by the H1a1a4b2 haplogroup) have been found at high frequencies in the Ychromosome genetic pool (approximately 50%), whereas their presence in the maternal counterpart (represented by the M haplogroups) is less frequent (less than 25%). This points to a higher male South Asian gene pool in the Romani groups [7]. Interestingly, comparing X chromosome ancestry proportions to those in the autosomes, South Asian ancestry had a higher result in the X chromosome than in the autosome [7].

Both uniparental South Asian lineages present clear evidence of founder effects [6], although the maternal lineages show higher levels of diversity, and their origin does not seem to be restricted to a single geographic location. In contrast, the Y-DNA lineages are more homogeneous, and their origin seems to be restricted to the region of Pakistan, reflecting a patrilocal pattern in the formation of the proto-Roma population [89].

Differences between males and females can also be observed in the West Eurasian lineages incorporated by the Roma during their diaspora. More than half of the mtDNA lineages in the Roma have a West Eurasian origin, belonging to different haplogroups (i.e., H7, J1b3 and J1c1) [6,21]. Some of these lineages present evidence of founder effects [6], and their origin seems to be explained by the independent assimilation of non-Roma females all along the Romani diaspora [6,21]. In contrast, the West Asian paternal lineage composition is dominated by Middle Eastern lineages with very low diversity (i.e., J2a1b). In a similar way, most European paternal lineages belong to a single haplogroup of Balkan origin (I1a-Z140) [89].

In agreement with the uniparental markers' data, higher Middle Eastern and Caucasian ancestry proportions were observed in the autosomes when compared to the X chromosomes, suggesting a higher influx of non-Roma males in the Middle Eastern and Caucasian regions through the Romani diaspora [7].

The differences between mtDNA and Y-DNA suggest that the inclusion of non-Romani people into Roma communities has been different according to sex: non-Roma females seem to have been adopted more easily, while the low diversity levels in male lineages suggest a sporadic inclusion of non-Roma males due to changes in the socio-historical context [2,90].

7. Biomedical Implications of the Complex Romani Population Demography

Due to the characteristic history of the Romani population, some genetic conditions have been reported to be more frequent than in non-Romani groups. Others have been reported to be less frequent or absent, while new founder mutations have been detected, highlighting a distinct genetic background in terms of disease occurrence [22,23,91,92]. The

possible impact of bottlenecks on the health of the Roma is discernible from the reduced genetic diversity (compared to other Europeans and South Asians), the depletion of rare alleles and the increased high frequency of slightly deleterious genomic variants [69,92]. The high level of homozygosity is another factor that can generate a higher frequency of harmful recessive mutations [91,92]. These demographic factors might have increased the mutational load in the Roma; however, this increase might have been counterbalanced by the extensive gene flow coming from the admixture with other populations. As a consequence, the excess of deleterious mutations is limited, as most did not have time to fixate due to rapid admixture events occurring shortly after founder events [69].

Regarding selection, several pathways related to immunity, metabolism, histone modification, cardiovascular traits, and xenobiotic response have been described to be subject to selective pressures in the Romani groups [69,93]. However, these signals of positive selection are also present in European or South Asian populations, pointing that the signals found in the Roma might be explained by selection processes that occurred before the admixture between these two groups, and that the signal is maintained in the Roma due to drift or weaker positive selection after the admixture [69].

Aside from genetic factors, previous reports suggest that the health status of the Roma is typically characterized by higher mortality, higher morbidity and shorter life expectancy (LE) than the non-Romani populations of Europe [94–96], where the most common causes of mortality are cardiovascular and respiratory diseases [96]. The causes of these are still unidentified and under research; however, they could likely be caused by a combination of inherited metabolic risk factors, socio-cultural and economic factors, discrimination or cultural constraints that prevent Romani people from accessing health care which has consequences, including limited access to preventive medicine [97].

As in other human populations that have experienced founder effects and bottlenecks linked to geographical, linguistic or cultural isolation, such as Finns [98], Sardinians [99] and French Canadians [100], among others, a number of private Romani mutations associated with specific single-gene disorders (Mendelian disorders) have been reported. For instance, among these private conditions, we found hereditary motor and sensory neuropathy Lom type and Russe type, congenital-cataracts facial-dysmorphism neuropathy, congenital myasthenic syndromes, limb-girdle muscular dystrophy type 2C and galactokinase deficiency to be the most common [22,23]. On average, one out of eight Romani people is a carrier of one of the five more common Mendelian mutations [22], with carrier rates for specific mutations often exceeding 5%, and sometimes 15%, depending on the analyzed group [22,23,101]. Many of the private mutations detected in the Roma have been described as ancient mutations present in the population prior to the out-of-India diaspora and are, therefore, shared by Romani patients throughout the European continent, Pakistan and India [22].

Regarding complex diseases, multiple studies have reported a higher prevalence of type 2 diabetes (T2DM) and prediabetes (PreDM) [97,102], metabolic syndrome, cardiovascular disease, a higher occurrence of obesity, overweight and hypertension [96,103], which all may contribute to their higher mortality [104]. All the risk factors were reported to be present more regularly in the Romani population compared with the majority populations in some of the studies, including smoking, sedentary lifestyle, unhealthy dietary habits, low socio-economic status and alcohol consumption, which was also associated with a higher incidence of obesity in many of them [97]. While studies have reported that the cardiovascular risk load in the Romani population differs from the non-Romani population of the same countries, the genetic risk for cardiovascular diseases appears to be in line with other European populations [105], if not lower [106]. Obesity, hypertension and smoking are also associated with metabolic syndrome, insulin resistance, high blood pressure and elevated serum lipids that can lead to the development of metabolic disorders, such as T2DM and cardiovascular disease [96]. Considering the fact that no firm conclusions can be made with the current knowledge, a metabolic hypothesis has been proposed in an effort to identify some of the causes of obesity in the Roma. The hypothesis suggests that the Roma

may have experienced inadequate maternal nutrition throughout their diaspora, which would have led to fetal undernutrition. As a result, genes that would alter glucose–insulin metabolism and help the body store more energy reserves might have been selected. After settling in Europe, they acquired better nutrition and substantially lowered their energy expenditure (due to the end of their nomadic lifestyle), which may have contributed to obesity, T2DM and metabolic syndrome, all of which negatively affect LE [93,107].

For some cardiovascular/metabolic conditions, the Roma show a higher average risk allele frequency (RAF) compared to other Europeans, but the difference in RAF has so far only been found to be statistically significant only for insulin response [92]. A similar but weaker tendency was observed when comparing the Roma to Indian groups [91,92]. Generally, it is necessary to keep in consideration that Asians are genetically more susceptible to insulin resistance and diabetes than Europeans [97], and although some of the existing studies suggest a substantial prevalence of diabetes among Romani populations, and even a higher risk of developing diabetes in Romani individuals compared to non-Roma individuals, the amount of published literature on the topic remains scarce and insufficient to draw any conclusions.

In light of these aspects, it is likely that public interventions that improve general and preventive healthcare access for the Roma would have a large impact on their health conditions and life expectancy.

8. Future Perspectives

Despite the efforts performed in recent times to provide a complete panorama of the genetic diversity of our species and to overcome the lack of genetic data of some human populations, ethnic minorities are still underrepresented in large genomic projects [108,109]. In this sense, one important challenge in studying the population genetics of the Roma is a lack of data, the sampling biases and the underrepresentation of some of the Romani groups analyzed so far. Most population genetic studies have only considered the countries of origin of the Roma individuals, regardless of their socio-cultural and ethnic characteristics. This bias might mask part of the genetic diversity of Romani groups. Genetic data are absent for some of them, such as the scanty data from Central European Sinti groups, not to mention the complete lack of knowledge of the Middle Eastern and American Roma gene pools. In relation to social issues, it is not known to what extent socio-cultural customs are a factor in limiting gene flow and population admixture in the Roma, and sociocultural norms may promote assortative mating [110]. So far, no studies on the assortative mating of the Roma have been undertaken, and this is another matter that needs to be addressed.

Minority communities regularly experience discriminatory practices in healthcare and receive inferior medical care [111]. Therefore, as in other minoritarian populations, in contrast to the one-fits-all model, a right-treatment-to-the-right-patient approach should be sought and broader access to personalized medicine should be built. It is particularly important to evaluate how to assess the frequencies of functional genetic variants in key drug responses and metabolism genes of the Roma, as these significantly influence drug response differences in different populations [112]. To conclude, for the interpretation of ethnic inequalities, further research is required to determine the impact of how genetic, inherited, environmental effects and other environmental factors interact. As ethnicity is not only influenced by genetic ancestry but also by socio-cultural factors, a multidisciplinary approach is needed to unravel the population history of Romani groups.

Future genetic studies including Romani groups should follow the general ethical guidelines required for genetic studies of human populations, including those where minority populations participate, ensuring appropriate informed consent is obtained and data sharing policies, among other ethnic regulations, are followed [113,114]. However, some recent articles have casted a shadow over the entire field by questioning the ethical review and consenting process in studies of the Roma [115]. This patronizing attitude assumes that only members of human minorities are unable to exercise their free will when consenting to participation in a genetic project, which can be considered a form of racism.

As in any other underrepresented group, the engagement of the Roma community in future genomic approaches is of pivotal importance to maximize benefits, minimize potential harms and protect the rights and interests of the participants [116,117].

Author Contributions: Conceptualization, G.F.E. and D.C.; writing—review and editing, G.F.E., J.A.-I., N.F.-P., D.C. and F.C. All authors have read and agreed to the published version of the manuscript.

Funding: This work was supported by the Spanish Ministry of Science, Innovation and Universities (MCIU) and the Agencia Estatal de Investigación (AEI) [PID2019-106485GB-I00/AEI/10.13039/50110 0011033]; and "Unidad de Excelencia María de Maeztu" [AEI, CEX2018-000792-M].

Acknowledgments: We would like to thank the Roma FAGiC association (Federació d'Associacions Gitanes de Catalunya) and all its members and associates for its continuous support and collaboration.

Conflicts of Interest: The authors declare no conflict of interest.

References

- 1. Hancock, I. On Romani Origins and Identity. The Romani Archives and Documentation Center, University of Texas at Austin, USA. 2006. Available online: https://www.oocities.org/romani_life_society/indexpics/HistoryOrigins.pdf (accessed on 1 June 2022).
- 2. Fraser, A.M. The Gypsies (The Peoples of Europe); Blackwell Pub.: Hoboken, NJ, USA, 1992; ISBN 978-0-631-19605-1.
- 3. Gresham, D.; Morar, B.; Underhill, P.A.; Passarino, G.; Lin, A.A.; Wise, C.; Angelicheva, D.; Calafell, F.; Oefner, P.J.; Shen, P.; et al. Origins and Divergence of the Roma (Gypsies). *Am. Soc. Hum. Genet.* **2001**, *69*, 1314–1331. [CrossRef]
- 4. Moorjani, P.; Patterson, N.; Loh, P.-R.; Lipson, M.; Kisfali, P.; Melegh, B.I.; Bonin, M.; Kádaši, L.; Rieß, O.; Berger, B.; et al. Reconstructing Roma History from Genome-Wide Data. *PLoS ONE* **2013**, *8*, e58633. [CrossRef]
- Mendizabal, I.; Lao, O.; Marigorta, U.M.; Wollstein, A.; Gusmão, L.; Ferak, V.; Ioana, M.; Jordanova, A.; Kaneva, R.; Kouvatsi, A.; et al. Report Reconstructing the Population History of European Romani from Genome-Wide Data. *Curr. Biol.* 2012, 22, 2342–2349. [CrossRef]
- Martínez-Cruz, B.; Mendizabal, I.; Harmant, C.; Pablo, R.D.; Ioana, M.; Angelicheva, D.; Kouvatsi, A.; Makukh, H.; Netea, M.G.; Pamjav, H.; et al. Origins, Admixture and Founder Lineages in European Roma. *Eur. J. Hum. Genet.* 2016, 24, 937–943. [CrossRef] [PubMed]
- Font-Porterias, N.; Arauna, L.R.; Poveda, A.; Bianco, E.; Rebato, E.; Prata, M.J.; Calafell, F.; Comas, D. European Roma Groups Show Complex West Eurasian Admixture Footprints and a Common South Asian Genetic Origin. *PLOS Genet.* 2019, 15, e1008417. [CrossRef] [PubMed]
- 8. Kenrick, D. Historical Dictionary of the Gypsies (Romanies); Scarecrow Press: Lanham, MD, USA, 2007; ISBN 978-0-8108-6440-5.
- 9. Covrig, A. Why Roma Do Not Declare Their Identity—Careful Decision or Unpremeditated Refusal? J. Study Relig. Ideol. 2010, 3, 90–101.
- 10. Roma Equality, Inclusion and Participation in the EU. Available online: https://ec.europa.eu/info/policies/justice-and-fundamental-rights/combatting-discrimination/roma-eu/roma-equality-inclusion-and-participation-eu_en (accessed on 8 June 2022).
- 11. Matras, Y. Romani A Linguistic Introduction; UK Cambridge University Press: Cambridge, UK, 2002; ISBN 0-511-03580-2.
- Zatreanu, M.; Halwachs, D.W. Romani in Europe. Ministry of Education and Research–Bucharest Romania, Department of Linguistics at the University of Graz, Austria. 2003. Available online: https://www.coe.int/t/dg4/education/roma/Source/ RomaniEurope_EN.pdf (accessed on 8 June 2022).
- 13. Iovita, R.P.; Schurr, T.G. Reconstructing the origins and migrations of diasporic populations: The case of the European Gypsies. *Am. Anthropol.* **2004**, *106*, 267–281.
- 14. Marushiakova, E.; Popov, V. Gypsies (Roma) in Bulgaria (Volume 18); Peter Lang Pub Incorporated: New York, NY, USA, 1997.
- 15. Grierson, G.A. Doms, Jâts, and the Origin of the Gypsies. J. Gypsy Lore Soc. 1888, 1, 71.
- 16. Frossard, E.B.D. The Gypsies in France. Saturday Mag. 1835, 6, 39–40.
- 17. Crawfurd, J. On the Origin of the Gypsies. Trans. Ethnol. Soc. Lond. 1865, 3, 25–36. [CrossRef]
- 18. Colocci, A. The Gypsies: History of an Errant People; Arnaldo Forni Editore: Bologna, Italy, 1889.
- 19. Pamjav, H.; Zalán, A.; Béres, J.; Nagy, M.; Chang, Y.M. Genetic Structure of the Paternal Lineage of the Roma People. *Am. J. Phys. Anthropol.* **2011**, *145*, 21–29. [CrossRef] [PubMed]
- 20. Regueiro, M.; Rivera, L.; Chennakrishnaiah, S.; Popovic, B.; Andjus, S.; Milasin, J.; Herrera, R.J. Ancestral Modal Y-STR Haplotype Shared among Romani and South Indian Populations. *Gene* **2012**, *504*, 296–302. [CrossRef]
- Mendizabal, I.; Valente, C.; Gusmão, A.; Alves, C.; Gomes, V.; Goios, A.; Parson, W.; Calafell, F.; Alvarez, L.; Amorim, A.; et al. Reconstructing the Indian Origin and Dispersal of the European Roma: A Maternal Genetic Perspective. *PLoS ONE* 2011, 6, e15988. [CrossRef]
- 22. Kalaydjieva, L.; Gresham, D.; Calafell, F. Genetic Studies of the Roma (Gypsies): A Review. BMC Med. Genet. 2001, 2, 5. [CrossRef]

- 23. Morar, B.; Gresham, D.; Angelicheva, D.; Tournev, I.; Gooding, R.; Guergueltcheva, V.; Schmidt, C.; Abicht, A.; Lochmüller, H.; Tordai, A.; et al. Mutation History of the Roma/Gypsies. *Am. J. Hum. Genet.* **2004**, *75*, 596–609. [CrossRef] [PubMed]
- 24. Kalaydjieva, L.; Morar, B.; Chaix, R.; Tang, H. A Newly Discovered Founder Population: The Roma/Gypsies. *BioEssays* 2005, 27, 1084–1094. [CrossRef]
- 25. Bloch, J.; David, H.; Foletier, F.D.V. The Gypsies; FeniXX: Singapore, 1969; ISBN 978-2-7059-1051-8.
- 26. Brearley, M. The Persecution of Gypsies in Europe. Am. Behav. Sci. 2001, 45, 588–599. [CrossRef]
- 27. Taylor, B. Another Darkness, Another Dawn: A History of Gypsies, Roma and Travellers; Reaktion Books: London, UK, 2014; ISBN 978-1-78023-297-3.
- 28. Barany, Z. The East European Gypsies in the Imperial Age. Ethn. Racial Stud. 2001, 24, 50–63. [CrossRef]
- 29. Crowe, D.; Kolsti, J.; Hancock, I. The Gypsies of Eastern Europe; Routledge: London, UK, 2016; ISBN 978-1-315-49024-3.
- 30. Kamusella, T.; Jaskułowski, K. Nationalisms Today; Peter Lang: New York, NY, USA, 2009; ISBN 978-3-03911-883-0.
- Marushiakova, E.; Popov, V.Z.; Popov, V. *Gypsies in the Ottoman Empire: A Contribution to the History of the Balkans*; University of Hertfordshire Press: Hatfield, UK, 2001; Volume 22, ISBN 978-1-902806-02-0.
- 32. Lewy, G. The Nazi Persecution of the Gypsies; Oxford University Press: Oxford, UK, 2000; ISBN 978-0-19-802904-5.
- 33. Ioanid, R. The Holocaust in Romania: The Destruction of Jews and Gypsies Under the Antonescu Regime, 1940–1944; Ivan R. Dee: Chicago, IL, USA, 2008; ISBN 978-1-4616-9490-8.
- 34. Kenrick, D.; Puxon, G. Gypsies Under the Swastika; University of Hertfordshire Press: Hatfield, UK, 2009; ISBN 978-1-902806-80-8.
- 35. Trevisan, P. 'Gypsies' in Fascist Italy: From Expelled Foreigners to Dangerous Italians. Soc. Hist. 2017, 42, 342–364. [CrossRef]
- Kotljarchuk, A. World War II and the Registration of Roma in Sweden: The Role of Experts and Census-Takers. *Holocaust Genocide* Stud. 2017, 31, 457–479. [CrossRef]
- Palmer, M.D.; Burgess, S.M. The Wiley-Blackwell Companion to Religion and Social Justice; John Wiley & Sons: Hoboken, NJ, USA, 2020; ISBN 978-1-119-57210-7.
- A Persisting Concern: Anti-Gypsyism as a Barrier to Roma Inclusion. Available online: https://fra.europa.eu/en/publication/20 18/persisting-concern-anti-gypsyism-barrier-roma-inclusion (accessed on 5 September 2022).
- Council of Europe Strategic Action Plan for Roma and Traveller Inclusion (2020–2025). Available online: https://edoc.coe.int/en/ roma-and-travellers/8508-council-of-europe-strategic-action-plan-for-roma-and-traveller-inclusion-2020-2025.html (accessed on 14 September 2022).
- ECRI General Policy Recommendation No. 13. Available online: https://www.coe.int/en/web/european-commission-againstracism-and-intolerance/recommendation-no.13 (accessed on 5 September 2022).
- Bernasovský, I.; Suchý, J.; Bernasovská, K.; Vargová, T. Blood Groups of Roms (Gypsies) in Czechoslovakia. Am. J. Phys. Anthropol. 1976, 45, 277–279. [CrossRef] [PubMed]
- 42. Bernasovský, I.; Jurícková, J.; Ferák, V. Population Genetic Study in Gypsies (Roms) from Slovakia: Distribution of Blood Group Genetic Markers. *Anthropol. Sci.* 1994, 102, 409–419. [CrossRef]
- Gáliková, J.; Vilímová, M.; Ferák, V.; Mayerová, A. Haptoglobin Types in Gypsies from Slovakia (Czechoslovakia). *Hum. Hered.* 1969, 19, 480–485. [CrossRef]
- Piccolo, F.; Jeanpierre, M.; Leturcq, F.; Dodé, C.; Azibi, K.; Toutain, A.; Merlini, L.; Jarre, L.; Navarro, C.; Krishnamoorthy, R.; et al. A Founder Mutation in the γ-Sarcoglycan Gene of Gypsies Possibly Predating Their Migration Out of India. *Hum. Mol. Genet.* 1996, 5, 2019–2022. [CrossRef] [PubMed]
- Lasa, A.; Piccolo, F.; de Diego, C.; Jeanpierre, M.; Colomer, J.; Rodríguez, M.J.; Urtizberea, J.A.; Baiget, M.; Kaplan, J.C.; Gallano, P. Severe Limb Girdle Muscular Dystrophy in Spanish Gypsies: Further Evidence for a Founder Mutation in the γ-Sarcoglycan Gene. *Eur. J. Hum. Genet.* **1998**, *6*, 396–399. [CrossRef]
- Abicht, A.; Stucka, R.; Karcagi, V.; Herczegfalvi, A.; Horváth, R.; Mortier, W.; Schara, U.; Ramaekers, V.; Jost, W.; Brunner, J.; et al. A Common Mutation (E1267delG) in Congenital Myasthenic Patients of Gypsy Ethnic Origin. *Neurology* 1999, 53, 1564. [CrossRef] [PubMed]
- Álvarez, A.; del Castillo, I.; Villamar, M.; Aguirre, L.A.; González-Neira, A.; López-Nevot, A.; Moreno-Pelayo, M.A.; Moreno, F. High Prevalence of the W₂₄X Mutation in the Gene Encoding Connexin-26 (GJB₂) in Spanish Romani (Gypsies) with Autosomal Recessive Non-Syndromic Hearing Loss. *Am. J. Med. Genet.* 2005, 137A, 255–258. [CrossRef]
- Bouwer, S.; Angelicheva, D.; Chandler, D.; Seeman, P.; Tournev, I.; Kalaydjieva, L. Carrier Rates of the Ancestral Indian W₂₄X Mutation in GJB₂ in the General Gypsy Population and Individual Subisolates. *Genet. Test.* 2007, 11, 455–458. [CrossRef]
- Azmanov, D.N.; Dimitrova, S.; Florez, L.; Cherninkova, S.; Draganov, D.; Morar, B.; Saat, R.; Juan, M.; Arostegui, J.I.; Ganguly, S.; et al. LTBP₂ and CYP₁B₁ Mutations and Associated Ocular Phenotypes in the Roma/Gypsy Founder Population. *Eur. J. Hum. Genet.* 2011, 19, 326–333. [CrossRef]
- Maheshwari, M.; Vijaya, R.; Ghosh, M.; Shastri, S.; Kabra, M.; Menon, P.S.N. Screening of Families with Autosomal Recessive Non-Syndromic Hearing Impairment (ARNSHI) for Mutations in GJB₂ Gene: Indian Scenario. *Am. J. Med. Genet.* 2003, 120A, 180–184. [CrossRef]
- 51. Gómez-Carballa, A.; Pardo-seco, J.; Fachal, L.; Vega, A.; Cebey, M.; Salas, A. Indian Signatures in the Westernmost Edge of the European Romani Diaspora: New Insight from Mitogenomes. *PLoS ONE* **2013**, *8*, e75397. [CrossRef]
- 52. Zalán, A.; Béres, J.; Pamjav, H. Paternal Genetic History of the Vlax Roma. *Forensic Sci. Int. Genet.* 2011, *5*, 109–113. [CrossRef] [PubMed]

- Rai, N.; Chaubey, G.; Tamang, R.; Pathak, A.K.; Singh, V.K.; Karmin, M.; Singh, M.; Rani, D.S.; Anugula, S.; Yadav, B.K.; et al. The Phylogeography of Y-Chromosome Haplogroup H1a1a-M82 Reveals the Likely Indian Origin of the European Romani Populations. *PLoS ONE* 2012, 7, e48477. [CrossRef] [PubMed]
- 54. Melegh, B.I.; Banfai, Z.; Hadzsiev, K.; Miseta, A.; Melegh, B. Refining the South Asian Origin of the Romani People. *BMC Genet*. **2017**, *18*, 82. [CrossRef]
- 55. Reich, D.; Thangaraj, K.; Patterson, N.; Price, A.L.; Singh, L. Reconstructing Indian Population History. *Nature* 2009, 461, 489–494. [CrossRef]
- 56. Narasimhan, V.M.; Patterson, N.; Moorjani, P.; Rohland, N.; Bernardos, R.; Mallick, S.; Lazaridis, I.; Nakatsuka, N.; Olalde, I.; Lipson, M. The Formation of Human Populations in South and Central Asia. *Science* **2019**, *365*, eaat7487. [CrossRef]
- 57. Bianco, E.; Laval, G.; Font-Porterias, N.; García-Fernández, C.; Dobon, B.; Sabido-Vera, R.; Stefanovska, E.S.; Kučinskas, V.; Makukh, H.; Pamjav, H.; et al. Recent Common Origin, Reduced Population Size, and Marked Admixture Have Shaped European Roma Genomes. *Mol. Biol. Evol.* 2020, *37*, 3175–3187. [CrossRef]
- 58. Matras, Y.; Bakker, P.; Kyuchukov, H. *The Typology and Dialectology of Romani*; John Benjamins B.V.: Amsterdam, The Netherlands, 1997.
- Bánfai, Z.; Ádám, V.; Pöstyéni, E.; Büki, G.; Czakó, M.; Miseta, A.; Melegh, B. Revealing the Impact of the Caucasus Region on the Genetic Legacy of Romani People from Genome-Wide Data. *PLoS ONE* 2018, 13, e0202890. [CrossRef]
- 60. Derenko, M.; Malyarchuk, B.; Bahmanimehr, A.; Denisova, G.; Perkova, M.; Farjadian, S.; Yepiskoposyan, L. Complete Mitochondrial DNA Diversity in Iranians. *PLoS ONE* **2013**, *8*, e80673. [CrossRef]
- Tarkhnishvili, D.; Gavashelishvili, A.; Murtskhvaladze, M.; Gabelaia, M.; Tevzadze, G. Human Paternal Lineages, Languages, and Environment in the Caucasus. *Hum. Biol.* 2014, 86, 113–130. [CrossRef]
- 62. Malyarchuk, B.A.; Perkova, M.A.; Derenko, M.V.; Vanecek, T.; Lazur, J.; Gomolcak, P. Mitochondrial DNA Variability in Slovaks, with Application to the Roma Origin. *Ann. Hum. Genet.* **2008**, *72*, 228–240. [CrossRef] [PubMed]
- 63. Gusmão, A.; Gusmão, L.; Gomes, V.; Alves, C.; Calafell, F.; Amorim, A.; Prata, M.J. A Perspective on the History of the Iberian Gypsies Provided by Phylogeographic Analysis of Y-Chromosome Lineages. *Ann. Hum. Genet.* **2008**, *72*, 215–227. [CrossRef] [PubMed]
- Klarić, I.M.; Salihović, M.P.; Lauc, L.B.; Zhivotovsky, L.A.; Rootsi, S.; Janićijević, B. Dissecting the Molecular Architecture and Origin of Bayash Romani Patrilineages: Genetic Influences from South-Asia and the Balkans. *Am. J. Phys. Anthropol.* 2009, 138, 333–342. [CrossRef] [PubMed]
- 65. Malyarchuk, B.A.; Grzybowski, T.; Derenko, M.V.; Czarny, J.; Miścicka-Śliwka, D. Mitochondrial DNA Diversity in the Polish Roma. Ann. Hum. Genet. 2006, 70, 195–206. [CrossRef]
- Kalaydjieva, L.; Calafell, F.; Jobling, M.A.; Angelicheva, D.; Knijff, P.D.; Rosser, Z.H.; Hurles, M.E.; Underhill, P.; Tournev, I.; Marushiakova, E.; et al. Patterns of Inter- and Intra-Group Genetic Diversity in the Vlax Roma as Revealed by Y Chromosome and Mitochondrial DNA Lineages. *Eur. J. Hum. Genet.* 2001, *9*, 97–104. [CrossRef]
- Pokupčić, K.; Cukrov, S.; Klarić, I.M.; Salihović, M.P.; Lauc, L.B.; Blažanović, A.; Janićijević, B. Y-STR Genetic Diversity of Croatian (Bayash) Roma. *Forensic Sci. Int. Genet.* 2008, 2, e11–e13. [CrossRef]
- 68. Ahmic, A.; Kalamujic, B.; Ismailovic, A.; Lasic, L.; Halilovic, E.; Mujkic, I.; Pojskic, N. Mitochondrial DNA Diversity of the Roma from Northeastern Bosnia, Bosnia and Herzegovina. *HOMO* **2018**, *69*, 347–356. [CrossRef]
- Font-Porterias, N.; Caro-Consuegra, R.; Lucas-Sánchez, M.; Lopez, M.; Giménez, A.; Carballo-Mesa, A.; Bosch, E.; Calafell, F.; Quintana-Murci, L.; Comas, D. The Counteracting Effects of Demography on Functional Genomic Variation: The Roma Paradigm. *Mol. Biol. Evol.* 2021, 38, 2804–2817. [CrossRef]
- Bánfai, Z.; Melegh, B.I.; Sümegi, K.; Hadzsiev, K.; Miseta, A.; Kásler, M.; Melegh, B. Revealing the Genetic Impact of the Ottoman Occupation on Ethnic Groups of East-Central Europe and on the Roma Population of the Area. *Front. Genet.* 2019, 10, 558. [CrossRef]
- Morar, B.; Azmanov, D.N.; Kalaydjieva, L. Roma (Gypsies): Genetic Studies. In *eLS*; John Wiley & Sons: Hoboken, NJ, USA, 2013; ISBN 978-0-470-01590-2.
- 72. Pereira, V.; Gusmão, L.; Valente, C.; Pereira, R.; Carneiro, J.; Gomes, I.; Morling, N.; Amorim, A.; João Prata, M. Refining the Genetic Portrait of Portuguese Roma through X-Chromosomal Markers. *Am. J. Phys. Anthropol.* **2012**, *148*, 389–394. [CrossRef]
- 73. Reidla, M.; Kivisild, T.; Metspalu, E.; Kaldma, K.; Tambets, K.; Tolk, H.-V.; Parik, J.; Loogväli, E.-L.; Derenko, M.; Malyarchuk, B.; et al. Origin and Diffusion of MtDNA Haplogroup X. *Am. J. Hum. Genet.* **2003**, *73*, 1178–1190. [CrossRef] [PubMed]
- 74. Chapman, N.H.; Thompson, E.A. The Effect of Population History on the Lengths of Ancestral Chromosome Segments. *Genetics* **2002**, *162*, 449–458. [CrossRef]
- 75. Chaix, R.; Austerlitz, F.; Morar, B.; Kalaydjieva, L.; Heyer, E. Vlax Roma History: What Do Coalescent-Based Methods Tell Us? *Eur. J. Hum. Genet.* **2004**, *12*, 285–292. [CrossRef]
- 76. Irwin, J.; Egyed, B.; Saunier, J.; Szamosi, G.; O'Callaghan, J.; Padar, Z.; Parsons, T.J. Hungarian MtDNA Population Databases from Budapest and the Baranya County Roma. *Int. J. Legal Med.* **2007**, *121*, 377–383. [CrossRef]
- Salihović, M.P.; Barešić, A.; Klarić, I.M.; Cukrov, S.; Lauc, L.B.; Janićijević, B. The Role of the Vlax Roma in Shaping the European Romani Maternal Genetic History. *Am. J. Phys. Anthropol.* 2011, 146, 262–270. [CrossRef] [PubMed]
- 78. Peričić, M.; Klarić, I.M.; Lauc, L.B.; Janićijević, B.; Đorđević, D.; Efremovska, L.; Rudan, P. Population Genetics of 8 Y Chromosome STR Loci in Macedonians and Macedonian Romani (Gypsy). *Forensic Sci. Int.* **2005**, *154*, 257–261. [CrossRef] [PubMed]

- Juyal, G.; Mondal, M.; Luisi, P.; Laayouni, H.; Sood, A.; Midha, V.; Heutink, P.; Bertranpetit, J.; Thelma, B.K.; Casals, F. Population and Genomic Lessons from Genetic Analysis of Two Indian Populations. *Hum. Genet.* 2014, 133, 1273–1287. [CrossRef]
- Mondal, M.; Bergström, A.; Xue, Y.; Calafell, F.; Laayouni, H.; Casals, F.; Majumder, P.P.; Tyler-Smith, C.; Bertranpetit, J. Y-Chromosomal Sequences of Diverse Indian Populations and the Ancestry of the Andamanese. *Hum. Genet.* 2017, 136, 499–510. [CrossRef]
- 81. Maria Theresia and Joseph II Policies of Assimilation in the Age of Enlightened Absolutism [Rombase]. Available online: http://rombase.uni-graz.at/cgi-bin/art.cgi?src=data/hist/modern/maria.en.xml (accessed on 9 June 2022).
- Gusmão, A.; Valente, C.; Gomes, V.; Alves, C.; Amorim, A.; Prata, M.J.; Gusmão, L. A Genetic Historical Sketch of European Gypsies: The Perspective from Autosomal Markers. *Am. J. Phys. Anthropol.* 2010, 141, 507–514. [CrossRef]
- Inotai, D.; Szilvasi, A.; Benko, S.; Boros-Major, A.; Illes, Z.; Bors, A.; Kiss, K.P.; Rajczy, K.; Gelle-Hossó, A.; Buhler, S.; et al. HLA Genetic Diversity in Hungarians and Hungarian Gypsies: Complementary Differentiation Patterns and Demographic Signals Revealed by HLA-A, -B and -DRB1 in Central Europe. *Tissue Antigens* 2015, *86*, 115–121. [CrossRef]
- Burton, M.L.; Moore, C.C.; Whiting, J.W.M.; Romney, A.K.; Aberle, D.F.; Barcelo, J.A.; Dow, M.M.; Guyer, J.I.; Kronenfeld, D.B.; Levy, J.E.; et al. Regions Based on Social Structure. *Curr. Anthropol.* **1996**, *37*, 87–123.
- 85. Gamella, J.F. Marriage, Gender and Transnational Migrations in Fertility Transitions of Romanian Roma Women. *Intersect. East Eur. J. Soc. Polit.* **2018**, *4*, 57–85. [CrossRef]
- 86. Weyrauch, W.O. Gypsy Law: Romani Legal Traditions and Culture; University of California Press: Berkeley, CA, USA, 2001.
- 87. Heyer, E.; Chaix, R.; Pavard, S.; Austerlitz, F. Sex-Specific Demographic Behaviours That Shape Human Genomic Variation. *Mol. Ecol.* **2012**, *21*, 597–612. [CrossRef] [PubMed]
- Goldberg, A.; Rosenberg, N.A. Beyond 2/3 and 1/3: The Complex Signatures of Sex-Biased Admixture on the X Chromosome. *Genetics* 2015, 201, 263. [CrossRef]
- García-Fernández, C.; Font-Porterias, N.; Kučinskas, V.; Sukarova-Stefanovska, E.; Pamjav, H.; Makukh, H.; Dobon, B.; Bertranpetit, J.; Netea, M.G.; Calafell, F.; et al. Sex-Biased Patterns Shaped the Genetic History of Roma. *Sci. Rep.* 2020, 10, 14464. [CrossRef]
- 90. Hancock, I.F. We Are the Romani People; University of Hertfordshire Press: Hatfield, UK, 2002; ISBN 978-1-902806-19-8.
- 91. Mendizabal, I.; Lao, O.; Marigorta, U.M.; Kayser, M.; Comas, D. Implications of Population History of European Romani on Genetic Susceptibility to Disease. *Hum. Hered.* **2013**, *76*, 194–200. [CrossRef]
- Font-Porterias, N.; Giménez, A.; Carballo-Mesa, A.; Calafell, F.; Comas, D. Admixture Has Shaped Romani Genetic Diversity in Clinically Relevant Variants. Front. Genet. 2021, 12, 683880. [CrossRef]
- 93. Dobon, B.; Ter Horst, R.; Laayouni, H.; Mondal, M.; Bianco, E.; Comas, D.; Ioana, M.; Bosch, E.; Bertranpetit, J.; Netea, M.G. The shaping of immunological responses through natural selection after the Roma Diaspora. *Sci. Rep.* **2020**, *10*, 1–12. [CrossRef]
- 94. Simko, V.; Ginter, E. Short Life Expectancy and Metabolic Syndrome in Romanies (Gypsies) in Slovakia. *Cent. Eur. J. Public Health* **2010**, *18*, 16. [CrossRef]
- 95. Parekh, N.; Rose, T. Health Inequalities of the Roma in Europe: A Literature Review. Cent. Eur. J. Public Health 2011, 19, 139–142.
- 96. Dobranici, M.; Buzea, A.; Popescu, R. The Cardiovascular Risk Factors of the Roma (Gypsies) People in Central-Eastern Europe: A Review of the Published Literature. *J. Med. Life* **2012**, *5*, 382–389. [PubMed]
- Nunes, M.A.; Kučerová, K.; Lukáč, O.; Kvapil, M.; Brož, J. Prevalence of Diabetes Mellitus among Roma Populations—A Systematic Review. Int. J. Environ. Res. Public. Health 2018, 15, 2607. [CrossRef] [PubMed]
- Sarantaus, L.; Huusko, P.; Eerola, H.; Launonen, V.; Vehmanen, P.; Rapakko, K.; Gillanders, E.; Syrjäkoski, K.; Kainu, T.; Vahteristo, P.; et al. Multiple Founder Effects and Geographical Clustering of BRCA1 and BRCA2 Families in Finland. *Eur. J. Hum. Genet.* 2000, *8*, 757–763. [CrossRef] [PubMed]
- Loudianos, G.; Dessi, V.; Lovicu, M.; Angius, A.; Figus, A.; Lilliu, F.; de Virgiliis, S.; Nurchi, A.M.; Deplano, A.; Moi, P.; et al. Molecular Characterization of Wilson Disease in the Sardinian Population—Evidence of a Founder Effect. *Hum. Mutat.* 1999, 14, 294–303. [PubMed]
- 100. Gosselin, I.; Thiffault, I.; Tétreault, M.; Chau, V.; Dicaire, M.-J.; Loisel, L.; Emond, M.; Senderek, J.; Mathieu, J.; Dupré, N.; et al. Founder SH3TC2 Mutations Are Responsible for a CMT4C French-Canadians Cluster. *Neuromuscul. Disord.* 2008, 18, 483–492. [CrossRef] [PubMed]
- Martínez-Frías, M.L.; Bermejo, E. Prevalence of Congenital Anomaly Syndromes in a Spanish Gypsy Population. J. Med. Genet. 1992, 29, 483–486.
- Werissa, N.A.; Piko, P.; Fiatal, S.; Kosa, Z.; Sandor, J.; Adany, R. SNP-Based Genetic Risk Score Modeling Suggests No Increased Genetic Susceptibility of the Roma Population to Type 2 Diabetes Mellitus. *Genes* 2019, 10, E942. [CrossRef]
- 103. Zeljko, H.; Škarić-Jurić, T.; Smolej Narančić, N.; Peričić Salihović, M.; Martinović Klarić, I.; Barbalić, M.; Starčević, B.; Barać Lauc, L.; Janićijević, B. Traditional CVD Risk Factors and Socio-Economic Deprivation in Roma Minority Population of Croatia. *Coll. Antropol.* 2008, 32, 667–676.
- 104. De Courten, B.V.; de Courten, M.; Hanson, R.L.; Zahorakova, A.; Egyenes, H.P.; Tataranni, P.A.; Bennett, P.H.; Vozar, J. Higher Prevalence of Type 2 Diabetes, Metabolic Syndrome and Cardiovascular Diseases in Gypsies than in Non-Gypsies in Slovakia. *Diabetes Res. Clin. Pract.* 2003, 62, 95–103.
- 105. Škarić-Jurić, T.; Zeljko, H.M.; Tomas, Ž.; Peričić Salihović, M.; Smolej Narančić, N.; Janićijević, B. Genetic Risk Factors for Cardiovascular Diseases in the Roma Minority Population of Croatia. *Eur. J. Hum. Genet.* **2009**, *17* (Suppl. S2), 262.

- 106. Soltesz, B.; Piko, P.; Sandor, J.; Kosa, Z.; Adany, R.; Fiatal, S. The Genetic Risk for Hypertension Is Lower among the Hungarian Roma Population Compared to the General Population. *PLoS ONE* **2020**, *15*, e0234547. [CrossRef]
- 107. Ginter, E.; Simko, V.; Wsolova, L. Fall of the Iron Curtain: Male Life Expectancy in Slovakia, in the Czech Republic and in Europe. *Cent. Eur. J. Public Health* **2009**, *17*, 171. [PubMed]
- 108. Popejoy, A.B.; Fullerton, S.M. Genomics Is Failing on Diversity. Nature 2016, 538, 161–164. [PubMed]
- 109. Sirugo, G.; Williams, S.M.; Tishkoff, S.A. The Missing Diversity in Human Genetic Studies. Cell 2019, 177, 26–31. [CrossRef] [PubMed]
- Alfonso-Sánchez, M.A.; Espinosa, I.; Gómez-Pérez, L.; Poveda, A.; Rebato, E.; Peña, J.A. Tau haplotypes support the Asian ancestry of the Roma population settled in the Basque Country. *Heredity* 2018, 120, 91–99. [PubMed]
- 111. Patel, K.; Patel, M.; Mukhi, H. Despite Advances in Precision Medicine, Disparities Persist. Target. Ther. Oncol. 2020, 9, 16–112.
- 112. Ingelman-Sundberg, M.; Mkrtchian, S.; Zhou, Y.; Lauschke, V.M. Integrating Rare Genetic Variants into Pharmacogenetic Drug Response Predictions. *Hum. Genomics* 2018, 12, 26. [CrossRef]
- 113. Knoppers, B.M. Framework for Responsible Sharing of Genomic and Health-Related Data. HUGO J. 2014, 8, 1–6. [CrossRef]
- 114. Claw, K.G.; Anderson, M.Z.; Begay, R.L.; Tsosie, K.S.; Fox, K.; Garrison, N. A Framework for Enhancing Ethical Genomic Research with Indigenous Communities. *Nat. Commun.* **2018**, *9*, 1–7. [CrossRef]
- 115. Lipphardt, V.; Rappold, G.A.; Surdu, M. Representing Vulnerable Populations in Genetic Studies: The Case of the Roma. *Sci. Context* **2021**, *34*, 69–100. [CrossRef]
- 116. Hudson, M.; Garrison, N.; Sterling, R.; Caron, N.R.; Fox, K.; Yracheta, J.; Anderson, J.; Wilcox, P.; Arbour, L.; Brown, A. Rights, Interests and Expectations: Indigenous Perspectives on Unrestricted Access to Genomic Data. *Nat. Rev. Genet.* 2020, 21, 377–384. [CrossRef] [PubMed]
- 117. Lemke, A.A.; Esplin, E.D.; Goldenberg, A.J.; Gonzaga-Jauregui, C.; Hanchard, N.A.; Harris-Wai, J.; Ideozu, J.E.; Isasi, R.; Landstrom, A.P.; Prince, A.E. Addressing Underrepresentation in Genomics Research through Community Engagement. *Am. J. Hum. Genet.* 2022, 109, 1563–1571. [CrossRef] [PubMed]