A new approach to estimating the prevalence of hereditary hearing loss: An analysis of the distribution of sign language users based on census data in Russia

Georgii P. Romanov¹,² Vera G. Pshennikova¹,², Sergey A. Lashin³,⁴, Aisen V. Solovyev¹,⁵, Fedor M. Teryutin¹,², Aleksandra M. Cherdonova¹, Tuyara V. Borisova¹, Nikolay N. Sazonov¹, Elza K. Khusnutdinova⁶,⁷, Olga L. Posukh³,⁴, Sardana A. Fedorova¹,², Nikolay A. Barashkov¹,²*¹

¹ Laboratory of Molecular Biology, MK Ammosov North-Eastern Federal University, Yakutsk, Russia, ² Laboratory of Molecular Genetics, Yakut Science Centre of Complex Medical Problems, Yakutsk, Russia, ³ Federal Research Center Institute of Cytology and Genetics, Siberian Branch of the Russian Academy of Sciences, Novosibirsk, Russia, ⁴ Novosibirsk State University, Novosibirsk, Russia, ⁵ Laboratory of the Human in the Arctic, Institute for Humanitarian Research and North Indigenous Peoples Problems, Federal Research Centre “The Yakut Scientific Centre of the Siberian Branch of the Russian Academy of Sciences”, Yakutsk, Russia, ⁶ Laboratory of Human Molecular Genetics, Institute of Biochemistry and Genetics, Ufa Federal Research Centre of the Russian Academy of Sciences, Ufa, Russia, ⁷ Department of Genetics and Fundamental Medicine, Bashkir State University, Ufa, Russia

Abstract

The absence of comparable epidemiological data challenges the correct estimation of the prevalence of congenital hearing loss (HL) around the world. Sign language (SL) is known as the main type of communication of deaf people. We suggest that the distribution of SL can be interpreted as an indirect indicator of the prevalence of congenital HL. Since a significant part of congenital HL is due to genetic causes, an assessment of the distribution of SL users can reveal regions with an extensive accumulation of hereditary HL. For the first time, we analyzed the data on the distribution of SL users that became available for the total population of Russia by the 2010 census. Seventy-three out of 85 federal regions of Russia were ranked into three groups by the 25th and 75th percentiles of the proportion of SL users: 14 regions—“low proportion”; 48 regions—“average proportion”; and 11 regions—“high proportion”. We consider that the observed uneven prevalence of SL users can reflect underlying hereditary forms of congenital HL accumulated in certain populations by specific genetic background and population structure. At least, the data from this study indicate that the highest proportions of SL users detected in some Siberian regions are consistent with the reported accumulation of specific hereditary HL forms in indigenous Yakut, Tuvinian and Altaian populations.
Introduction

Hearing loss (HL) is one of the most common sensory disorders that makes it a serious public health problem. According to the World Health Organization (WHO), the number of people with disabling HL is approximately 466 million worldwide, including 34 million children [1]. The WHO data are based on the Global Burden of Diseases (GBD) project, which has estimated the HL prevalence based on only 42 studies in 29 countries [2]. Among many studies on HL prevalence in the general population, only a small number of surveys are suitable for analysis, and more population-based surveys are needed in all regions of the world [2, 3]. For countries with missing or scarce data, the GBD calculates the HL prevalence using a Bayesian hierarchical model that is effective for sparse data [2]. Thus, uniform description of HL prevalence is now available for only particular populations or regions since different studies are based on highly specific data and resources.

Universal newborn hearing screening (UNHS) programs are potentially the best source of data on the prevalence of congenital deafness [4–6]. The rate of congenitally deaf infants is generally considered to vary from 1 to 3 per 1,000 newborns in developed countries and up to 24 per 1,000 newborns in developing countries due to the higher presence of risk factors [4–9]. Despite all benefits of the UNHS data, a very limited number of countries (mostly highly developed countries) have massively adopted such programs, and the reported rates of congenital deafness are not applicable for other parts of the world [4, 7, 9]. The lack of systematic reporting and analysis of data also greatly affects the correct evaluation of congenital HL prevalence by the UNHS [4, 7, 9].

At the same time, there are studies assessing HL prevalence by other approaches in particular countries. One of the most well known is the 1972 National Census of the Deaf Population (NCDP) in the USA, which gathered information on the size, geographic distribution, and characteristics of the deaf population in the USA [10]. However, it is difficult to assess whether the proportion of deaf people has been the same since 1972 because the NCDP was conducted only once. Currently, approximately 5–6% of the whole population of the USA is estimated to be deaf or hard of hearing [11, 12]. These data come from the national household surveys conducted by the federal government: the Behavioral Risk Factor Surveillance System (BRFSS), the National Health Interview Survey (NHIS), and the Survey of Income and Program Participation (SIPP). Despite certain general limitations in methodology, such as difficulties in sampling a rare population, the use of a telephone-mediated survey (BRFSS), the exclusion of a certain age or social group (SIPP), and unclear distinction between deaf and hard-of-hearing individuals (NHIS), these surveys are currently the only basis of estimation of HL prevalence in the USA [11, 12].

In Italy, the data from the Italian National Institute of Social Insurance (INPS) were applied as a basis for estimating the prevalence of HL [13]. As, in Italy, sickness benefits are granted to recognized cases, the INPS keeps a record of all cases of prelingual deafness (60 dB or more for 0.5-, 1.0- and 2.0-kHz frequency tones in the better ear). The analysis of available data from 18 of 20 regions of Italy (98.2% of Italian population) detected a total of 40,887 cases of prelingual profound sensorineural HL in Italy in 2003 (the prevalence rate of 0.72 per 1,000) with differences by sex and high accumulation in the southern regions, possibly due to the epidemic incidence of maternal rubella in the 40s and the 50s and due to frequent consanguineous marriages in the past [13]. Unfortunately, there are no other studies suitable for comparison with these data.

In this work, we propose to consider the data on the prevalence of the knowledge of sign language (SL) as an alternative indicator of the prevalence of congenital and prelingual HL. Congenital or early-onset HL is the most socially significant form of HL, greatly affecting the healthy development and socialization of deaf people, most of whom use SL as their only way of communication. Therefore, a high incidence of SL users in certain locations might indicate...
the accumulation of hereditary forms of deafness, as half of the congenital HL cases have a genetic etiology [5]. For the first time, the data on SL users have become available as the result of the 2010 total population census in Russia.

The prevalence of congenital HL in Russia was described previously only for certain populations either by broad surveys on inherited diseases [14–16] or by molecular genetic studies performed on specific samples of deaf patients [17–33]. In this study, we present the estimation of the prevalence of hereditary severe or profound HL in Russia based on the analysis of the 2010 census data concerning the knowledge of sign language.

Materials and methods
To analyze the prevalence of SL users in Russia, we used the data of the last (2010) comprehensive nationwide population census conducted simultaneously according to a unified statistical methodology in all regions of Russia and published by the Federal State Statistics Service (FSSS) of the Russian Federation [34]. The reports were published in thematic volumes: “Size and distribution of the population”, “Age, sex and marital status”, “Education”, “Ethnic composition and language skills, citizenship”, “Sources of livelihood”, “Number and composition of households”, “Economically active and economically inactive population”, “Length of residence of the population in the place of permanent residence”, “Housing conditions of the population”, “Fertility”, which were published as accordingly named volumes of report. The published results of the census are available separately for each region of Russia by local offices of FSSS which were aggregated in total comprehensive statistical report.

In 2010, the Russian Federation constitutionally consisted of 85 federal subjects, and the total population amounted to 142,946,788 people [34]. A distinctive feature of the 2010 census questionnaires that made this study possible was the inclusion of SL on the issue of the knowledge and use of languages. Data on the number of people reporting SL knowledge were extracted from the report tables available on the FSSS website [34]. The data in these tables were formed on the basis of the answers to the questions in the census form: “9.1 Do you speak Russian?”, where the respondents choose “yes” or “no” by putting a mark in the corresponding checkbox, and “9.2 What other languages do you speak?”, where respondents could indicate up to three languages other than Russian, and indicate knowledge of SL by putting a mark in a “sign language” checkbox. Data provided by regional departments of the FSSS were available in full for 73 out of 85 federal subjects of Russia and were analyzed to study the regional distribution of SL users. For each value, we calculated the 95% confidence interval (CI) using the BETAINV function in the Microsoft Excel table made by Mait Metspalu (Estonian Biocentre, Estonia). To categorize the data, we calculated the median, 25th and 75th percentiles for available values (the number and proportion of SL users). We divided regional values for SL users into three groups by lower and upper quartiles. Values significantly lower than the 25th percentile were considered “low”, values significantly higher than the 25th percentile and lower than the 75th percentile were considered “average”, and values significantly higher than the 75th percentile were considered “high”.

This study was approved by the local Biomedical Ethics Committee of Federal State Budgetary Scientific Institution “Yakut Science Centre of Complex Medical Problems”, Yakutsk, Russia (Protocol No. 16, April 16, 2015). The data were analyzed anonymously, and no informed consent forms were required.

Limitations
It should be noted that the number of SL users cannot be directly interpreted as the number of hard-of-hearing people since a certain part of hearing people also know and use SL (relatives,
social workers, interpreters, teachers). Nevertheless, we suggest that the proportion of hearing SL users can be neglected when analyzing the prevalence of SL across regions of Russia due to a relatively uniform distribution of educational institutes and accessibility of social services for deaf people in federal subjects of Russia.

Results

According to the final report of the 2010 census, 138,312,535 people answered the question about their language knowledge. Among them, 18,591,655 respondents indicated that they know and use another language besides Russian (Fig 1). In total, there are 172 languages spoken in Russia, although some of them are dialects, but, as having been specified, still counted as a language [34]. Fifty-three of them are considered foreign, such as English, French, German, Spanish, Japanese, Chinese, and other foreign languages. By excluding these languages, we counted 119 languages that respondents indicated as their native language, and SL, according to the number of SL users, ranks thirty-second in this list (Fig 1 and S1 Table).

To study the prevalence of SL users, we had two sets of relevant data: aggregated statistics and regional reports of local offices of the FSSS. According to the aggregated statistics from the census results, there were 120,528 SL users among 138,312,535 respondents, with a proportion of 0.087%. Information on SL users reported by local FSSS offices was available for 73 out of 85 federal subjects of Russia, and in this case, the total number of SL users was 107,064 out of 122,527,891 respondents, which corresponds to the same proportion of 0.087%. We assume that the “missing” 13,464 SL users are uniformly distributed among 12 regions with unavailable census data.

We analyzed the regional distribution of the number and proportion of SL users across different regions of Russia. According to the number of SL users, the regions of Russia were divided (by lower and upper quartiles) into three groups: “low number” - 19 regions; “average number” - 36 regions; and “high number” (18 regions). There were some regions with the largest number of SL users: Moscow – 9,342; Moscow Oblast’ – 4,162; the Republic of Bashkortostan – 4,059; Sverdlovsk Oblast’ – 3,887; Chelyabinsk Oblast’ – 3,732; Rostov Oblast’ – 3,557; and

![Fig 1. Distribution of different languages used in Russia according to the number of respondents (the 2010 census).](https://doi.org/10.1371/journal.pone.0242219.g001)
Saint-Petersburg– 3,553 (S1 Fig and S2 Table). The regions with the lowest number of SL users were Chukotka Autonomous Okrug– 29; Evrei Autonomous Oblast’– 111; Magadan Oblast’– 161; Kamchatka Krai– 215; Sevastopol– 243; Yamal-Nenets Autonomous Okrug– 255; and the Republic of Altai– 284 (S1 Fig and S2 Table).

The proportion of SL users varied across federal regions of Russia, from 0.045% to 0.261%. We subdivided all regions into three categories according to the proportion of SL users: “low proportion”– 14 regions; “average proportion”– 48 regions; and “high proportion”– 11 regions (Fig 2 and S2 Table). The highest proportion of SL users was registered in the Republic of Tyva (Fig 2A)– 0.261% (CI– 0.244–0.028%). Other regions with a high proportion of SL users were as follows: the Republic of Sakha– 0.180% (CI– 0.171–0.188%); the Republic of Adygeya– 0.149% (CI– 0.138–0.161%); the Republic of Altai– 0.140% (CI– 0.124–0.157%); the Republic of Khakasiya– 0.134% (CI– 0.124–0.144%); and Orel Oblast’– 0.133% (CI– 0.125–0.142%). The lowest proportion of SL users– 0.045% (CI– 0.042–0.049%)–was registered in Khanty-Mansi Autonomous Okrug, although there were no significant differences compared to some other regions: Yamal-Nenets Autonomous Okrug (0.050%, CI– 0.044–0.057%); the Republic of

https://doi.org/10.1371/journal.pone.0242219.g002
of Mordovia (0.052%, CI– 0.048–0.056%); the Republic of Chechnya (0.052%, CI– 0.049–0.057%); and Chukotka Autonomous Okrug (0.060%, CI– 0.041–0.086%). Some other regions also had overlapping CIs (Fig 2 and S2 Table).

Discussion

In this study, by using census data, we determined the proportion and regional distribution of SL users in the total population (~ 140 million people) of Russia. Sign language, according to the total number of SL users, is in the thirty-second position in the list of 119 different native languages used in Russia (Fig 1 and S1 Table). The total number of SL users in Russia in 2010 was 120,528 (0.087%) out of 138,312,535 respondents, and the proportion of SL users varied from 0.045% to 0.261% in different regions of Russia.

The number of SL users varied from 29 in Chukotka Autonomous Okrug to 9,342 in Moscow, with a mean number of 1,466.63 per region, and corresponded to the number of respondents in the studied regions (S2 Table). The number of SL users in regions with a high number of respondents was also high, as expected. For example, the proportion of SL users in Moscow (0.084%, 9,342 out of 11,133,239 respondents) is close to the general proportion of SL users in Russia (0.087%). Similarly, in other regions with over 3,000,000 respondents, the proportion of SL users varied from 0.062% to 0.102% (S2 Table). These data can be relevant for health organizers planning the work of social security authorities and state health institutions.

To identify regions with values different from the general proportion of SL users in Russia (0.087%), we divided them into three groups by lower and upper quartiles: “low proportion” (14 regions), “average proportion” (48 regions), and “high proportion” (11 regions) (Fig 2 and S2 Table). The highest proportions of SL users were revealed in the Republic of Tyva (0.261%) and the Republic of Sakha (Yakutia) (0.180%). The values for the Republic of Adygeya (0.149%), the Republic of Altai (0.140%), the Republic of Khakasiya (0.134%), Orel Oblast’ (0.133%), Bryansk Oblast’ (0.121%), Pskov Oblast’ (0.119%), the Republic of Kalmykiya (0.113%), Penza Oblast’ (0.111%), and Chelyabinsk Oblast’ (0.110%) were also above the 75th percentile. Some of these regions of Russia (the Republic of Tyva, the Republic of Sakha (Yakutia), the Republic of Adygeya, the Republic of Altai, the Republic of Khakasiya, and the Republic of Kalmykiya) are inhabited, in addition to Russians, by different small indigenous peoples (Tuvinians, Yakuts, Adyge, Altaians, Khakas and Kalmyks). We suggest that the observed uneven distribution of SL users in these regions can represent the underlying hereditary forms of hearing impairment accumulated by a certain genetic background, a particular ethnic history and specific demographic factors in different ethnic groups living in Russia.

Numerous studies have shown that the frequencies of Mendelian diseases and different pathogenic variants can vary significantly among different regions of Russia [35–42]. Although Russians are the major ethnic group in Russia (111 million out of the total population of 146 million), there is a significant number of indigenous ethnic populations living for hundreds of years in their historical locations (over 200 different ethnicities and ethnic groups, according to the 2010 Census) unevenly dispersed across the vast territory of Russia [34]. The small size of these populations, isolation and specific mating traditions led to significant levels of endogamy [42–47]. These factors can explain the high rates of specific hereditary pathologies detected in different regions of Russia. For example, the founder effect and long isolation of some local populations in the Volga-Ural region of Russia determined a high prevalence of autosomal-recessive osteopetrosis [35] and autosomal-recessive hypotrichosis [37]; in the Caucasus regions of Russia, monoethnic marriages led to the highest world incidence of phenylketonuria [48]; and in Eastern Siberia, the high frequency of some specific Mendelian disorders was found to be caused by the founder effect [40].
The obtained data on high proportions of SL users in some regions of Siberia (the Republic of Tyva, the Republic of Sakha (Yakutia), and the Republic of Altai) are consistent with available data on the high prevalence of unique pathogenic variants in the genes associated with hearing impairment among indigenous populations in these regions (Tuvinians, Yakuts, and Altaians) [17, 18, 25, 29, 30, 49]. We suggest that a certain accumulation of hereditary forms of hearing loss can also be detected in other regions of Russia with a high proportion of SL users. Therefore, while SL knowledge is not defined exclusively by hearing status, the prevalence of SL users might be used as an indirect indicator of the accumulation of congenital or early-onset hereditary deafness, which, in turn, would determine the direction for more detailed genetic and epidemiological studies.

Supporting information

S1 Fig. The prevalence (in numbers) of SL users in different regions of Russia. A–Number of SL users across federal regions of Russia. The regions without available information are shown in gray. B–Comparison of the number of SL users in different regions of Russia. The regions are shown in three different colors according to the number of SL users: high values (above the 75th percentile) in red, average values (between the 25th and the 75th percentile) in yellow, and low values (under the 25th percentile) in green. (TIF)

S1 Table. The list of 119 native languages in Russia according to 138,312,535 completed questionnaires of the 2010 national census. (DOCX)

S2 Table. Distribution of SL users across the Russian Federation according to the 2010 national census. (DOCX)

Author Contributions

Conceptualization: Georgii P. Romanov, Nikolay A. Barashkov.

Formal analysis: Georgii P. Romanov, Aisen V. Solovyev, Olga L. Posukh, Nikolay A. Barashkov.

Funding acquisition: Sardana A. Fedorova.

Investigation: Georgii P. Romanov.

Methodology: Georgii P. Romanov.

Project administration: Sardana A. Fedorova, Nikolay A. Barashkov.

Supervision: Nikolay A. Barashkov.

Visualization: Georgii P. Romanov.

Writing – original draft: Georgii P. Romanov, Olga L. Posukh, Nikolay A. Barashkov.

Writing – review & editing: Georgii P. Romanov, Vera G. Pshennikova, Sergey A. Lashin, Aisen V. Solovyev, Fedor M. Teryutin, Aleksandra M. Cherdonova, Tuyara V. Borisova, Nikolay N. Sazonov, Elza K. Khusnutdinova, Olga L. Posukh, Sardana A. Fedorova, Nikolay A. Barashkov.
References

1. World Health Organization. Deafness and hearing loss [cited 2020 04.06.2020]. Available from: https://www.who.int/health-topics/hearing-loss#tab=tab_2.

2. Stevens G, Flaxman S, Brunsllik E, Mascaenhas M, Mathers CD, Finucane M, et al. Global and regional hearing impairment prevalence: an analysis of 42 studies in 29 countries. Eur J Public Health. 2013; 23(1):146–52. Epub 2011/12/24. https://doi.org/10.1093/eurpub/ckr176 PMID: 22197756

3. Pascolini D, Smith A. Hearing impairment in 2008: a compilation of available epidemiological studies. Int J Audiol. 2009; 48(7):473–85. https://doi.org/10.1093/ije/dyp044 PMID: 19444763

4. Krever AM, Smith RJ, Van Camp G, Schleiss MR, Bitner-Glindzicz MA, Lustig LR, et al. Congenital hearing loss. Nat Rev Dis Primers. 2017; (3):16094. Epub 2017/01/12. https://doi.org/10.1038/nrdp.2016.94 PMID: 28079113

5. Morton C, Nance W. Current concepts: Newborn hearing screening —A silent revolution. New England Journal of Medicine. 2006; 354(20):2151–64. https://doi.org/10.1056/NEJMra0507090 PMID: 16707752

6. Wroblewska-Seniuk KE, Dabrowski P, Szyfter W, Mazela J. Universal newborn hearing screening: methods and results, obstacles, and benefits. Pediatr Res. 2017; 81(3):415–22. Epub 2016/11/18. https://doi.org/10.1093/pedrsk/a0i051 PMID: 27861465

7. Butcher E, Dezateux C, Cortina-Borja M, Knowles RL. Prevalence of permanent childhood hearing loss detected at the universal newborn hearing screen: Systematic review and meta-analysis. PLoS One. 2019; 14(7):e0219600. Epub 2019/07/11. https://doi.org/10.1371/journal.pone.0219600 PMID: 31295316

8. Chibisova SS, Markova TG, Alekseeva NN, Yasinskaya AA, Tsygankova ER, Bliznetz EA, et al. [Epidemiology of hearing loss in children of the first year of life]. Vestn Otorinolaryngol. 2018; 83(4):37–42. https://doi.org/10.7116/otorino201883437 PMID: 30113578

9. Harris MS, Dodson EE. Hearing health access in developing countries. Curr Opin Otolaryngol Head Neck Surg. 2017; 25(5):353–8. https://doi.org/10.1097/MOO.0000000000000392 PMID: 28618006

10. Shein JD, Delk MT. The deaf population of the United States. First edition ed. Silver Spring, Md.: National Association of the Deaf; 1974. 336 p.

11. Li CM, Zhao G, Hoffman HJ, Town M, Themann CL. Hearing Disability Prevalence and Risk Factors in Two Recent National Surveys. Am J Prev Med. 2018; 55(3):326–35. Epub 2018/07/19. https://doi.org/10.1016/j.amepre.2018.03.022 PMID: 30031639

12. Mitchell RE. How many deaf people are there in the United States? Estimates from the Survey of Income and Program Participation. J Deaf Stud Deaf Educ. 2006; 11(1):112–9. Epub 2005/09/21. https://doi.org/10.1093/deafed/enn004 PMID: 16177267

13. Bubbico L, Rosano A, Spagnolo A. Prevalence of prelingual deafness in Italy. Acta Otorhinolaryngol Ital. 2007; 27(1):17–21 PMID: 17601206

14. Zinchenko R, El'chinova G, Baryshnikova N, Polyakov A, Ginter E. Prevalences of hereditary diseases in different populations of Russia. Russian Journal of Genetics. 2007; 43(9):1038–45. https://doi.org/10.1134/S1022795407090104 PMID: 17990523

15. Zinchenko R, Elchinova G, Gavrilina S, Ginter E. Analysis of diversity of autosomal recessive diseases in populations of Russia. Russian Journal of Genetics. 2001; 37(11):1312–22. https://doi.org/10.1023/A:1012569411698

16. Barashkov NA, Romanov GP, Borisova JP, Solovyov AV, Pshennikova VG, Teryutin FM, et al. A novel pathogenic variant c.975G>T (p.Trp325*) in the POU3F4 gene in Yakut family (Eastern Siberia, Russia) with the X-linked deafness-2 (DFNX2). International Journal of Pediatric Otorhinolaryngology. 2018; 104:94–7. https://doi.org/10.1016/j.ijporl.2017.11.001 PMID: 29287890

17. Barashkov NA, Pshennikova VG, Posukh OL, Teryutin FM, Solovyov AV, Klarov LA, et al. Spectrum and Frequency of the GJB2 Gene Pathogenic Variants in a Large Cohort of Patients with Hearing Impairment Living in a Subarctic Region of Russia (the Sakha Republic). Plos One. 2016; 11(5): e0156300. https://doi.org/10.1371/journal.pone.0156300 PMID: 27224056

18. Barashkov NA, Romanyuk GP, Borisova JP, Solovyov AV, Pshennikova VG, Teryutin FM, et al. A rare case of Waardenburg syndrome with unilateral hearing loss caused by nonsense variant c.772C>T (p.
21. Bliznetz EA, Lalayants MR, Markova TG, Balanovskiy OP, Balanovskya EV, Skhalyakho RA, et al. Update of the GJB2/DFNB1 mutation spectrum in Russia: a founder Ingush mutation del(GJB2-D13S175) is the most frequent among other large deletions. J Hum Genet. 2017; 62(8):789–95. Epub 2017/04/13. https://doi.org/10.1038/jhg.2017.42 PMID: 28405014

22. Bliznetz EA, Galkina VA, Matushchenko GN, Kisina AG, Markova TG, Poliakov AV. Changes in the connexin 26 gene (GJB2) in Russian patients with hearing loss: Results of long-term molecular diagnostics of hereditary nonsyndromic hearing loss. Russian Journal of Genetics. 2012; 48(1):101–12. https://doi.org/10.1134/S1022795412010036

23. Bliznetz EA, Makienko ON, Okuneva EG, Markova TG, Poliakov AV. New recurrent large deletion, encompassing both GJB2 and GB6 genes, results in isolated sensorineural hearing impairment with autosomal recessive mode of inheritance. Russ J Genet. 2014; 50(4):415–20. https://doi.org/10.1134/S1022795414020045

24. Bozhkova VP, Khashaev ZH, Umanskaya TM. Frequency and the mutation spectrum of GJB2-related hearing loss in children of Dagestan as compared with the central European part of Russia. Biophysics. 2010; 55(3):453–62. https://doi.org/10.1134/S0006350910030176

25. Churbanov A, Karafet T, Morozov I, Mikhalskia V, Zytsar M, Bondar A, et al. Whole Exome Sequencing Reveals Homozygous Mutations in RA1, OTOF, and SLC26A4 Genes Associated with Nonsyndromic Hearing Loss In Altai Families (South Siberia). Plos One. 2016; 11(4). https://doi.org/10.1371/journal.pone.0153841 PMID: 27082237

26. Zhuravskii S, Grinchik O, Taraskina A, Ivanov S, Galkin V. Mutatsiya 35delG gena konneksina-26 kak prichina prelingual'noy tukoukhosti v Arkhangelskoy oblasti [35delG mutation of the connexin-26 gene as a cause of prelingual sensorineural hearing loss in the Arkhangelsk region]. Ekologiya cheloveka [Human ecology]. 2008;(7):53–6.

30. Posukh O, Pallares-Ruiz N, Tadinova V, Osipova L, Claustres M, Roux A. First molecular screening of deafness in the Altai Republic population. Bmc Medical Genetics. 2005; 12;(6). https://doi.org/10.1186/1471-2350-6-12 PMID: 15790391

31. Posukh O, Zytsar M, Bady-Khoo M, Danilchenko V, Maslova E, Barashkov N, et al. Unique Mutational Spectrum of the GJB2 Gene and Its Pathogenic Contribution to Deafness in Tuvinians (Southern Siberia, Russia): A High Prevalence of Rare Variant c.516G > C (p.Trp172Cys). Genes. 2016; 7(7):688–97. https://doi.org/10.3390/genes70700688 PMID: 27407781

33. Zhuravskii S, Taraskina A, Podlesnyi E, Baldakova O, Ivanov S. Mutatsiya 35delG gena konneksina-26 kak prichina prelingual’noy tukoukhosti v Arkhangelskoy oblasti [35delG mutation of the connexin-26 gene as a cause of prelingual sensorineural hearing loss in the Arkhangelsk region]. Ekologiya cheloveka [Human ecology]. 2008;(7):53–6.

34. Federal State Statistics Service of Russian Federation. All-Russian Census 2010 data. Available from: https://glk.ru/free_doc/new_site/perepis2010/croc/perepis_itogi1612.htm.

35. Bliznetz E, Tverskaya S, Zinchenko R, Abrukova A, Savaskina E, Nikulin M, et al. Genetic analysis of autosomal recessive osteopetrosis in Chuvashia: the unique splice site mutation in TICIR1 gene spread by the founder effect. European Journal of Human Genetics. 2009; 17(5):664–72. https://doi.org/10.1038/ejhg.2008.294 PMID: 19172990
36. Gundorova P, Zinchenko R, Makaov A, Polyakov A. The Spectrum of Mutations in the PAH Gene in Patients with Hyperphenylalaninemia from the Karachay-Cherkess Republic. Russian Journal of Genetics. 2017; 53(7):813–9. https://doi.org/10.1134/S1022795417070043

37. Kazantseva A, Goltsov A, Zinchenko R, Grigorenko A, Abrukova A, Moliak Y, et al. Human hair growth deficiency is linked to a genetic defect in the phospholipase gene LIPH. Science. 2006; 314(5801):982–5. https://doi.org/10.1126/science.1133276 PMID: 17095700

38. Marakhonov AV, Konovalov FA, Makaov AK, Vasilyeva TA, Kadyshchev VV, Galkina VA, et al. Primary microcephaly case from the Karachay-Cherkess Republic poses an additional support for microcephaly and Seckel syndrome spectrum disorders. BMC Med Genomics. 2018; 11(8):91–95. Epub 2018/02/13. https://doi.org/10.1186/s12920-018-0326-1 PMID: 29504900

39. Petrova NV, Kashirskaya NY, Vasilyeva TA, Timkovskaya EE, Voronkova AY, Shabalo LA, et al. High prevalence of W1282x mutation in cystic fibrosis patients from Karachay-Cherkessia. J Cyst Fibros. 2016; 15(3):e28–32. https://doi.org/10.1016/j.jcf.2016.02.003 PMID: 26948992

40. Puzyrev VP, Maximova NR. Hereditary diseases among Yakuts. Russian Journal of Genetics 2008: 44 (10):1141–47. https://doi.org/10.1134/S1022795408100037 PMID: 19062529

41. Tarskaia L, Zinchenko R, Elchinova G, Egorov A, Korotov M, Basova E, et al. The structure and diversity of hereditary pathology in Sakha Republic (Yakutia). Russian Journal of Genetics. 2004; 40 (11):1264–72. https://doi.org/10.1023/B:RUGE.0000048669.22362.1c PMID: 15612572

42. Zinchenko R, Elchinova G, Petrova N, Osipova E, Malyshov P, Polyakov A, et al. Genetic structure of the Udmurt population. Russian Journal of Genetics. 2007; 43(8):918–28. https://doi.org/10.1134/S1022795407080145

43. El’chinova G, Shakmanov M, Revazova Y, Zinchenko R. Population and genetic characteristics of Abazins in Karachay-Cherkessia (marital migrations and surname frequency distribution). Russian Journal of Genetics. 2015; 51(10):1020–5. https://doi.org/10.1134/S1022795415100051 PMID: 27169233

44. El’chinova G, Simonov Y, Vafina Z, Zinchenko R. Endogamy and isolation by distance in the Tatarstan population. Russian Journal of Genetics. 2011; 47(8):999–1003. https://doi.org/10.1134/S1022795411080059 PMID: 21954622

45. Goltsov T, Osipova L. Genetic demography structure of aborigine Siberian populations in connection with problems of microevolution. VOGIS Herald. 2006; 10(1):126–54.

46. Kurbatova O, Yankovsky N. Migration as the main factor of the Russia’s urban population dynamics. Russian Journal of Genetics. 2016; 52(7):726–45. https://doi.org/10.1134/S1022795416070061 PMID: 29368870

47. Kurbatova OL, Pobedonostseva EY, Privalova VA. Strategies of adaptation: interpopulation selection differentials. J Physiol Anthropol Appl Human Sci. 2005; 24(4):363–5. https://doi.org/10.2114/jpa.24.363 PMID: 16079581

48. Gundorova P, Zinchenko R, Kuznetsova I, Bliznetz E, Stepanova A, Polyakov A. Molecular-genetic causes for the high frequency of phenylketonuria in the population from the North Caucasus. Plos One. 2018; 13(8). https://doi.org/10.1371/journal.pone.0201489 PMID: 30067850

49. Zytsar MV, Bady-Khoo MS, Danilenko VY, Maslova EA, Barashkov NA, Morozov IV, et al. High Rates of Three Common GJB2 Mutations c.516G>C, c.-23+1G>A, c.235delC in Deaf Patients from Southern Siberia Are Due to the Founder Effect. Genes. 2020; 11(7):833. https://doi.org/10.3390/genes11070833