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Abstract

A total of 112 Nagaybaks, a Turkic ethnoconfessional group living mainly in the Nagaybak district of the Chelyabinsk Region of Russian South Urals, were genotyped for HLA-A, -B, -DRB1, -DQA1 and -DQB1 loci using PCR-SSP (low-resolution) and HLA-A29 (high-resolution). All loci were in Hardy-Weinberg equilibrium (all \( p \) values >0.1 thus showing no locus-level deviations. The genotype data are available in the Allele Frequencies Net Database under the population name “Russia, South Ural, Chelyabinsk Region, Nagaybaks” and the identifier (AFND0003397).
Nagaybaks (Nağaybäks, Nogaybaks, Nogoybaqs) are a Turkic ethnoconfessional group that is close to the Kryashen Tatars. They live mainly in the Nagaybak district of the Chelyabinsk Region. According to the latest census (National Population Census, 2010) some 7,679 Nagaybaks, comprising 94.2% of all Nagaybaks in Russia, live in the Chelyabinsk Region.

There are several theories regarding Nagaybak ethnogenesis. The first states that Nagaybaks originated from the Volga region and supposedly are descendants of Nogay and Kipchak peoples, the general population of the Nogai Horde [1]. The Nogai Horde was formed in the 15th/16th Centuries after the collapse of the Golden Horde on the left bank of the lower Volga, the territory of today's Bashkortostan, Chelyabinsk and Orenburg Regions and western and central Kazakhstan. Another version of Nagaybaks’ ethnogenesis suggests that ancestors of Nagaybaks came from Kazan Khanate, another state formed after the collapse of the Golden Horde. The Arsk region was a part of Kazan Khanate. A further theory states that the Nagaybaks were Tatarized Finno-Ugric peoples that kept the Kazan Khanate's borders [1].

Early historical records indicate that since ancient times and up until 1736 Nagaybaks lived separately from Bashkirs and Tatars away from Russian cities and villages in Ufa Country (today's Bashkiria). They spoke a dialect of the Tatar language, professed orthodoxy and paid tribute to the Russian Tsar. In 1736 the Russian tsarist government built a town for them and named it the "Nagaybak’s Fortress”. The inhabitants were called "Nagaybaks" and they became Cossacks under the decree of the empress.

In 1773 the Nagaybak population of the Fortress and surrounding 14 villages was about 2,000. But among them there were some 200 other nationalities, including Bashkirs and others, notably Persians, Arabs, Arabians, Armenians, Afghans and Turks. Cossack-Nagaybaks, as a military caste, took part in all internal Russian conflicts and foreign campaigns of Russia since the 18th century. In 1864, according to imperial decree, the Nagaybaks were ordered to move to the territory of the modern Nagaybak’s districts of the Chelyabinsk Region called Parizh, Cassel and Fershampenuaz. They moved due to changes in the borders of Russia and the establishment of fortresses and Cossack settlements on the "New Orenburg line". Thus, the majority of Nagaybaks have lived in the Chelaybinsk Region from around 1864 to the present day.

It is clear that the Nagaybaks were in geographic and cultural isolation from their main body of close relatives, the Volga-Urals Tatars. By the beginning of the 20th century Nagaybaks had acquired traits expected of an ethnologically-independent unit and spoke a sub-dialect of the Tatar language [1].
All subjects were normal, healthy unrelated blood donors, of between 18 and 55 years of age, living in the Chelyabinsk Region (South Urals, Russia). Their ethnic origin was determined by a comprehensive questionnaire. Only subjects with a Nagaybaks origin spanning at least three generations were included in the study. All 112 Nagaybaks (54 men and 58 women) were Russian language speakers and some 95% spoke the Nagaybaks language (a dialect of the Tatar language). One hundred and nine lived in the Nagaybak’s districts of the Chelyabinsk Region (38 from Kassel'skij, 31 from Fershampenuaz, 19 from Parizh, 16 from Ostrolenka and 5 from Nagajbakskij) (Supplementary Figure 1). The remaining 3 came from other cities of the Region. DNA was obtained from EDTA-anticoagulated peripheral blood using AxyPrep spin columns (AxyPrep_Blood Genomic DNA Miniprep Kit; Axygen Biosciences, Union City, CA, USA) according the manufacturer’s protocol.

HLA-A, -B, -DRB1 and -DQB1 typing was performed by PCR using sequence-specific primers (SSP). The primers and primer mixtures used were essentially those of Downing et al. [2]. HLA-DQA1 typing to the 2nd field and HLA-A*29 typing, to the 2nd field, was undertaken using commercial kits (One Lambda, Canoga Park, CA, USA).

Population genetics analysis was performed as described by Schipper et al. [3]. The validity of Hardy–Weinberg equilibrium and homozygosity was tested for each locus. Carriage frequencies (cf) and gene frequencies (gf) were determined by direct counting and maximum likelihood, respectively. Two- and three-locus haplotype frequency (HF) estimates were calculated by maximum likelihood. The linkage disequilibrium (LD) parameter (Δ), the relative magnitude of the delta value (Δrel) and the significance of delta values for two- and three-locus haplotypes were determined as previous described [4].

HLA-A, -B and -DRB1 frequency data were used for the construction of a dendrogram using the maximum likelihood method (PHYLIP version 3.68) [5]. Thus, the Nagaybak population was compared with populations of Russians, Tatars and Bashkirs of the Chelyabinsk Region [6] and with 18 other populations [6, 7, 8, 9, 10]. The findings of the Hardy-Weinberg and homozygosity analyses for HLA-A, -B, -DRB1, -DQA1 and -DQB1 all showed an acceptable goodness-of-fit for both phenotype distribution and the number of likely homozygotes identified (all p values >0.1). Allele family frequencies are presented in Supplementary Table 1, 2-loci HF and LD data in Supplementary Table 2, 3-loci HF and LD data in Supplementary Table 3, and 5-loci HF data in Supplementary Table 4. Supplementary Table 5 shows the findings of HLA-A*29 typing to the 2nd field. Prior to taking blood samples full informed consent for the collection and use of specimens was given by all
donors. All obtained data are publically available at Allele Frequencies Net Database AFND0003397.

A dendrogram, comparing the Nagaybak population with other populations of the Chelyabinsk Region and populations worldwide, is presented in Supplementary Figure 2.

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References


