

## ARTICLE

## Genetic analysis of mitochondrial ND5 gene of siberian ibex (*Capra Sibirica*, Pallas, 1776) population in Mongolia

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**Abstract:** The Siberian ibex (*Capra sibirica*) from Central Asia is believed to be the most ancient species of the genus *Capra*. In Mongolia, it is distributed in the areas of Mongolian Altai, Gobi-Altai, Dzungaria, Altai, Khan Khuhii, Khoridal Saridag and Ulaan Taiga as well as in the desert and semi-desert steppe zones of Dundgobi and Dornogobi aimags (provinces). In the current study, we investigated the mitochondrial ND5 gene fragments of the Siberian ibex population from different parts of Mongolia. Nine haplotypes, including 6 shared and 3 unique haplotypes were identified among these populations. Furthermore, Tajima's statistics and Fu's statistics did not reveal significant positive value across the population, indicating population decline and balancing selection. In the phylogenetic tree by 9 haplotypes, no separated clusters were generated. In addition, nucleotide diversity was 0.015, haplotype diversity was 0.86 and the average number of differences in nucleotides was 8.2 in the overall population. These results suggest that genetic diversity across all the populations was low, while haplotype diversity and the average number of differences in nucleotides were high.

**Keywords:** Siberian ibex; mitochondrial ND5 gene; haplotype; population genetic diversity; *Capra sibirica*;

### INTRODUCTION

Classification of the genus *Capra* has been a complicated issue in modern taxonomy studies of ungulates. In a recent molecular genetic study, the genus *Capra* contains eight different species: bezoar (*C. aegagrus*), Alpine ibex (*C. ibex*), Siberian ibex (*C. sibirica*), Nubian ibex (*C. nubiana*), Spanish ibex (*C. pyrenaica*), markhor (*C. falconeri*), Kuban tur (*C. caucasica*), and Dagestan tur

(*C. cylindricornis*). The Siberian ibex from Central Asia is believed to be the most ancient species of the genus *Capra* [5]. According to the International Union for the Conservation of Nature Red List Categories, this species is listed as Least Concerned at the global level and Near Threatened at the regional level [1]. There are two different subspecies of Siberian ibex (*C. sibirica*) in terms of geographical

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and genetical characteristics: which are *C. sibirica* and *C. s. sakeen* [5,6]. In Mongolia, the Siberian ibex is unevenly distributed in the mountainous areas of Mongolia Altai, Gobi-Altai, Dzungaria, Altai, Khan khuhii, Khoridal Saridag and Ulaan Taiga as well as in desert and semi-desert steppe zones of Dundgobi and Dornogobi aimags [1]. In the early 1980s, Siberian ibex population was estimated at 80.0 thousand in the country, of which around one thousand were found in the red taiga mountains of Khoridal Saridag in Khuvsgul aimag [4].

A 2009 national estimation report of Mongolian mountain ungulates showed that the Siberian ibex was distributed over a total area of 55,985 km<sup>2</sup> and the population was estimated as 36.0 thousand in the country [9]. Since ancient times, its population has been declining steadily due to constant tropical hunting, growing acreage of domestic livestock pasture, parasites as well as severe

cold winters with heavy snowfall, causing mass deaths [3,10,13]. Mitochondrial DNA is one of the most reliable tools for population genetic studies thanks to its relatively high and rapid mutational rate [17]. Also, mitochondrial DNA contains the hypervariable and conserved regions, making them suitable for taxonomic studies [12]. In addition, mitochondrial ND5 gene has been used as a powerful genetic marker to reveal phylogenetic divergence and relationships between and within closely related species in some studies [8,15].

In order to address the issue of protecting endangered species, it is important to study their classification, phylogenetic linkages and population genetic structure at the molecular level. In the current study, we aimed to conduct molecular genetic analysis by mitochondrial ND5 gene of Siberian ibex from different parts of Mongolia.

## MATERIALS AND METHODS

In order to investigate the genetic differentiation among Siberian ibex from different parts of Mongolia, phylogenetic analysis of mitochondrial ND5 gene, which encodes NADH-ubiquinone oxidoreductase chain 5 protein of the electron transport chain, was carried out. Samples of Siberian ibex (*Capra sibirica*) population in different parts of Mongolia were analyzed. 25 samples from the Altai region, including 8 individuals (sample code: AB1-AB8) from Bayankhongor, 15 individuals (sample code: AH9-AH23) from Khovd, and 2 individuals (sample number: AU24-AU25) from Uvs aimags, were analyzed. Also, 8 samples from the Gobi region and the Khoridal saridag protected are in Khuvsgul, which included 5 individuals (sample code: GI26-GI30) from Ikh nart, Dornogobi and 3 individuals (sample number: KhS31-KhS33) from Khoridal Saridag were analyzed. DNA was isolated from fecal pellets, skin and muscle samples. DNA extraction from fecal pellets was performed using DNA extraction

method from the feces of *Caprinae* [16].

DNA extraction from other types of tissue samples, such as skin and muscles, was performed using QIAgen tissue kit. Mitochondrial ND5 gene fragment was amplified using primers **ND5L** (5'-AATAGTTTATCCAGTTGGTCTTAGG-3'), **ND5RI** (5'-AAGATTTGTTGGAGATCTCAGGTG-3'). PCR amplifications were performed in a total volume of 50 µl containing 2.5 units of *iTaq*<sup>TM</sup> DNA polymerase, 5 µl of 10x PCR buffer, 3 µl of 25 mM MgCl<sub>2</sub>, 5 µl of 2.5 mM dNTPs, 2 µl of 10µM each of the primers and 1µl of genomic DNA. Thermal conditions for PCR are as follows: 94°C for 3 min, followed by 35 cycles of 94°C for 30 sec, 55°C for 30 sec, 72°C for 1 min, and final extension at 72°C for 5 min. The PCR products were sequenced in Macrogen Inc, Korea. Sequence data were edited and compiled using Bioedit v.7.2.5 software. Alignment of the sequences was performed by NCBI Blast program (<https://blast.ncbi.nlm.nih.gov/Blast>).

cgi). Construction of phylogenetic tree by neighbor-joining method was performed using MEGA v.10 software. Genetic diversity at the

molecular level was calculated using DnaSP v.5.2. Statistical analysis of population genetics was performed using Arlequin v. 2.000.

## RESULTS AND DISCUSSION

We have successfully amplified 656 bp fragments of ND5 gene and 554 bp fragments of which was analyzed for the current phylogenetic study. Within ND5 gene fragments, 22 variable sites were identified with transition/transversion ratio of 22:1, and 20 parsimony informative sites and 2 singleton sites were among these variable sites. Additionally, constitutions of nitrogenous bases across the populations were C: 42.56%, T: 23.00%, A : 14.6%, G : 19.83%. Nine haplotypes, which includes 6 shared (H1, H2, H3, H4, H7, H9) and 3 unique haplotypes (H5, H6, H8) were detected among 33 individuals of all the populations from Mongolia. Relative frequency of each haplotypes in overall population was between 3% and 27.3%. The most common haplotype H2 (27.3%) was found in 7 individuals of the Siberian ibex population in Bayankhongor aimag, 2 individuals of the Siberian ibex populations in Uvs aimag in the Altai region. The most number of haplotypes (H3, H5, H6, H7, H8, H9) were detected in 15 individuals of Siberian ibex population in Khovd aimag in the Altai region. In addition, unique haplotypes

H1 and H4 were found in 3 individuals of Khoridal Saridag, Khuvsgul and 5 individuals of Ikh Nart, Dornogobi, respectively (Table 1). In the Siberian ibex population in Mongolia, nucleotide diversity presented a low value, while haplotype diversity and average number of differences in nucleotides presented high values (Table 2). For that reason, Altai population, including large numbers of individuals of Siberian ibex populations from different provinces such as Uvs, Khovd, Bayankhongor might have influenced the high value of haplotype diversity and the average number of nucleotide differences due to high number of pairwise differences between individuals arising from a large number of base substitutions. Furthermore, Tajima's statistics' and Fu's statistics did not reveal significant positive value for the populations (Table 2). When positive Tajima's *t* value is signified, it means low levels of both low and high frequency polymorphisms, indicating population decline and balancing selection [14]. Similarly, positive Fu's *F<sub>st</sub>* indicates a lack of low frequency polymorphism [7].

Table 1. Variable sites of mitochondrial ND5 gene in Siberian ibex (*Capra sibirica*) samples. 22 Polymorphic sites in 554bp fragment of ND5 gene

Haplotype	11783	11815	11836	11865	11870	11876	11891	11894	11902	11915	11960	11966	11990	11999	12014	12027	12072	12134	12200	12204	12233	12269	Sample code (number of samples)	Reference sequence
AB743824	G	G	C	G	G	T	A	C	T	C	C	C	A	T	C	C	C	T	C	C	A	C	C	KhS31, KhS32, KhS33 (3)
H1	.	.	.	.	A	C	G	.	C	T	T	T	.	.	T	.	.	.	G	.	.	.	.	KhS31, KhS32, KhS33 (3)
H2	.	.	.	.	.	.	.	.	.	.	.	.	.	.	.	.	.	.	.	.	.	.	.	AB1 ,AB2, AB3, AB4, AB6, AB7, AB8, AU24, AU25 (9)
H3	.	A	T	A	.	C	G	.	C	T	.	.	G	C	T	T	.	.	A	T	.	.	AB5, AH13, AH19, AH20, AH21, AH23 (6)	
H4	.	.	.	.	.	C	G	.	C	T	T	T	.	.	T	T	C	C	G	.	.	.	GI26, GI27, GI28, GI29, GI30 (5)	
H5	.	.	.	.	.	.	.	.	.	.	.	.	.	.	.	.	.	.	.	.	.	T	.	AH22 (1)
H6	.	.	.	.	C	.	.	.	.	.	.	.	.	.	.	.	.	.	.	.	G	.	.	AH18, (1)
H7	.	.	.	.	C	.	.	.	.	.	.	.	.	.	.	.	.	.	.	.	.	.	.	AH9, AH11, AH17 (3)
H8	.	.	.	.	C	G	.	.	C	T	T	.	.	.	T	T	C	C	G	.	.	.	.	AH16 (1)
H9	A	A	T	A	.	C	G	T	C	T	.	.	G	C	T	T	.	.	A	T	.	.	.	AH10, AH12, AH14, AH15 (4)

Table 2. Result of genetic diversity analysis of Siberian Ibex (*Capra sibirica*) in Mongolia.

N	H	S	Hd	$\pi$	k	Tajima D	Fu's Fst
33	22	9	0.862	0.01481	8.025	1.55444	3.94

N - Number of Samples, S - Segregating sites, H - Haplotype number, Hd - Haplotype diversity;

$\pi$  - Nucleotide diversity, k - Average number of nucleotide differences

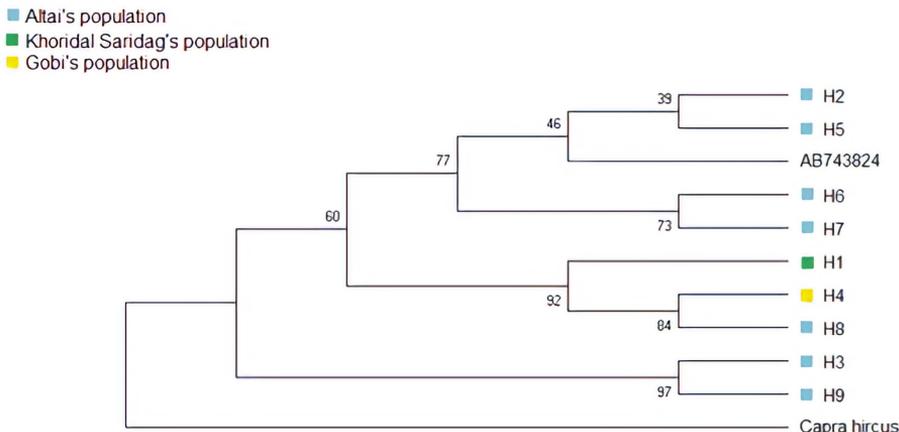


Figure 1. Phylogenetic tree of mitochondrial ND5 haplotypes (554 bp fragment) constructed using Neighbor-joining method. Numbers in nodes indicate bootstrap values >39 with 1000 replicates.

In the current study, we conducted the mitochondrial ND5 gene analysis of Siberian ibex (*Capra sibirica*) population in Mongolia for the first time. We detected a total of 9 haplotypes, H1-H9, and except haplotype H2, none of which have been registered in the NCBI Genbank. In addition, there are some genetic studies based on the mitochondrial DNA segments, including protein coding regions, to uncover the domestication process of domestic goats (*Capra hircus*) and their genetic characterization [2, 11]. As a result of these studies, haplotype H2, detected from Siberian ibex in Mongolia was registered in the Genbank. Besides, haplotype H2 was identified as the most common shared haplotype in our

study. So, this could be a broadly distributed haplotype in the Siberian ibex population in Mongolia. Also, the mitochondrial ND5 gene analysis of Siberian ibex is still not widely studied. We, therefore, hope that our data will represent Mongolian Siberian ibex population and contribute to the future population genetic studies of genus *Capra* based on the mitochondrial ND5 gene. Furthermore, a few samples from the Gobi region and the protected area of Khoridal Saridag was included in this study. Therefore, It is highly recommended to conduct further research on the mitochondrial ND5 gene of Siberian ibex with added number of samples from Gobi and Khoridal Saridag's populations to verify the result.

## CONCLUSIONS

In conclusion, genetic diversity across all the populations of Siberian ibex in Mongolia presented a low value, while haplotype diversity and the average number of differences in nucleotides presented high values. Nevertheless, the Altai population, which includes a large number of samples from different provinces, could have influenced the result.

Furthermore, the Siberian ibex population in Khovd aimag of Altai region had the most haplotype diversity. Moreover, both the populations from the protected area of

Khoridal Saridag in Khuvsgul and Ikh Nart, in Dornogobi aimags had their own one unique haplotypes. Thus, the geographical fragmentation might have caused the genetic differentiation across all the populations.

In the future, high sensitive genetic markers such as mitochondrial control region and microsatellite marker studies are recommended in order to obtain more information on the phylogenetic relationships of all the population.

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