Genetic characterization of *Echinococcus* isolates from various intermediate hosts in the Qinghai-Tibetan Plateau Area, China

Xumin Han1,*, Yingna Jian2,*, Xueyong Zhang2, Liqing Ma2, Wenjun Zhu1, Qigang Cai2, Shile Wu1, Xiangqian Wang3 and Bingqiang Shi3

1Clinical Research Institute of Hydatid Disease, Qinghai Provincial People’s Hospital, Xining 810007, China and 2Qinghai Academy of Animal Sciences and Veterinary Medicine, Qinghai University, State Key Laboratory of Plateau Ecology and Agriculture, Qinghai University, Xining 810016, China

**Abstract**

This study examined *Echinococcus* spp. genotypes and genetic variants isolated from humans as well as domestic and wild animals from the Qinghai-Tibetan Plateau Area using the cox1 gene. All samples except the pika isolates were identified as the *Echinococcus granulosus* sensu stricto. Sixteen different haplotypes with considerable intraspecific variation were detected and characterized in mitochondrial cox1 sequences. The parsimonious network of cox1 haplotypes showed star-like features, and the neutrality indexes computed via Tajima’s D and Fu’s Fs tests showed high negative values in *E. granulosus* s.s., indicating deviations from neutrality; the Fst values were low among the populations, implying that the populations were not genetically differentiated. The pika isolates were identified as *E. multilocularis* and *E. shiquicus*. Only one haplotype was recognized in the pika isolates. *E. granulosus* s.s. was the predominant species found in animals and humans, followed by *E. multilocularis* and *E. shiquicus*, with high genetic diversity circulating among the animals and humans in this area. Further studies are needed to cover many sample collection sites and larger numbers of pathogen isolates, which may reveal abundant strains and/or other haplotypes in the hydatid cysts infecting human and animal populations of the QTPA, China.

**Introduction**

Echinococcosis is caused by the metacestodes of *Echinococcus* spp. and is parasitic in the livers, lungs and/or other organs of humans and animals, resulting in serious zoonotic parasitic disease. Echinococcosis epidemic diseases not only threaten and harm the health of humans and animals but also seriously hinder the production of animal husbandry and affect the development of the national economy (Budke et al., 2006; Qian et al., 2017). This disease is prevalent in Asia, South America, North Africa, Central Europe and other regions and causes harm throughout the world (Grosso et al., 2012). China, especially the Qinghai-Tibetan Plateau Area (QTPA), has one of the highest prevalence rates in the world (Wang et al., 2014). Echinococcosis identified as a zoonotic disease with a major impact on the public health of rural populations in Qinghai-Tibetan Plateau. As our previous study showed a high seropositive rate (37.0%) of echinococcosis in Qinghai-Tibetan primary school students (Han et al., 2018). Prevalence of *E. granulosus* in yaks, pigs and Tibetans investigated in Qinghai-Tibetan Plateau was 6.49, 7.27 and 1.83%, and Prevalence in yaks was 3.61, 9.66 and 6.33% in 2014, 2015 and 2016, respectively (Li et al., 2017). The action plan for prevention and treatment of echinococcosis was carried out by the People’s Republic of China. A series of work was being implemented, such as the management and deworming of source of infection-dog, the vaccine immunization of the livestock, the implementation of livestock slaughter standard in slaughtering house (safety disposal of diseased organs), the management of patients (patient care and rescue), health education, people training and providing safe drinking water. The seven provinces and autonomous regions in northwestern China are epidemic areas of hydatid disease, endangering nearly 50 million people and 70 million livestock, which has resulted in direct economic losses of up to 30 billion Yuan (Qian et al., 2017). *Echinococcus* spp. exhibit a fixed life cycle of definitive and intermediate hosts. Carnivores, such as dogs, foxes and wolves, are the definitive hosts of the parasites. The intermediate hosts involve different species and change in different environments (Romig et al., 2017). *Echinococcus* spp. can be infective in metacestode hosts, such as humans, livestock and some wild animals (Thompson, 2017). With the continuous improvement of molecular and genetic knowledge of parasites, identification and classification research has surpassed morphological study (Nakao et al., 2013). Genetic analysis employing mitochondrial genes (cox1, nad1, cob and nad5 gene) and ribosomal genes might reveal the biological relationships between *Echinococcus* spp. and strains (Marinova et al., 2017; Kinkar et al., 2018a). Mitochondrial DNA (mtDNA) is the most suitable genetic marker for the analysis of genetic diversity, genetic differentiation and evolution of species and is widely used in species classification (Umhang et al., 2014). There are five epidemic *Echinococcus* spp.: *E. granulosus* s.l., *E. multilocularis*, *E. oligarthrus*, *E. vogeli* and *E. shiquicus* (Nakao et al., 2013).
**DNA extraction and PCR amplification**

To extract genomic DNA from cysts, including the protoscoleces and germinal layers, each individual cyst was washed at least three times with sterile distilled water and centrifuged to remove the salt ions from PBS, after which genomic DNA was extracted according to the manufacturer’s instructions (TIANamp Micro DNA Kit, Code: DP316, Tiangen, Beijing, China). The genomic DNA concentration was measured using a spectrophotometer (Merck Millipore, Frankfurt, Germany), and the DNA was then stored at −20 °C until being used for PCR amplification. A fragment of the mitochondrial genes was amplified from each sample using the primers described by Liu et al. and Bowles et al. for detection and analysis (Bowles et al., 1992, 1994; Liu et al., 2015). The PCR conditions and procedures were modified slightly, and the primers were cox1F: 5′- CCTGATTGTGTTAATAATTCGA-3′ and cox1R: 5′- ATCATGAAATTACATTATCA-3′ (product = 366 bp). Each PCR mixture had a total volume of 50.0 µL containing 25.0 µL of Premix Taq™ (TaKaRa Taq™ Version 2.0, Code: R004, Takara Bio Inc, Tokyo, Japan), 2.0 µL of each primer (10.0 µM), 2.0 µL of template DNA, and 19.0 µL of deionized distilled water. Positive and negative controls were run in parallel with the PCR amplification of the DNA samples. PCR amplification was carried out in a thermocycler (Masterecycler nexus GXX1, Eppendorf, Saxony, Germany) with a 5.0 min initial denaturation step at 95.0 °C; 35 cycles of a 35 s denaturation at 94.0 °C, a 45 s of annealing at 54.5 °C and a 40.0 s of extension at 72.0 °C; and a 10.0 min final extension at 72.0 °C. In the next step, the PCR products of the cox1 gene were electrophoresed in 1.2% agarose gels and stained with nucleic acid dyes. The PCR product bands were observed under UV light and recorded as digital images with a gel documentation system (BEIJING LIUYI BIOTECHNOLOGY CO., LTD., Beijing, China).

**Sequencing and alignment analysis**

A total of 244 PCR products (product = 366 bp) from different hosts were sequenced using both the forward and reverse primers by the GENEWIZ Company (Suzhou, China). Sequences were identified and compared in the GenBank database through BLAST analysis (https://blast.ncbi.nlm.nih.gov/Blast.cgi?), developed by the National Center for Biotechnology Information (https://www.ncbi.nlm.nih.gov/). Amino acid sequences were inferred from the nucleotide sequences based on echinoderm and flatworm mitochondrial genetic codes using the ExPaSy translate tool (https://web.expasy.org/translate/). Then, the sequences were subjected to multiple alignment by using the Clustal Omega alignment programme (http://www.ebi.ac.uk/Tools/msa/clustalo/) with reference sequences from different E. granulosus s. s. genotypes and Echinococcus spp.
**Data analysis**

DnSP 5.0 software was used to convert the fasta format (.fas) to the Network data format (.rdt) ARLEQUIN data format (.arp), and population diversity indexes (number of haplotypes and haplotype diversity) could also be calculated with this software. Then, we used NETWORK 5.0 (http://www.fluxus-engineering.com) to construct haplotype median-joining networks (Bandelt et al., 1999). Networks were constructed from the nucleotide sequences of the mitochondrial cox1 gene of Echinococcus spp. from all samples based on parameters of weights = 10 and epsilon = 0; nuclear data showed minimal variation and were not included. We computed population diversity indexes (number of haplotypes, haplotype and nucleotide diversity, and mean number of pairwise differences) within the different host groups identified from the phylogenetic analyses with the programme ARLEQUIN 3.5 (Excoffier and Lischer, 2010), which was also employed to calculate the neutrality indexes of Tajima (1989) and Fu’s Fs (Fu, 1997); finally, the degree of gene flow among the three host populations (human, yak and sheep) was estimated using a pairwise fixation index (Fst) as a relative measure of population differentiation, which was determined with the ARLEQUIN package.

**Results**

**Sequencing analysis**

PCR amplification of the cox1 gene was successfully performed for all of the isolated hydatid cysts. According to the cox1 gene nucleotide sequences obtained from the isolated samples, all the human isolates (n = 93), yak isolates (n = 91) and sheep isolates (n = 38) were identified as the E. granulosus s. s. G1 genotype. In addition, the pika isolates were identified as E. multilocularis (n = 16) and E. shiquicus (n = 6). E. granulosus s. s. G1 was clearly the most prevalent species in the animal and human isolates of hydatid cysts. None of the isolates from a given patient/animal occurred in a coinfection.

**Variation in nucleotide and amino acid sequences of E. granulosus s. s.**

A 366-nucleotide consensus cox1 sequence was used to compare and to obtain the haplotypes (Farhadi et al., 2015). Based on the comparison data on cox1 gene sequences, 16 different haplotypes (G1s) were detected, which were designated as EgQH1 to EgQH16 (GenBank accession numbers MG674403-MG674418 (Table 1)). A total of 34 point mutations were found within the haplotypes, consisting of 27 (79.4%) synonymous and 7 (20.6%) non-synonymous substitutions. For the non-synonymous mutations, the number of transition mutations was 5, and the number of transversions was 2; interestingly, the same transition mutation was found at position 56 of EgQH4 (C to T) and EgQH11 (C to T), and the same transition mutation was found at position 111 of ten haplotypes (T to C). Based on the sequence alignment, the EgQH7 haplotype was completely identical to the G1 (AF297617) reference genotype without base mutations; EgQH1, EgQH2, EgQH4, EgQH6, EgQH8, EgQH9, EgQH11, EgQH12, EgQH13, EgQH15 and EgQH16 showed very small nucleotide variations at either one or two positions, and seven haplotypes all included the non-synonymous substitution position 111 (T to C); another five haplotypes, EgQH3, EgQH5, EgQH7, EgQH10 and EgQH14, showed a large number of nucleotide variations (3 to 6) parallel to this, resulting in 1 to 2 different amino acid substitutions. Six nucleotide variations in EgQH10 led to only 2 amino acid substitutions, but two nucleotide variations in EgQH8 also generated 2 substitutions. There was only one transition mutation in EgQH4, EgQH5 and EgQH9, leading to the substitution of an amino acid. Among all 16 haplotypes, EgQH1 was the most common variant and was observed in 53 (21.7%) isolates, comprising 19 human, 21 yak and 13 sheep isolates. EgQH2 was the second most common variant, being found in 40 (16.4%) isolates: 14 human, 15 yak and 11 sheep isolates (Table 1). The next fourteen haplotypes

| Haplotype of isolated Qinghai strains | Host origin (number) | Accession no. | Haplotype in the Fig. 1 |
|--------------------------------------|---------------------|---------------|------------------------|
| EgQH1 Human (19), yak (21), sheep (13) | MG674403 | EGH1 |
| EgQH2 Human (14), yak (15), sheep (11) | MG674404 | EGH3 |
| EgQH3 Human (11), yak (13), sheep (5) | MG674405 | EGH4 |
| EgQH4 Human (9), yak (11), sheep (2) | MG674406 | EGH7 |
| EgQH5 Human (7), yak (9), sheep (2) | MG674407 | EGH8 |
| EgQH6 Human (6), yak (2), sheep (1) | MG674408 | EGH9 |
| EgQH7 Human (5), yak (4), sheep (0) | MG674409 | EGH10 |
| EgQH8 Human (3), yak (3), sheep (0) | MG674410 | EGH11 |
| EgQH9 Human (2), yak (1), sheep (1) | MG674411 | EGH12 |
| EgQH10 Human (3), yak (2), sheep (1) | MG674412 | EGH15 |
| EgQH11 Human (2), yak (2), sheep (0) | MG674413 | EGH16 |
| EgQH12 Human (2), yak (2), sheep (0) | MG674414 | EGH17 |
| EgQH13 Human (2), yak (2), sheep (0) | MG674415 | EGH18 |
| EgQH14 Human (3), yak (1), sheep (1) | MG674416 | EGH19 |
| EgQH15 Human (3), yak (1), sheep (0) | MG674417 | EGH22 |
| EgQH16 Human (2), yak (2), sheep (0) | MG674418 | EGH23 |
| EmQH1 Pika (16) | MG674419 | EMH1 |
| EsQH1 Pika (6) | MG674420 | ESH1 |
(EgQH3–EgQH16) were found in 129 (52.9%) isolates. In addition, while haplotype EgQH1 was the most prevalent variant (22.5%) in the animal isolates, it was also observed (20.4%) in human isolates, and the second most frequent haplotype in the animal isolates was EgQH2.

**Haplotype networks**

In *E. granulosus* s.s., 16 mtDNA haplotypes were found in 222 isolates from the QTPA of China. Nine haplotypes (EgQH1- EgQH 6, EgQH9, EgQH10 and EgQH14) were found in the populations of all three species (Fig. 1). Assuming that the ancestral haplotypes were still present in the recent populations, we constructed a statistical parsimony network to discriminate the genealogical relationships of the haplotypes among the hosts. The network showed a star-like configuration, with the common haplotype (G1) occupied the centre of the network, while G2 (EGH22) and G3 (EGH23) were linked to G1 via one mv1 (median vector) (Fig. 1). The population network presented a high-complexity structure, which included two main sub-groups (EGH4-EGH5 and EGH7-EGH12). However, the majority of haplotypes (EGH1-EGH16) and (EGH18-EGH21) all contained Chinese isolates from different hosts. Additionally, the haplotypes from different hosts were the same in the populations of other countries.

In *E. multilocularis*, 12 mtDNA haplotypes were found in all the isolates, which were also plotted a star-like network with one major haplotype (EMH2) (Fig. 2). The EMH1 haplotype came from three different hosts from two countries, and the EMH2 haplotype came from six different hosts from five countries, while the EMH4-EMH8 haplotypes all came from China but originated from different hosts.

In *E. shiquicus*, 15 mtDNA haplotypes were found in all isolates from China originating from four different hosts. The network was complicated with one star-like (ESH1-7 an ESH9) configuration and two main sub-groups (ESH10-ESH13 and ESH8, ESH11-ESH12, ESH14-ESH15) (Fig. 3).

**Diversity and neutrality indexes**

The diversity indexes for the China QTPA isolates of *E. granulosus* s.s. from three different host species are shown in Table 2. Haplotype diversity was high for all *E. granulosus* s. s. populations within these three host species and was highest in *E. granulosus* s. s. from humans and lowest in those from sheep. In contrast, nucleotide diversity was low for all host species and ranged from 0.006 to 0.008 because of the richness of single nucleotide substitutions.

The neutrality indexes of *E. granulosus* s.s. populations from host species calculated with Tajima’s D and Fu’s Fs tests are also shown in Table 2; the values for these two indexes were all negative, which indicates an excess of rare polymorphic sites and a significant deviation from neutrality.

**Pairwise fixation index values**

The pairwise fixation indexes (Fst) for the *cox1* sequences of populations of *E. granulosus* s.s. from different host species are shown in Supplementary Table 2. Low Fst values were observed for the majority of *E. granulosus* s. s. populations when compared in a pairwise manner with some populations showing negative values (humans/yaks). Since one common haplotype existed predominantly in the three host species, the Fst values between the
populations were very small, ranging from 0.004 to 0.01. These low values implied that the populations were not genetically differentiated among these host species.

Discussion

In this study, the genetic diversity and population structure of Echinococcus spp. in QTPA of China was investigated. Data were obtained via sequencing of the cox1 mitochondrial gene, which had historically been demonstrated to show intraspecific variability and had been used for the study of the population structure of Echinococcus spp. from other parts of the world (Bowles et al., 1992; Nakao et al., 2010; Casulli et al., 2012; Yanagida et al., 2012). Although the new data published in October, 2018, showed that nad5 gene (680 bp) was a highly suitable marker used for the differentiation of E. granulosus s. s. genotypes (Kinkar et al., 2018a). Initially, we used the partial cox1 (366 bp) to distinguish three genotypes (G1−G3) described within E. granulosus s. s. (Bowles et al., 1992). The results presented in this report indicated that G1 was the most dominant distinct species of E. granulosus s. s. in the hydatid cyst samples from humans and animals in regions of the QTPA, China. The epidemiological data of this present study were in line with some previous molecular studies from China that had demonstrated G1 was the most common and dominant species of E. granulosus s. s. in humans, livestock and wild animals in China (Ma et al., 2008, 2015; Yang et al., 2009; Liu et al., 2013; Yan et al., 2013; Zhong et al., 2014; Hu et al., 2015), Turkey (Erdogan et al., 2017), Iran (Farhadi et al., 2015; Arbabi et al., 2017), and Bulgaria (Marinova et al., 2017). Likewise, many epidemiological studies conducted in the majority of regions of the world have also indicated E. granulosus s. s. G1 as the predominant species (Laurimae et al., 2016; Roinioti et al., 2016; Avila et al., 2017; Debiaggi et al., 2017; Ehsan et al., 2017; Thapa et al., 2017).

The cox1 haplotypes of E. granulosus s. s. found in this study did not reveal apparent systematic phylogeographic structuring in the QTPA of China. The parsimony network analysis revealed that the haplotypes exhibited a star-like expansion from a main definitive hosts, which further suggested it might cycle among these host species. The high frequency of the dominant E. granulosus s. s. haplotypes in the QTPA suggested that it may be the ancestral haplotype in the QTPA of China. In another surveillance report of Echinococcus isolates from the QTPA of China, a total of 105 haplotypes (H1-H105) were detected, and 177 variable sites were recorded in 521 samples using the cox1 mtDNA marker gene (Ma et al., 2015). Our results were different from these, our animal isolates collected from the slaughterhouse and the transported animals usually came from the same places, so the results were also different.

In the current study, E. multilocularis (n = 16) and E. shiquicus (n = 6) were present in pika isolates, and no co-infections were observed in individual isolated samples from the Golog Tibetan Autonomous Prefecture. Unfortunately, we were only allowed to conduct sampling in the Golog prefecture at this time, which provided an explanation for why only the same genotype was found here. In our future work, we will conduct sampling and analyses elsewhere, and more samples will be collected and analysed. The identified sequences were highly similar to the referenced E. multilocularis (AB033406) and E. shiquicus sequences (AB208064). E. shiquicus was first identified in the Tibetan fox Vulpes ferrilata (adult stage) and the plateau pika Ochotona curzoniae (larval stage) and was characterized based on its morphological, genetic and ecological features (Xiao et al., 2005, 2006). Thereafter, E. shiquicus infections in dogs were also reported in the eastern QTPA; E. shiquicus was being transferred from its natural host, the Tibetan fox to the domestic dog (Boufana et al., 2013), which will threaten human health, although no cases of E. shiquicus human infection have yet been reported. Fan et al., conducted a genetic diversity analysis of E. shiquicus isolates from the plateau pika in Darlag County of Qinghai Province, and the genetic diversity of the nad1 and cox1 genes was shown to vary by 0.1−1.2% and 0.1−1.0%, respectively, with 6 haplotypes ranging from 4.2

| Host origin | n | S | K | H | Hdh ± s.e. | π ± s.e. | Tajima’s D | Fu’s Fs |
|-------------|---|---|---|---|----------|--------|----------|--------|
| Humans      | 93| 21| 2.74708| 16| 0.9028 ± 0.0140| 0.007506 ± 0.004447| −0.97737| −3.47949|
| Yaks        | 91| 21| 2.60555| 16| 0.8781 ± 0.0161| 0.007127 ± 0.004264| −1.06740| −3.89570|
| Sheep       | 38| 15| 2.35155| 9 | 0.7909 ± 0.0416| 0.006424 ± 0.003982| −1.09838| −1.06127|
| Total       | 222| 21| 2.619420| 16| 0.8731 ± 0.0109| 0.007157 ± 0.004250| −0.67886| −1.84956|

Tunisia (Boufana et al., 2014), Jordan, Iran (Yanagida et al., 2012) and Europe (Casulli et al., 2012). On the other hand, the population genetic structures of E. granulosus s. s. comparing among various endemic areas clarified its worldwide dispersal. The neutrality indexes of Tajima’s D obtained in the current study were negative, which suggested a bias towards the presence of nucleotide variants and was a feature of recent population expansion. Additionally, the neutrality index values Fu’s Fs values was also negative for all populations, which indicated that the incidence of rare haplotypes was lower than expected under neutrality, and the values pointed to bottleneck events and/or purifying selection events that might have occurred in the past (Nakao et al., 2010; Boufana et al., 2014). Furthermore, the extremely low values of the fixation index Fst also supported genetic non-differentiation between the local populations, indicating limited gene flow.

The occurrence of most haplotypes (EgQH1, 3, 4, 7, 12, 13, 16) of E. granulosus s. s. in the QTPA of China was consistent with previously reported results (Ma et al., 2008, 2015). This distribution of the haplotypes indicates the importance of sheep and yak in maintaining potential reservoir infections for humans and definitive hosts, which further suggested it might cycle among these host species. The high frequency of the dominant E. granulosus s. s. haplotypes in the QTPA suggested that it may be the ancestral haplotype in the QTPA of China. In another surveillance report of Echinococcus isolates from the QTPA of China, a total of 105 haplotypes (H1-H105) were detected, and 177 variable sites were recorded in 521 samples using the cox1 mtDNA marker gene (Ma et al., 2015). Our results were different from these, our animal isolates collected from the slaughterhouse and the transported animals usually came from the same places, so the results were also different.

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to 29.6% (Fan et al., 2016); however, only one E. shiquicus haplotype was found in this study. Fan et al. trapped 71 plateau pika samples; we obtained 22 samples and conducted one gene (cox1) sequence genetic diversity analysis. The number of samples and sample collection sites for analysis may need to be expanded in future studies. E. multilocularis represents the second greatest echinococcosis threat to the local people following E. granulosus s. l., and some E. multilocularis haplotypes are also shared by domestic animals (sheep, yaks, and dogs) and humans (Ma et al., 2015); however, only one E. multilocularis haplotype was found in the plateau pika and not in the yaks, sheep and humans in this study, but the threat of E. multilocularis to humans and livestock should not be neglected. E. multilocularis has been detected in domestic animals (dogs and cats) and wild hosts (deer mice, meadow voles and southern red backed voles), not detected in domestic animals (dogs and cats) and wild hosts

**Supplementary material**

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**Author ORCIDs.** Zhang Xueyong, 0000-0001-5076-9211.

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**Conflict of interest.** The authors declare that they have no conflict of interest.

**Ethical standards.** This study was reviewed strictly by the ethics committee of Qinghai Provincial People’s Hospital in accordance with relevant medical ethics regulations, because of the study of hydatid cysts isolated from patients, which was related to the category of medical ethics. Therefore, the study was performed under the supervision of ethics committee, to ensure that the study met the relevant requirements of the ethics committee and maintained the rights of patient. This study was only focus on the hydatid cysts isolated from patients, it was not involved any personal privacy information (individual details, images or videos), and before hydatid cysts removal surgery, the patients were all informed that the operation would be further identified or used for scientific research, they all knew and agreed to publish the hydatid cysts biological information data.

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