

SHORT REPORT

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High prevalence of pathogenic *Leptospira* in alien American mink (*Neovison vison*) in Patagonia

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Abstract

Background: Leptospirosis is an important zoonosis with worldwide distribution caused by pathogenic bacteria of the genus *Leptospira*. The North American mink (*Neovison vison*) has an important role in the environmental contamination with *Leptospira*, as minks live in aquatic environments and are the predators of rodents.

Findings: Blood and kidney samples were obtained from 57 minks in Southern Chile 39° S to 45° S. Pathogenic species of *Leptospira* were detected by PCR on 31/57 minks. To determine the species, we sequenced the 16S ribosomal RNA (rRNA) gene on nine of the positive samples. We predicted two pathogenic species: *Leptospira interrogans* (five samples) and *Leptospira borgpetersenii* (four samples).

Conclusions: This study showed that the American mink presents pathogenic species of *Leptospira* and confirm important environmental contamination of Patagonian rivers and lakes with pathogenic *Leptospira*.

Keywords: Pathogen *Leptospira*; Alien North American mink; Patagonia

Introduction

Leptospirosis is the most widespread zoonosis in the world. Usually, it is transmitted between wild domestic mammals and humans through contaminated water or direct exposure to urine of infected animals. Leptospire can persist in natural environments for long periods of time; while this largely depends on the ability of the bacteria to adapt to a wide range of animals, the pathogen can also survive in water as well (Smith and Zochowski 2011).

Studies on leptospirosis in Chile have reported a high prevalence of infection: 37% in dogs, 88.7% in cattle, 24.9% in sheep, 7.1% in horses, and 69.9% in swine; see review by (Zunino and Pizarro 2007). However, only a few studies have investigated the prevalence of *Leptospira* spp. in wild animals. For example, only the prevalence of *Leptospira* spp. in wild rodents is currently known (47.2%); see reviews by (Zamora and Riedemann 1999; and Zunino and Pizarro 2007).

The North American mink (*Neovison vison*) is a semi-aquatic mustelid that was introduced in Chile in the 1930s (Jaksic et al. 2002). Currently, the species has outgrowth considerably in number and distribution. The

mink is now widely distributed throughout the riverine and marine habitats of the Patagonian lacustrine in both Argentina and the south of Chile (from 38° S latitude to Tierra del Fuego Island and adjacent archipelagos 55° S) (Medina 1997; Fasola et al. 2011). Mustelids are among the animals considered susceptible to infection with *Leptospira*; importantly, rodents are not only the primary reservoir of *Leptospira*, but also are an important part of the diet of minks. Additionally, the fact that minks often come into direct or indirect contact with different domestic animals suggests that inter-animal disease transmission events or potential human-transmission events are possible (Ullmann and Langoni 2011). While the role of minks as potential reservoir of *Leptospira* has not been determined yet, this role cannot be discarded as many emerging human, domestic animal, and wildlife diseases are usually maintained in specific reservoirs, many of which are rarely identified see reviews by (Adler and de la Peña. 2010; and by Smith and Zochowski 2011, Lau et al. 2012). Furthermore, *Leptospira* transmission is very complex, in fact, this is a pathogen of multiple hosts, which often resides in one or more epidemiologically connected populations (Haydon et al. 2012). The complexity of *Leptospira* transmission along with the abundance of mink populations indicates that minks may be an important link in the

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ecology of leptospires, especially due to their wide distributions, semiaquatic live, and diet (Medina 1997; Millán et al. 2009; Moinet et al. 2010; Sepúlveda et al. 2011). In this study, we determined and characterized the presence of pathogenic *Leptospira* species in North American minks in Southern Chile.

Materials and methods

Sample collection

During September 2011 to March 2012 we trapped 57 minks (39 males and 18 females) using leg hold soft catch traps and cage traps. The animals were captured in ten study areas within three Southern Chilean districts (Los Rios, Los Lagos, and Aysén), between 39° S to 45° S (Figure 1). These areas support a large diversity of habitats and species, including domestic and native animals, such as cows (*Bos taurus*), horses (*Equus ferus caballus*), sheep (*Ovis aries*), dogs (*Canis lupus familiaris*), cats (*Felis silvestris catus*), foxes (e.g. *Lycalopex* spp.), and otters (*Lontra provocax*). The coordinates of the traps were determined with a global positioning system (GPS). Once

under anesthesia, with ketamine-dexmedetomidine in a dose of 10–0.025 mg/kg IM respectively, two blood samples (3–5 ml each) were collected by jugular venipuncture (one blood sample was collected using tubes with EDTA and the other sample was collected in tubes with cell buffer lysis). The minks were then euthanized with Tiopental® (Biosano S.A., Santiago, Chile) doses for postmortem kidney tissue collection (collected in alcohol). All animal trapping and handling were performed according with ethical protocols of the Committee of Ethic of the Universidad Andrés Bello and the National Commission for Science and Technology. Samples of blood and kidney collected in the field were frozen and transported to the laboratory for further analysis.

Laboratory analysis

Total DNA from blood and kidney was extracted with the QIAamp DNA Mini Kit (Qiagen®, Germany). On three samples per individual (two blood and one kidney), we used PCR to screen for *Leptospira* spp. as previously described by Lester and LeFebvre, 2003. To determine

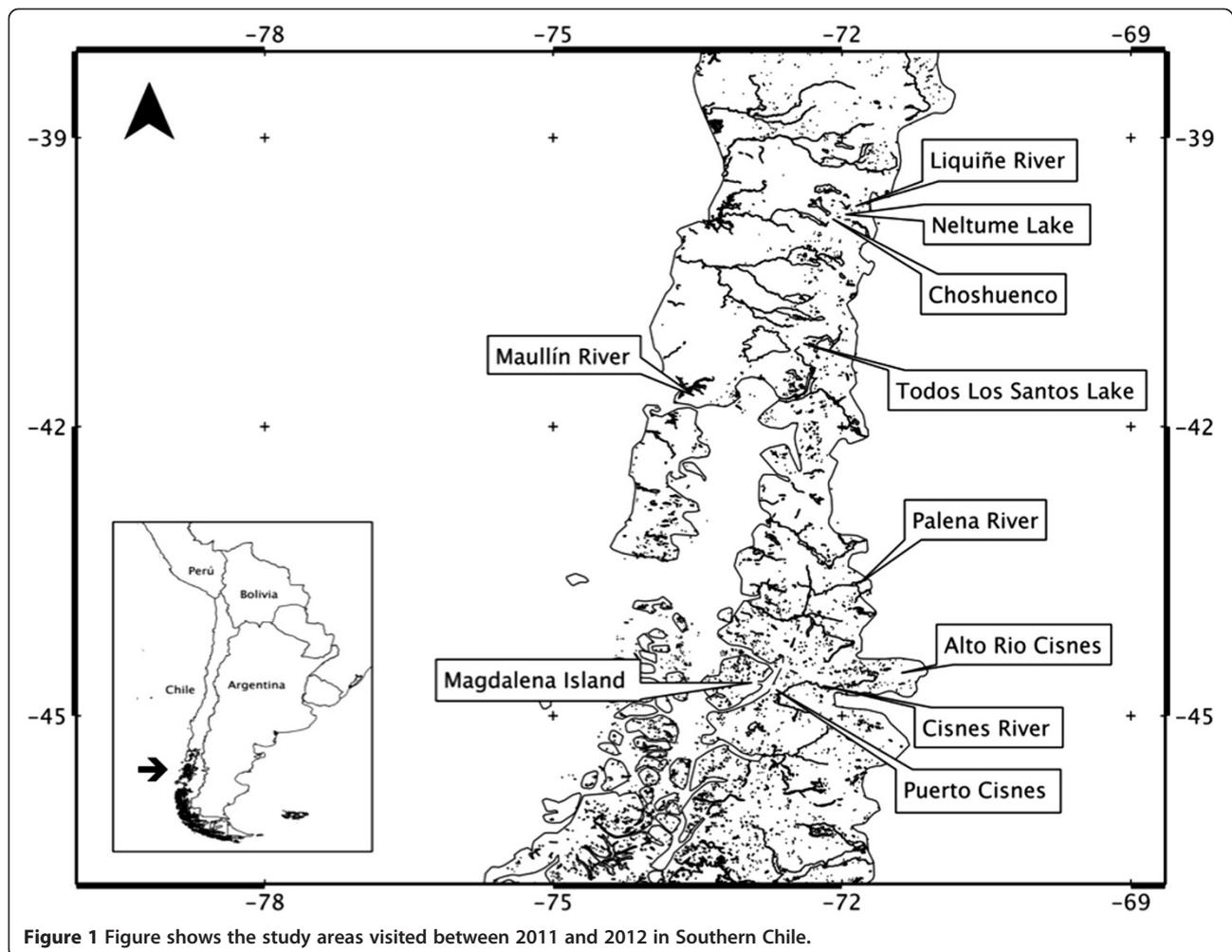


Figure 1 Figure shows the study areas visited between 2011 and 2012 in Southern Chile.

the species of *Leptospira* present in the positive samples, the amplified DNA (571 bp) of nine representative samples were sequenced at GenYtec (Santiago, Chile) using the ABI PRISM 310. To predict the species, sequences were compared with the GenBank database using the BLAST algorithm (<http://www.ncbi.nlm.nih.gov/BLAST>).

Findings

This study reports for the first time the prevalence of pathogenic *Leptospira* species in the American mink in the Chilean Patagonia. Among all the mink tested, 31 individuals corresponding to 55.6% (95% confidence interval = 24, 8–86, 4) (21 males and 10 females) were positive for the PCR detection of pathogenic *Leptospira* spp. (Table 1). Seven animals tested positive for both, blood and kidney samples; eighteen of them tested positive for blood and only six for kidney samples. This suggests that more than one sample per individual facilitates the detection of *Leptospira* by PCR. In eight of the ten locations where samples were collected, we obtained positive results; specifically, only the location of Choscuenco and Rio Cisnes were negative (Table 1 and Figure 1). The location with the higher number of positive samples ($n = 8$) was Isla Magdalena; however, we collected samples from 15 individuals in this location.

Nine positive samples from different minks and from different locations were selected for sequencing of the 571 bp amplicon of the 16S ribosomal RNA (rRNA) (Table 1). The BLAST algorithm was used to (i) validate that the amplified PCR products correspond with *Leptospira* species and to (ii) predict the putative pathogenic species. We identified sequences closely related to two of the *Leptospira* pathogenic species. *Leptospira interrogans*

was putatively predicted in five samples from three locations (i.e., Palena, Alto Rio Cisnes, and Islas Magdalena) and samples closely related to *Leptospira borgpetersenii* in four samples from four locations (i.e., Liquiñe, Neltume, Todos los Santos, and Alto Rio Cisnes) (Table 1). Our matches showed approximately 97% of nucleotide identity with the *Leptospira* species mentioned above. However, these findings need to be validated through the analysis of the complete 16 S rRNA gene.

We report for the first time evidence of pathogenic *Leptospira* in alien North American mink in South America. In this study, we used PCR to detect the pathogen; previous studies have used traditional detection methods (e.g., microscopic agglutination test [MAT]); while these methods are not easily compared, here we discuss the prevalence previously identified by other authors in comparison with our findings, regarding of the method used. Importantly, we identified a higher prevalence (55.6%, with 95% confidence interval = 24, 8–86, 4) compared with previous reports of 23.5% of *Leptospira* in wild animals in Spain (Millán et al., 2009), which used the MAT to detect the presence of *Leptospira*. Another study reported by Moinet et al. (2010) identified a prevalence of 86% in wild American mink in France MAT detection. The same study detected by PCR a renal carriage of 26% on free-ranging American minks. The fact that we detected pathogenic species of *Leptospira* is relevant to both human and wild species health. Prevalence was high and transversal to almost all study sites, which are characterized by a mixture of natural forest and human activities. A previous study by Ghneim et al. (2007) showed that dogs in rural areas were more likely to have been in contact with contaminated water

Table 1 Number of positive samples and predicted *Leptospira* species in each study location

Location	Environment	Sample size	Number of positive minks ^a (sequenced samples)	Putative species of <i>Leptospira</i> ^b (number of samples)
Liquiñe	Riverine	3	2 (1)	<i>L. borgpetersenii</i> (1)
Neltume	Lacustrine	5	3 (1)	<i>L. borgpetersenii</i> (1)
Choscuenco	Lacustrine	3	0	-
Todos los Santos	Lacustrine	8	5 (1)	<i>L. borgpetersenii</i> (1)
Maullín	Palustre	6	5	Not sequenced
Palena	Riverine	1	1 (1)	<i>L. interrogans</i> (1)
Alto Rio Cisnes	Riverine	5	4 (3)	<i>L. interrogans</i> (2), <i>L. borgpetersenii</i> (1)
Isla Magdalena	Marine	15	8 (2)	<i>L. interrogans</i> (2)
Puerto Cisnes	Marine	6	3	Not sequenced
Rio Cisnes	Riverine	5	0	-
Total		57	31	

^aPCR was conducted in three samples per individual (two blood and one kidney sample).

^b*Leptospira* species were predicted according to the amplification of a 571 fragment of the 16S RNA gene. The best hit of BLAST was used to assign the putative species. A complete sequence of the 16S rRNA gene would validate these findings.

with leptospires, as compared with dogs in urban areas. This may indicate that the complexity of animal species, along with environmental factors (e.g., rainfall and standing water) in rural areas, could facilitate the contact with the pathogen. Minks likely become infected after preying on rodents. In fact, studies carried out in the same region as our study demonstrate that mink preys heavily on rodents (Medina-Vogel et al. 2013), and that rodents may become infected by contact with garbage and contaminated water (Smith and Rochowski 2011). In the south of Chile, rodents present high rates of *Leptospira* infection (Zunino and Pizarro 2007). In fact Muñoz-Zanzi et al. (2014) detected, by PCR, a 19.7% *Leptospira* prevalence in rodents associated with agricultural fields, 25.9% associated with rural villages, 21.1% associated with wild rodents, and 12.3% associated with slums.

In this study, we found *L. interrogans* in Palena, Alto Rio Cisnes, and Isla Magdalena, which is a *Leptospira* species largely associated to rodents (Himsworth et al. 2013). In addition, we found *L. borgpetersenii* in Liquiñe, Neltume, Lake Todos los Santos, and Alto Rio Cisnes, which is a *Leptospira* species associated with cattle farming. Importantly, in locations where we found *L. borgpetersenii*, livestock production systems are common. Thus, it is tempting to speculate that minks could be infected by preying on wild native and alien rodents in locations where livestock is rare, as well by contact with garbage and contaminated water from horse, cattle, sheep, and pig urine in locations where livestock is common.

In 13 individuals, samples from kidneys were positive; this not only demonstrates the presence of chronic disease in minks, but also that they present renal carrier status, which is the central point of the epidemiology of leptospirosis. Consequently, minks in the south of Chile may excrete leptospires through the urine to the environment (Ullmann and Langoni 2011). These initial data indicates that American mink can serve as hosts for leptospirosis, thus serving as a tool for measuring environmental contamination with this pathogen. This study represents a first insight of the mink's role in the epidemiology of *Leptospira* in the south of Chile. This initial evidence suggests that mink abundance may pose a threat to human aquatic activities (e.g., canoeing, aquaculture, fisheries) and may as well affect the survival of aquatic species of conservation concern, such as the southern river otter (*L. provocax*) (Gaydos et al. 2007). Further investigation in order to identify source of infections, wild and domestic mammalian species involved, and the mechanism of transmission are necessary to provide evidence of the role of minks in the transmission of this important zoonotic pathogen.

Competing interests

The authors declare that they have no competing interests.

Authors' contributions

LS participated in the sequence alignment and have been involved in drafting the manuscript. LL participated in the sequence alignment and drafted the manuscript. CN carried out the molecular genetic studies and participated in the sequence alignment. MB carried out the acquisition of data and analyzed and wrote the manuscript. GM conceived of the study and participated in its design and coordination and have been involved in drafting the manuscript. All authors read and approved the final manuscript.

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