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# Zoonotic Parasites of Bobcats around Human Landscapes

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# Zoonotic Parasites of Bobcats around Human Landscapes

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**We analyzed *Lynx rufus* fecal parasites from California and Colorado, hypothesizing that bobcats shed zoonotic parasites around human landscapes. *Giardia duodenalis*, *Cryptosporidium*, *Ancylostoma*, *Uncinaria*, and *Toxocara cati* were shed. *Toxoplasma gondii* serology demonstrated exposure. *Giardia* and *Cryptosporidium* shedding increased near large human populations. Genotyped *Giardia* may indicate indirect transmission with humans.**

Pathogen spillover is a major disease source for humans and wildlife (8, 10, 16). Studies have suggested that wild felids might contribute to spillover of a variety of high-profile pathogens to humans (3, 5, 22, 24, 27, 39). However, owing to their secretive nature and the effort required to sample, few studies can draw significant inferences about wild felid pathogen exposure and shedding. In this study, we screened a large number of bobcat (*Lynx rufus*) fecal samples for zoonotic enteric parasites and evaluated determinants of shedding and exposure in relation to human-occupied landscapes. We hypothesized that bobcats become exposed and shed zoonotic parasites around human-occupied landscapes.

Bobcats are widespread in North America and persist across a continuum of natural to heavily urbanized habitats, often existing in sympatry with humans, domestic animals, and urban-adapted wildlife along urban boundaries (9, 28, 31). Fecal zoonotic pathogens shed by felids, and considered most likely to be detected in this study, include *Toxoplasma gondii*, *Cryptosporidium*, *Giardia*, roundworms, and hookworms. In certain circumstances, these cosmopolitan pathogens infect and cause disease to humans and cause outbreaks, primarily associated with environmental contamination (7, 13, 29, 34, 41). Comparably, domestic animals, urban wildlife, and, in some cases, humans are also permissive hosts for such common parasites (11, 17, 32, 40). Thus, along urban boundaries, abundant sympatric populations of permissive hosts may amplify the pathogen circulation and force of infection (18, 32).

We screened samples from three sites in California (Ventura County [VC]) and Colorado (Front Range [FR] and Western Slope [WS]) that vary in the degree of urbanization (Fig. 1). Bobcat sex, age (young, <2 years; adult, ≥2 years), and location were recorded at the time of capture (4). Fecal ( $n = 141$ ) and matching blood ( $n = 73$ ) samples were collected in the field, refrigerated, and then frozen until analyses. From a subset of WS animals (ca. 60%), fecal samples were collected from hunter-killed animals and thoracic fluid was collected instead of serum.

Fecal flotation was used for detection of parasite eggs, cysts, and oocysts, and immunofluorescence assays were used for *Giardia* cysts and *Cryptosporidium* oocysts (36). PCR assays for *Giardia* sp., *Cryptosporidium* sp., and *T. gondii* were performed on fecal dilutions from fluorescence assays (19, 26, 30, 33, 37). Positive amplicons were sequenced in forward and reverse directions (ABI3100 genetic analyzer; Applied Biosystems). Sequences were compared with sequences from GenBank by BLAST, and new se-

quences were placed in GenBank. Additionally, IgM and IgG antibodies against *T. gondii* were detected using indirect enzyme-linked immunosorbent assays (ELISAs) on sera (20, 42) and modified agglutination tests on thoracic fluid (2).

Statistical analyses focused on the most-abundant pathogens and those with greatest zoonotic potential (*T. gondii*, *Cryptosporidium*, and *Giardia*). Estimates of prevalence (and 95% confidence intervals [CIs]) were determined using maximum likelihood. The effects of host (sex and age) and geographic (study site) determinants of shedding/serostatus were evaluated using individual odds ratios (based on conditional maximum likelihood estimation). Analyses were conducted using R (v.12.14.1; [www.r-project.org](http://www.r-project.org)) with the stats and epitools packages.

Fecal flotation identified a variety of parasites (Table 1). In support of our hypothesis, we found evidence of zoonotic parasite shedding (*Giardia duodenalis*, *Cryptosporidium* sp., *Ancylostoma* sp., *Uncinaria* sp., *Toxascaris leonina*, and *Toxocara cati*) or exposure (*T. gondii*) across all sites (Table 1). There are a few studies of enteric parasites of bobcats, but those that exist collectively report similar parasite communities (e.g., see references 12, 23, 38, and 43). The prevalences (1 to 13.5%) of the parasites examined here are also broadly comparable to those of adjacent domestic cat populations (6, 14), suggesting common exposure among the felids.

No fecal samples were positive for *T. gondii* oocysts or DNA. However, 17.8% (95% CI, 10.2 to 27.6) of bobcats were seropositive by IgG only, demonstrating past exposure and suggesting shedding patterns similar to domestic cats (25). Seroprevalence was not predicted by sex, but adults were 6.1 (95% CI, 1.2 to 61.8) times more likely to be seropositive than young animals. Bobcats from VC and FR, the more-urban sites, were 5.8 (1.2 to 33.2) and 6.3 (0.7 to 51.2) times more likely to be seropositive than bobcats from WS, respectively, but this effect was partially driven by more young animals being sampled from WS. In general, *T. gondii* exposure appears greater among wild felid populations than among domestic cats in the United States, suggesting that wild felids may

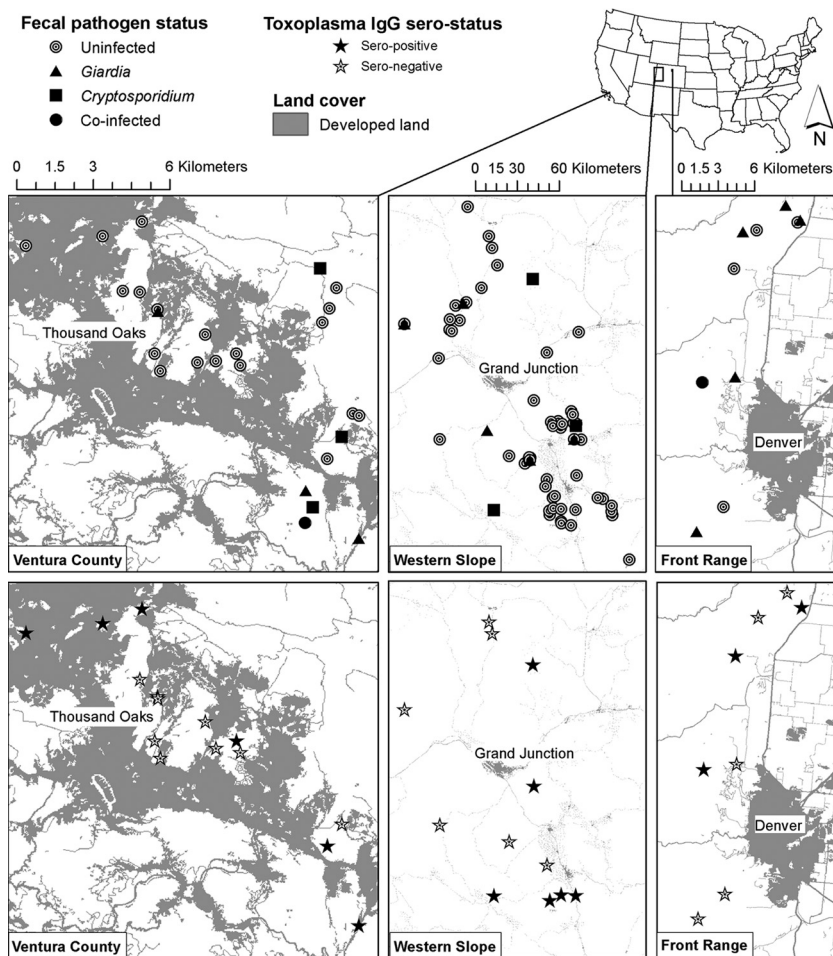
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**FIG 1** Capture locations of bobcats and associated developed landscape features (primarily urban areas and major highways) across study locations in California (Ventura County [VC]) and Colorado (Front Range [FR] and Western Slope [WS]). VC is a highly urbanized landscape within the Santa Monica Mountains, north of Los Angeles, CA. FR is an urbanized area along the Front Range of the Rocky Mountains, northwest of the Denver metropolitan area and immediately adjacent to Boulder. WS is a primarily rural and exurban region along the Western Slope of the Rocky Mountains around Montrose and Grand Junction. Upper panels illustrate detection of *Giardia*, *Cryptosporidium*, and their cooccurrence in feces. Lower panels illustrate *T. gondii* IgG serostatuses of bobcats. *T. gondii* IgM antibodies, oocysts, and DNA (shedding) in feces were not detected in any of the bobcats.

**TABLE 1** Enteric parasites identified through microscopic examination of fecal flotation filtrate across study sites

Phylum	Group	Parasite(s)	No. of samples with indicated parasite in each location (n) <sup>a</sup>		
			VC (33)	WS (103)	FR (10)
<i>Nematoda</i>	Hookworms <sup>b</sup>	<i>Ancylostoma</i> sp. and <i>Uncinaria</i> sp.	2	2	1
	Roundworms <sup>b</sup>	<i>Toxascaris leonina</i>	1	3	5
		<i>Toxocara cati</i>	0	5	0
	Whipworms	<i>Trichuris</i> sp.	2	0	0
	Stomach worms	<i>Physaloptera felis</i>	0	9	0
	Lungworms	<i>Eucoleus</i> sp.	0	1	1
	Unidentified	Soil nematodes	3	0	0
<i>Platyhelminthes</i>	Flatworms	Fluke eggs	1	0	0
<i>Apicomplexa</i>	Coccidias	<i>Eimeria</i> sp.	1	0	0
		<i>Isospora</i> sp.	0	7	0
		<i>Cystoisospora felis</i>	0	2	0

<sup>a</sup> VC, Ventura County; WS, Western Slope; FR, Front Range. Numbers in parentheses indicate numbers of fecal samples examined.

<sup>b</sup> Zoonotic potential.

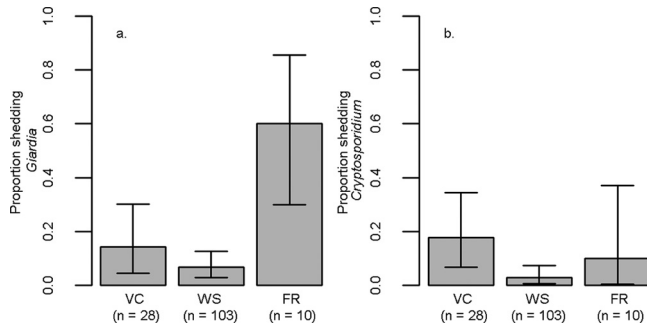


FIG 2 Proportions (maximum likelihood estimates  $\pm$  95% CIs) of bobcats shedding *Giardia* (a) and *Cryptosporidium* (b) across the study locations Ventura County (VC), Western Slope (WS), and Front Range (FR).

be a source for domestic cats (4), although *T. gondii* can persist in regions in which wild felids are absent (1). The force of exposure to *T. gondii* for domestic cats, humans, and urban wildlife might be enhanced directly adjacent to the urban-wildland interface, where there is sympatry among wild and domestic felids and amplification of *T. gondii* through the food chain (21). For example, sympatry between pumas and domestic cats might have been the driver of the 1995 human toxoplasmosis outbreak in British Columbia (3).

*Cryptosporidium parvum* and *Giardia duodenalis* account for the majority of waterborne disease outbreaks in humans, with approximately 2 million and 750,000 annual cases in the United States, respectively (11, 15, 17, 35). Both pathogens have several species and subtypes, with some host specific and others zoonotic. Little is known about the shedding of these pathogens by bobcats. We found shedding across all sites (Fig. 2). Neither *Giardia* nor *Cryptosporidium* was predicted by sex or age. The prevalence of *Giardia* shedding was greater at FR than at either VC or WS, and there was a trend for greater shedding at VC than WS (Table 2 and Fig. 2a). Similarly, shedding of *Cryptosporidium* oocysts was greater at VC than WS, and a trend for greater shedding at FR than WS was noted (Table 2 and Fig. 2b). Overall, results indicate a greater probability of shedding of *Giardia* and *Cryptosporidium* at VC and FR, which are situated near large human population centers, than at the more-remote WS site.

We typed four *Giardia* isolates as *G. duodenalis* assemblage A1, which is commonly found in humans and appears maintained in multispecies complexes, including domestic animals and wildlife

(11). Though *Cryptosporidium* shedding patterns were similar to those of *Giardia*, we were unable to determine if positive samples were felid, human, or other animal in origin. A direct fluorescence assay is the gold standard for generic *Giardia* and *Cryptosporidium* identification, and our inability to identify more isolates to the species level was likely due to PCR inhibitors in feces or possibly due to the volume of sample available. Notably, a cluster of bobcats shedding *Giardia* and *Cryptosporidium* at VC corresponds to the highly urbanized Malibu creek watershed (Fig. 1). Given shedding patterns among sites and within VC as well as genetic identities, our results may suggest anthroponotic transmission of *Giardia* to bobcats, likely through contaminated water or other environmental sources.

An alternative interpretation for our results is the impact of climatic and ecological drivers among sites, particularly in comparing California and Colorado sites. VC is characterized by a Mediterranean climate and associated vegetative communities, whereas the Colorado study sites are cooler, semiarid, and dominated by coniferous woodlands. If climate- and habitat-related factors are primary determinants of enteric pathogens across these sites, we might expect Colorado sites (WS and FR) to be more similar to each other than to VC. However, the similarity of the most-urbanized sites (VC and FR) in the prevalence of pathogen shedding and the differences between these sites and the least-urbanized site (WS) suggest other factors to be important determinants of enteric pathogen shedding and exposure.

Our results are supportive of abundant populations of permissive hosts around urban areas leading to amplification of pathogen circulation and force of infection and exposure (18). We found support for our hypothesis that bobcats become exposed to and shed zoonotic parasites in fecal material around human-occupied landscapes. Future research evaluating parasite exposure and shedding, along with landscape genetic structure along urban-wildland gradients, among bobcats, other wildlife (particularly *Puma concolor*), domestic animals, and humans will contribute a better empirical understanding of multihost parasite dynamics, pathogen spillover, and human exposure risk.

**Nucleotide sequence accession numbers.** New sequences were added to GenBank under accession numbers JQ688299 to JQ688300 for *Giardia* and JQ740600 to JQ740605 for *Cryptosporidium*.

TABLE 2 Individual odds ratios and 95% confidence intervals for predictors of *Giardia* sp. and *Cryptosporidium* sp. shedding in bobcats<sup>a</sup>

Compared parameters	Odds ratios and 95% CIs for predictors of shedding of:					
	<i>Giardia</i> sp.			<i>Cryptosporidium</i> sp.		
	Odds ratio	95% CI		Odds ratio	95% CI	
		Lower	Upper		Lower	Upper
Male vs female	1.272	0.385	4.919	0.389	0.073	1.909
Adult vs young	1.600	0.456	7.162	1.685	0.303	17.303
VC vs WS	2.269	0.450	9.815	7.094	1.277	48.909
FR vs VC	8.320	1.326	63.870	0.141	0.020	0.783
FR vs WS	19.385	3.673	117.946	0.275	0.020	15.733

<sup>a</sup> Based on conditional maximum likelihood estimations. The predictors include sex (male or female), age (young or adult), and location (Ventura County [VC], Western Slope [WS], or Front Range [FR]).

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Any use of trade, product, or firm names is for descriptive purposes only and does not imply an endorsement by the U.S. Government.

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