spp. isolates in US food animals (10). Given these linkages, the transfer of an MDR plasmid from *Salmonella* spp. to *Y. pestis* seems possible. However, we emphasize that to date no evidence supports this type of event.

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David M. Wagner,
Janelle Runberg, Amy J. Vogler,
Judy Lee, Elizabeth Driebe,
Lance B. Price,
David M. Engelthaler,
W. Florian Fricke, Jacques Ravel,
and Paul Keim

Author affiliations: Northern Arizona University, Flagstaff, Arizona, USA (D.M. Wagner, J. Runberg, A.J. Vogler, J. Lee, P. Keim); Translational Genomics Research Institute, Flagstaff (E. Driebe, L.B. Price, D.M. Engelthaler, P. Keim); and University of Maryland School of Medicine, Baltimore, Maryland, USA (W.F. Fricke, J. Ravel)

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References


Address for correspondence: David M. Wagner, Center for Microbial Genetics and Genomics, Northern Arizona University, Flagstaff, AZ 86011-4073, USA; email: dave.wagner@nau.edu

**Triatoma infestans**

**Bugs in Southern Patagonia, Argentina**

To the Editor: *Triatoma infestans* bugs, the main vector of Chagas disease, historically occupied a large area from northeastern Brazil to Chubut Province in Patagonia, Argentina (1). Large-scale insecticide spraying during the 1980s and 1990s reduced its geographic range and abundance and interrupted transmission of *Trypanosoma cruzi*, mainly in Uruguay, Chile, and Brazil (2). However, *T. infestans* and transmission of *T. cruzi* persist in the Gran Chaco, a large ecoregion in Argentina, Bolivia, and Paraguay (3).

Chubut Province has historically been an area with no risk for vector-mediated transmission of *T. cruzi*, only its extreme northern region was categorized as having a low transmission risk (4,5). However, increased immigration from disease-endemic rural areas in Argentina and Bolivia into Chubut has raised concerns about accidental introduction of *T. infestans* in travelers’ luggage (1) and establishment of a transmission cycle.

In January 2007, we conducted a province-wide survey of 21 villages in Chubut Province previously infested with *T. infestans* bugs by using 0.2% tetramethrin as a dislodgent agent (1 person-hour/house); no *T. infestans* bugs were detected (online Appendix Figure, www.cdc.gov/EID/content/16/4/887-appF.htm). Only *T. patagonica* bugs were found in 11% of peridomestic structures, and none were infected with *T. cruzi*. In June 2007, a *T. infestans*-like bug was found in a primary healthcare center in Comodoro Rivadavia (45°51’S, 67°28’W), a city in southern Chubut Province (online Appendix Figure). Healthcare center staff reported visits by immigrants from Bolivia a few days before this finding and suspected them to be the source. The bug was identifi-
fied morphologically as a *T. infestans* female and it laid 6 eggs. PCR amplification of kinetoplast DNA showed that it was not infected by *T. cruzi*.

DNA sequence analysis is useful for investigating evolutionary history and population structure within Triatominae (6). *T. infestans* bugs from Bolivia and Argentina showed genetic differences for nuclear (7) and mitochondrial markers (6), including mitochondrial cytochrome oxidase I (mtCOI) (8). We used our mtCOI haplotype database, which includes published (8) and new domestic, peridomestic, and sylvatic *T. infestans* from 65 locations in 13 provinces in Argentina (n = 346) and 3 departments in Bolivia (n = 144), to analyze the mtCOI sequence of the bug found in southern Patagonia and determine if it could be assigned to a known haplotype from Bolivia or Argentina. We investigated phylogenetic relationships with other haplotypes by using neighbor-joining and Bayesian approaches.

Our mtCOI database included 53 haplotypes: 42 were found in Argentina, 9 in Bolivia, and 2 in both countries (Figure). The bug from southern Patagonia had haplotype *x*, which has been found in only 3 western or southern provinces in Argentina (San Juan, San Luis, and Rio Negro) (8; online Appendix Figure).

Results of phylogenetic analyses were congruent (Figure). The neighbor-joining tree showed that haplotype *x* formed a cluster with haplotype *h* (Argentina) and haplotypes from Bolivia clustered in 3 other groups: 1) two groups with bootstrap values >70% (one with haplotypes at, n, c, and 33 haplotypes from Argentina, and the other with haplotypes ab, ac, ad, ae, ap, and az); and 2) one group with a bootstrap value of 68% (haplotypes ax and aa). The Bayesian tree showed that haplotypes from Bolivia were arranged in 2 well-supported clades (posterior probabilities ≥83%) and that haplotype *x* was not included within any of them. Thus, haplotype *x* of the bug from southern Patagonia was found only in Argentina and was not closely related to haplotypes from Bolivia.

We investigated the geographic origin of non-native putative attendees of the healthcare center in San Cayetano. These persons were immigrants from Bolivia and from northern (Salta and Jujuy), western (Mendoza, San Juan, and San Luis), and southern Argentina (Rio Negro), i.e., from the 3 putative sources of the bug. These immigrants typically pay extended visits to their home towns at least once a year and transport luggage in which the bug could have traveled. In 2006, San Juan had the highest levels of domestic and

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**Figure.** Phylogenetic relationships between mitochondrial cytochrome oxidase I gene haplotypes of *Triatoma infestans* from Argentina and Bolivia. The neighbor-joining tree was constructed by using MEGA 4.1 (www.megasoftware.net) and bootstrap values (based on 1,000 replications) >50% are shown above the branches. A Bayesian maximum clade credibility tree was similar, and clade posterior probabilities >50% are shown below the branches of the neighbor-joining tree. MRBAYES 3.1 (http://mrbayes.csit.fsu.edu) default priors were assumed and run for 4 million generations. Convergence of the Markov chain Monte Carlo analysis was investigated with the SD of split frequencies and diagnostics implemented in AWTY (http://ceb.csit.fsu.edu/awty). The model of evolution (Hasegawa-Kishino-Yano + invariable sites + Γ [HKY + I + Γ]) was chosen with Mrmodeltest 2.3 (www.abc.se/~nylander). Because MEGA 4.1 does not support HKY, the more inclusive Tamura-Nei method (www.megasoftware.net/WebHelp/part_iv___evolutionary_analysis/computing_evolutionary_distances/distance_models/nucleotide_substitution_models/hc_tamura_nei_distance.htm) was used for the neighbor-joining analysis. Haplotypes *al*, *an*, *ao*, *aq*, *at*, *au*, *ax*, *az*, *aaa*, and *aab* are reported. DNA sequences are available in GenBank (accession nos. EF451005-EF451041, FJ439768, FJ811845–8, and GQ 478993–GQ 479005). *Two provinces in Argentina; †Tarja, Bolivia; ‡10 provinces in Argentina. Scale bar indicates nucleotide substitutions per site.
peridomestic infestation with *T. infestans* (35% and 21%, respectively), including urban infestation (9). Mendoza (not in our database) had considerable domestic and peridomestic infestations (both 7%), and San Luis (0.5% and 5.3%, respectively) and Rio Negro (both <0.1%) had low infestations in 2001 (4) and thereafter (C. Spellmann, unpub. data). Bolivia, Salta, and Jujuy are excluded as potential sources of the bug because haplotypes closely related to haplotype x were not found in these places. Active dispersal from a local source can be ruled out because there is no precedent for *T. infestans* in Comodoro Rivadavia, and the mean temperature in June (8°C) is below the known threshold for flight initiation (23°C) (10).

Our results show that molecular phylogenetics can identify passive transport of insects into areas where a disease is not endemic and rule out putative sources supported only by circumstantial evidence. Our findings reinforce the need for sustained and coordinated vector surveillance and control at a regional level (3).

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Romina V. Piccinali, Delmi M. Canale, Alejandra E. Sandoval, Marta V. Cardinal, Oscar Jensen, Uriel Kitron, and Ricardo E. Gürtler

Author affiliations: Universidad de Buenos Aires, Buenos Aires, Argentina (R.V. Piccinali, M.V. Cardinal, R.E. Gürtler); Coordinación Nacional de Control de Vectores, Córdoba, Argentina (D.M. Canale); Secretaría de Salud de Chubut, Chubut, Argentina (A.E. Sandoval, O. Jensen); and Emory University, Atlanta, Georgia, USA (U. Kitron)

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References


Address for correspondence: Romina V. Piccinali, Laboratorio de Eco-Epidemiología, Departamento de Ecología, Genética y Evolución, Facultad de Ciencias Exactas y Naturales, Universidad de Buenos Aires, Pabellón 2, Ciudad Universitaria, Int. Güiraldes 2160, C1428EGA, Buenos Aires, Argentina; email: rpicei@ege.fcen.uba.ar

Serologic Survey of Hantavirus Infection, Brazilian Amazon

To the Editor: Since the etiology of hantavirus cardiopulmonary syndrome (HCPS) was recognized in 1993 in the United States (1), various hantaviruses have been associated with the syndrome in South America (2,3). Depending on the viral genotype involved, hantavirus infection can take a wide variety of forms, from asymptomatic or oligosymptomatic to the classic clinical form (4,5).

The first cases of HCPS in Brazil were reported in the state of São Paulo in 1993 (6,7). In 2000, an outbreak of HCPS was reported in the municipality of Anajatuba in the state of Maranhão in the Maranhão western