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Zoonotic Pathogens among White-Tailed Deer, Northern Mexico, 2004–2009

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To the Editor: Intense wildlife management for hunting affects risks associated with zoonotic pathogens ([1](#)). White-tailed deer (*Odocoileus virginianus*) are increasingly managed by fencing, feeding, watering, and translocation to increase incomes from hunting in northern Mexico ([2](#)). These deer also play a major role in dissemination and reintroduction of pathogens and vectors from Mexico into the United States ([3,4](#)). White-tailed deer are suitable reservoir hosts for *Mycobacterium bovis* ([1](#)), and an *M. bovis*-positive white-tailed deer was recently found in Tamaulipas in northeastern Mexico ([2](#)). Brucellosis is widespread in many animal hosts in Latin America ([5](#)) and thus of interest in white-tailed deer. Another major zoonosis, sometimes linked to raw deer meat consumption, is hepatitis E, which is caused by genotypes of hepatitis E virus (HEV) ([6](#)). HEV is increasingly prevalent in red deer (*Cervus elaphus*) ([7](#)), but its prevalence in white-tailed deer is unknown.

The objective of this study was to determine the prevalence of zoonotic pathogens in white-tailed deer in northern Mexico. This study was conducted under a scientific collecting permit issued by the Mexican Division of Animal and Wildlife Health and on 8 ranches in 3 states in northern Mexico ($\approx 26\text{--}28^\circ\text{N}$, $99\text{--}100^\circ\text{W}$).

Serum samples ($n = 347$) were collected during 2004–2009 in a cross-sectional survey for antibodies against HEV, *Brucella* spp., and mycobacteria. Deer were opportunistically sampled during live-capture operations as described by Cantú et al. ([8](#)). Bleeding was performed by using jugular venipuncture and vacuum tubes without anticoagulant. Samples were allowed to clot and centrifuged to collect serum that was stored at -20°C .

Serum samples were tested for IgG against HEV by ELISA as described ([7](#)). Serum samples were also tested for antibodies against *Brucella* spp. by using a commercial ELISA (Ingezim Brucella Compac 2.0 Ingenasa, Madrid, Spain), according to the manufacturer's instructions. Detection of antibodies cross-reacting with 2 widely used mycobacterial antigens, bovine purified protein derivative (PPD) and paratuberculosis protoplasmatic antigen 3 (PPA3), was conducted as described ([9](#)). The sensitivity and specificity of this assay have not been established for white-tailed deer, but it has been used in seroprevalence studies of wild boar and fallow deer ([9,10](#)).

Insufficient volumes of serum samples prevented testing for antibodies against all pathogens ([Table](#)). Limited serum volume and lack of other (organ) samples also precluded additional analyses to verify presence of pathogens.

[Table](#)

Prevalence of serum antibodies against zoonotic pathogen antigens among white-tailed deer on 8 ranches, northern Mexico, 2004–2009*