## Pathogenic *Leptospira* infecting wild and captive crocodiles from the Yucatan Peninsula

Jonathan Pérez Flores\*1, Daniel Atilano López 2, Efrén Díaz Aparicio 3, Angélica Olivo Díaz4, Pierre Charruau5, J. Rogelio Cedeño-Vázquez1 and Erika M. Carrillo-Casas4

<sup>1</sup>El Colegio de la Frontera Sur, Unidad Chetumal, C.P. 77014, Chetumal, Quintana Roo, México (johnspf77@gmail.com; rogeliocedeno@gmail.com)

<sup>2</sup>Laboratorio de Diagnóstico, área Leptospirosis, Depto. Microbiología e Inmunología, Facultad de Medicina Veterinaria y Zootecnia

(daniel.atilano@yahoo.com.mx)

<sup>3</sup>CENID Microbiología, Instituto Nacional de Investigaciones Forestales y Agropecuarias (efredia@yahoo.com)

<sup>4</sup>Departamento de Biología Molecular e Histocompatibilidad, Subdirección de investigación, Hospital General "Dr. Manuel Gea González" (erikam.carrillo@salud.gob.mx; aolivod@yahoo.com)

<sup>5</sup>Centro del Cambio Global y la Sustentabilidad A.C., CP 86080, Villahermosa, Tabasco,

México (charruau\_pierre@yahoo.fr)

Abstract: Leptospira is a worldwide zoonosis that affects a large number of mammals, amphibians, and reptiles such as crocodilians. In this work, samples were collected from American crocodiles (*Crocodylus acutus*; n = 4) captured in Banco Chinchorro Biosphere Reserve, and from captive Morelet's crocodiles (*Crocodylus moreletii*; n = 14) on a farm in Chetumal, Quintana Roo, Mexico. Blood samples were obtained by venipuncture of the postoccipital sinus and serum was separated for subsequent analyses. Each sample was seeded in EMJH medium and observed in darkfield at 250X, and subsequently, DNA extraction was performed. For detection of the genus Leptospira, endpoint PCRs based on the 23S subunit and on the LipL32 lipoprotein were applied. For the detection of pathogenic species, PCR based on IS1500 insertion sequence was applied. Additionally, PCR based on the 16S subunit was used to identify if saprophytic Leptospira species were carried by the crocodiles. Cultures for isolation were subcultured in EMJH medium and followed up weekly for Leptospira isolation. Five crocodiles (5/18=27.77%) were identified as positive by darkfield observation, and four of them were confirmed to carry DNA from the genus Leptospira by PCR, and that they belonged to pathogenic species. No Saprophytic species were identified to be carried by the crocodiles by the 16S PCR. In conclusion, since The DNA of pathogenic species was identified in the peripheral blood of the crocodiles sampled, it is recommended to avoid contact with peridomestic animals that may function as carriers of Leptospira capable of infecting the water bodies and soil where the crocodiles remain.

<u>Keywords:</u> Crocodylus acutus, Crocodylus moreletii, Leptospirosis, Yucatan Peninsula, Zoonosis

*Type of presentation:* Poster

*Thematic area: Ex situ* conservation (Veterinary)