

# Antinociception and anti-inflammation induced by simvastatin in algeseometric assays in mice

Miranda, Hugo F.

Noriega, Viviana

Olavarria, Loreto

Zepeda, Ramiro J.

Sierralta, Fernando

Prieto, Juan C.

Statins, belonging to a well-known drug class used for lowering cholesterol through competitive inhibition of 3-hydroxy-3-methylglutaryl coenzyme A (HMG-CoA) reductase, also have other pleiotropic properties, such as anti-inflammatory action. The purpose of this study was to evaluate the antinociceptive and anti-inflammatory effects of simvastatin in five models of nociceptive behaviour. Oral gavage administration of simvastatin induced a dose-dependent inhibition of nociception for 1 day in the acetic acid writhing ( $ED_{50}=5.59\pm 0.07$ ), tail-flick ( $ED_{50}=112.96\pm 8.00$ ), hot-plate ( $ED_{50}=134.87\pm 2.20$ ), formalin hind paw ( $ED_{50}=19.86\pm 1.12$  in phase I and  $23.30\pm 2.05$  in phase II) and orofacial formalin ( $ED_{50}=5.54\pm 2.74$  in phase I and  $11.48\pm 1.88$  in phase II) tests. However, after 3 days, the values were in the acetic acid writhing ( $ED_{50}=6.14\pm 0.51$ ), tail-flick ( $ED_{50}=154\pm 8.88$ ), hot-plate ( $ED_{50}=136.14\pm 2.94$ ), formalin hind paw ( $ED_{50}=15.93\pm 0.42$  in phase I and  $17.10\pm 1.80$  in phase II) and orofacial f