

Molecular Biotechnology Programme

Uppsala University School of Engineering

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| Title (English) | |
| Effects of luminal hypertonicity on some duodenal functions | |
| Title (Swedish) | |
| as 40 to as high as 1000 mOsmol/kg H ₂ O, depending we responses to a hypotonic milieu. The aim of the present hypertonicity. Methods: The proximal duodenum was care min and the effects on duodenal wall contractions, me permeability studied in anesthetized rats, in the absence inhibitor parecoxib. Results: Perfusion of the duodenum mucosal permeability in COX-2 inhibited animals but ha parecoxib with duodenal contractions. Further, luminal hypinduced duodenal motility and reduced duodenal mucocontrols. Luminal hypertonicity increased bicarbonate sed duodenal responses to luminal hypertonicity varied contractions, while the inhibitory effect is due to inhibition of motors. | ed to contents with different osmolality, ranging from as low that we drink and eat. Previous studies have examined the t study was to examine the duodenal responses to luminal nnulated and perfused with a 350 mM NaCl solution for 30 ucosal bicarbonate secretion, net fluid flux and mucosal and presence of the selective cyclooxygenase-2 (COX-2) with 350 mM NaCl induced a delayed nine-fold increase in d no effect in controls or in animals that did not respond to expertonicity induced net fluid secretion, inhibited parecoxibosal bicarbonate secretion. No contractions were seen in secretion and induced net fluid secretion. Conclusions: The siderable between controls and COX-2 inhibited animals. Certion. The stimulatory effect is passive and due to solvent cility. |
| Keywords Duodenum, hypertonicity, cyclooxygenase, parecoxil | o, ⁵¹ Cr-EDTA permeability, fluid flux |
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