

# **Effects of lipopolysaccharide on the gastrointestinal transit in mice: role of nitric oxide and prostaglandin**

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摘要

## **Abstract**

AIM: To investigate the effect of lipopolysaccharide (LPS) on the diarrheogenic activity, gastrointestinal transit (GIT), and intestinal fluid content and the possible role of nitric oxide (NO) and prostaglandin E<sub>2</sub> (PGE<sub>2</sub>) in gastrointestinal functions of endotoxin-treated mice. METHODS: Diarrheogenic activity, GIT, and intestinal fluid content as well as nitric oxide and PGE<sub>2</sub> products were measured after intraperitoneal administration of LPS in mice. RESULTS: LPS dose-dependently accumulated abundant fluid into the small intestine, induced diarrhea, but decreased the GIT. Both nitric oxide and PGE<sub>2</sub> were found to increase in LPS-treated mice. Western blot analysis indicated that LPS significantly induced the protein expression of inducible nitric oxide synthase (iNOS) and cyclooxygenase-2 in mice intestines. Pretreatment with NG-nitro-L-arginine-methyl ester (L-NAME, a non-selective NOS inhibitor) or indomethacin (an inhibitor of prostaglandin synthesis) significantly attenuated the effects of LPS on the diarrheogenic activity and intestine content, but reversed the GIT. CONCLUSION: The present study suggests that the pathogenesis of LPS treatment may mediate the stimulatory effect of LPS on nitric oxide and PGE<sub>2</sub> production and NO/prostaglandin pathway may play an important role on gastrointestinal function