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interleukin-1beta. Oral administration of l-arginine, a precursor of nitric oxide (NO), has been shown to modulate immune responses to various agents. In the current study, we examined the effects of l-arginine on LPS-induced systemic inflammatory response and polymorphonuclear neutrophil (PMN) infiltration into the peritoneal cavity and lung. Oral l-arginine at a dose of 200 mg/kg inhibited the production of TNF-alpha and interleukin-6 in sera, thereby inhibiting the delayed-type hypersensitivity reaction induced by footpad injections of the antigen ovalbumin in mice. The suppressive effect of l-arginine was observed not only in LPS-treated mice, but also in mice that were challenged with LPS and lipoteichoic acid, suggesting that l-arginine administration has broad-spectrum anti-inflammatory

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properties. Furthermore, we observed that l-arginine strongly inhibited PMN migration into the peritoneal cavity and lung in mice. The results of the current study strongly indicate that oral administration of l-arginine can suppress the systemic inflammatory response and PMN migration induced by the immunostimulant LPS, suggesting a novel mechanism of action of this agent. Transcription and replication of phage P22 DNA in transformed E. coli K12. Replication and transcription of phage P22 DNA in E. coli K12 transformed with the helper plasmid pBR322, carrying the P22 transfer genes, have been compared

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