

论著

## 牛磺酸对大鼠胸主动脉的舒张作用及其机制的研究

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**摘要** 目的 研究牛磺酸舒血管作用的可能机制。方法 记录苯肾上腺素 (PE) 和KCl预收缩的离体大鼠主动脉环张力变化, 观察牛磺酸的舒血管作用及不同工具药对其作用的影响。结果 牛磺酸 ( $20\sim 80\text{ mmol}\cdot\text{L}^{-1}$ ) 对PE ( $1\text{ }\mu\text{mol}\cdot\text{L}^{-1}$ ) 或KCl ( $60\text{ mmol}\cdot\text{L}^{-1}$ ) 预收缩的大鼠主动脉环均有非内皮依赖的、浓度依赖性的舒张作用。在内皮完整的血管环, 左旋硝基精氨酸甲酯 ( $0.1\text{ mmol}\cdot\text{L}^{-1}$ ) 对牛磺酸的舒血管作用无明显影响;  $\beta$ -丙氨酸 ( $60\text{ mmol}\cdot\text{L}^{-1}$ ) 在PE预收缩的血管环增强牛磺酸的舒血管作用, 而在KCl预收缩的血管环则降低牛磺酸的舒血管作用; 在KCl预收缩基础上, 钾通道阻断剂格列本脲 ( $10\text{ }\mu\text{mol}\cdot\text{L}^{-1}$ ) 和四乙胺 ( $10\text{ mmol}\cdot\text{L}^{-1}$ ) 明显抑制牛磺酸的舒血管作用, 而4-氨基吡啶 ( $1\text{ mmol}\cdot\text{L}^{-1}$ ) 和BaCl<sub>2</sub> ( $1\text{ mmol}\cdot\text{L}^{-1}$ ) 无影响。结论 牛磺酸有浓度依赖性的血管舒张作用, 此作用不依赖血管内皮, 可能与其跨细胞膜转运有关, 可能有钙依赖性钾通道和ATP敏感性钾通道的参与。

**关键词** [牛磺酸](#) [主动脉, 胸](#) [血管舒张](#) [丙氨酸](#) [硝基精氨酸](#) [钾通道阻滞剂](#)

分类号

## Vasodilative effect and mechanism of taurine on thoracic aorta of rats

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### Abstract

**AIM** To investigate the vasodilative roles and the possible mechanisms of taurine on thoracic aorta of rats.

**METHODS** Isotonic tension of thoracic aortic rings precontracted by phenylephrine (PE,  $1\text{ }\mu\text{mol}\cdot\text{L}^{-1}$ ) or KCl ( $60\text{ mmol}\cdot\text{L}^{-1}$ ) was recorded. The vasorelaxing action of taurine and the influence of various drugs on it were observed in the rings with endothelium intact or endothelium denuded. **RESULTS** Taurine ( $20\text{-}80\text{ mmol}\cdot\text{L}^{-1}$ ) caused concentration-dependent relaxation in thoracic aortas with or without endothelium, and there was no significant difference between them.  $N^G$ -nitro-*L*-arginine methyl ester ( $0.1\text{ mmol}\cdot\text{L}^{-1}$ ) had no effect on the vasorelaxing action of taurine on thoracic aortas precontracted by PE or KCl.  $\beta$ -Alanine ( $60\text{ mmol}\cdot\text{L}^{-1}$ ) diminished the vasorelaxing action of taurine in KCl- precontracted rings, but enhanced the action in PE-precontracted rings. Tetraethylamine, an antagonist of calcium activated potassium channels ( $K_{Ca}$ ), and glibenclamide, an antagonist of ATP sensitive potassium channels ( $K_{ATP}$ ) attenuated the vasorelaxing effect of taurine, but 4-aminopyridine and BaCl<sub>2</sub> had no significant effect on the vasorelaxing action of taurine.

**CONCLUSION** The vasorelaxing action of taurine is endothelium-independent and associated with taurine transmembrane transportation;  $K_{Ca}$  and  $K_{ATP}$  may be involved in the action of taurine.

**Key words** [taurine](#) [aorta](#) [thoracic](#) [vasodilation](#) [alanine](#) [nitroarginine](#) [potassium channels blockers](#)

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