

[1]陈筱山,何选丽,朱丽娟,等.外源性硫化氢对原代神经元早老素1和B淀粉样蛋白的影响[J].第三军医大学学报,2013,35(19):2060-2064.

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## 外源性硫化氢对原代神经元早老素1和B淀粉样蛋白到:

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**Title:** Effects of exogenous hydrogen sulfide on expression of presenilin 1 and B-amyloid in primary cultured neurons

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**关键词:** [硫化氢](#); [原代神经元](#); [早老素1](#)

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**摘要:** **目的** 探讨外源性H<sub>2</sub>S对原代培养皮质神经元早老素1(presenilin 1, PS1)和B淀粉样蛋白(B-amyloid peptide, AB)的影响及机制。 **方法** ①NaHS为外源性H<sub>2</sub>S的供体,不同浓度NaHS(0、5、10、20、30、40、50 μmol/L)作用于原代培养的皮质神经元, TUNEL法检测细胞凋亡, 酶联免疫吸附实验(ELISA)检测AB1-42水平, Western blot检测PS1蛋白表达水平。②神经元分为对照组(0 μmol/L NaHS)、20 μmol/L NaHS组、A干预组(NaHS 20 μmol/L加PI3K通路阻断剂LY294002 20 μmol/L)和B干预组(NaHS 20 μmol/L加MAPK通路阻断剂PD98059 20 μmol/L), Western blot检测PS1蛋白表达。 **结果** ①5-20 μmol/L浓度NaHS干预时,不引起明显神经元凋亡; 30-50 μmol/L时,神经元凋亡明显增加(P<0.05); 10、20 μmol/L浓度NaHS显著降低AB1-42水平和PS1蛋白表达水平(P<0.05)。②与对照组相比,20 μmol/L NaHS组和B干预组PS1蛋白表达降低(P<0.05),而A干预组PS1蛋白表达无改变(P>0.05);而20 μmol/L NaHS组和B干预组间无差异(P>0.05);与A干预组相比,B干预组PS1蛋白表达显著降低(P<0.05)。 **结论** 低浓度外源性H<sub>2</sub>S(30 μmol/L浓度NaHS范围内)不引起神经元明显凋亡,降低AB1-42水平和PS1蛋白表达,并可能通过PI3K信号通路发挥降低PS1蛋白表达的作用。

**Abstract:** **Objective** To determine the effect of exogenous hydrogen sulfide (H<sub>2</sub>S) on the expression of presenilin 1 (PS1) and B-amyloid (AB) in primary cultured neurons. **Methods** Sodium hydrosulfide (NaHS), as H<sub>2</sub>S donor, was used

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to treat primary cultured cortical neurons derived from newborn SD rats at a dose of 0, 5, 10, 20, 30, 40 and 50  $\mu\text{mol/L}$  respectively for 24 h. Cell apoptosis was observed by TUNEL assay. AB1-42 level in the supernatant was measured by ELISA. The expression of PS1 protein was determined by Western blotting. The neurons was treated respectively by 0  $\mu\text{mol/L}$  NaHS (control), 20  $\mu\text{mol/L}$  NaHS, 20  $\mu\text{mol/L}$  NaHS+20  $\mu\text{mol/L}$  LY294002 (PI3K signal inhibitor), and 20  $\mu\text{mol/L}$  NaHS+20  $\mu\text{mol/L}$  PD98059B (MAPK signal inhibitor). Western blotting was used to detect the protein level of PS1.

**Results** The apoptotic rate had no obvious change when the dose of NaHS was 5, 10 and 20  $\mu\text{mol/L}$ , and was significantly increased at the dose of 30-50  $\mu\text{mol/L}$  ( $P<0.05$ ). NaHS treatment at 10 and 20  $\mu\text{mol/L}$  resulted in a decrease in AB1-42 concentration in the supernatant ( $P<0.05$ ) and in the protein level of PS1 ( $P<0.05$ ). Compared to the control, the expression of PS1 in the neurons treated by 20  $\mu\text{mol/L}$  NaHS in presence or absence of PD98059B was significantly decreased ( $P<0.05$ ), but had no change in the cells treated in presence of LY294002 ( $P>0.05$ ). There was also no difference in the protein level between the cells in presence or absence of PD98059B ( $P>0.05$ ). The PS1 protein level was decreased in the cells treated by PD98059B than those by LY294002.

**Conclusion** Low-dosed exogenous  $\text{H}_2\text{S}$  (<30  $\mu\text{mol/L}$  NaHS) does not induce obvious apoptosis, and decreases the protein levels of AB1-42 and PS1 in primary cultured neurons, which probably through PI3K signal pathway.

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