

药理学专栏

烟酰胺单核苷酸对RIN-m5f细胞中胰岛素分泌及PDX-1和FoxO1基因表达的影响

盛飞凤<sup>1,2</sup>, 任贤<sup>3</sup>, 戴幸平<sup>1</sup>, 徐潇静<sup>1</sup>, 董敏<sup>1</sup>, 裴奇<sup>1</sup>, 屈健<sup>1</sup>, 周智广<sup>4</sup>, 周宏灏<sup>1</sup>, 刘昭前<sup>1</sup>

1. 中南大学临床药理研究所, 遗传药理学湖南省重点实验室, 长沙 410078;
2. 广东省妇幼保健院药剂科, 广州 510010;
3. 上海绿谷制药有限公司, 上海 2012203;
4. 中南大学湘雅二医院内分泌科, 长沙 410011

摘要:

目的: 在细胞水平研究烟酰胺单核苷酸(nicotinamide mononucleotide, NMN)对胰岛素分泌的调节作用及其对与胰岛素分泌相关的重要转录因子胰十二指肠同源盒基因(pancreatic and duodenal homeobox-1, *PDX-1*)和分叉头框家族转录因子1(forkhead box-containing protein O-1, *FoxO 1*)基因表达的影响。方法: 采用大鼠胰岛素ELISA试剂盒检测RIN-m5f细胞胰岛素分泌水平。用Real-time PCR检测RIN-m5f细胞*PDX-1*和*FoxO 1*的mRNA表达水平。用Western印迹检测RIN-m5f细胞*PDX-1*蛋白表达水平。结果: 用瑞格列奈10 nmol/L+NMN 100 μmol/L处理RIN-m5f细胞48 h, 与空白对照及DMSO对照组相比, 胰岛素分泌量均显著增高( $P<0.05$ ); 与NMN 50 μmol/L组比较, 胰岛素分泌量的增高也有统计学意义( $P<0.05$ )。10, 50和100 μmol/L的NMN作用RIN-m5f细胞36 h, *PDX-1*的mRNA表达量均上调(依次为 $P<0.05$ ,  $P<0.01$ ,  $P<0.001$ )。100 μmol/L剂量组与10 μmol/L和50 μmol/L剂量组比较差异也有统计学意义( $P<0.001$ )。50, 100和200 μmol/L的NMN作用RIN-m5f细胞36或48 h, *PDX-1*的蛋白表达量与对照组比较差异无统计学意义( $P>0.05$ )。结论: NMN可以调控RIN-m5f细胞中胰岛素的分泌及*PDX-1*的mRNA表达水平。

关键词: 烟酰胺单核苷酸 胰十二指肠同源盒基因 分叉头框家族转录因子1 RIN-m5f

Effect of nicotinamide mononucleotide on insulin secretion and gene expressions of *PDX-1* and *FoxO1* in RIN-m5f cells

SHENG Feifeng<sup>1,2</sup>, REN Xian<sup>3</sup>, DAI Xingping<sup>1</sup>, XU Xiaojing<sup>1</sup>, DONG Min<sup>1</sup>, PEI Qi<sup>1</sup>, QU Jian<sup>1</sup>, ZHOU Zhiquan<sup>4</sup>, ZHOU Honghao<sup>1</sup>, LIU Zhaoqian<sup>1</sup>

1. Institute of Clinical Pharmacology, Hunan Key Laboratory of Pharmacogenetics, Xiangya School of Medicine, Central South University, Changsha 410078;
2. Department of Pharmacy, Maternal and Child Health Hospital of Guangdong, Guangzhou 510010;
3. Department of Medicine, Shanghai Green Valley Pharmaceutical Co., Ltd., Shanghai 2012203;
4. Department of Endocrinology, Second Xiangya Hospital, Central South University, Changsha 410011, China

Abstract:

Objective To investigate the effect of nicotinamide mononucleotide (NMN) on insulin secretion and gene expressions of pancreatic and duodenal homeobox 1 (*PDX-1*) and forkhead box-containing protein O-1 (*FoxO1*), which were important transcription factors for insulin secretion. Methods Insulin secretion level in RIN-m5f cells was detected by rat insulin ELISA detection kit. The mRNA expression levels of *PDX-1* and *FoxO1* in RIN-m5f cells were analyzed by real-time PCR. The protein expression of *PDX-1* was measured by Western blot. Results Insulin secretion levels in RIN-m5f cells treated with repaglinide (10 nmol/L) plus NMN (100 μmol/L) was significantly higher than those in the blank control, the DMSO control group, and the NMN (50 μmol/L) treated group ( $P<0.05$ ). The mRNA expression levels of *PDX-1* in RIN-m5f cells treated with NMN (10, 50 and 100 μmol/L) for 36 h were significantly higher than those in the control group ( $P<0.05$ ,  $P<0.01$ , and  $P<0.001$ , respectively). There was marked differences in the mRNA expression levels of *PDX-1* among different concentrations of NMN ( $P<0.001$ ), but no significant differences in the mRNA expression level of *FoxO1* ( $P>0.05$ ). No significant difference was found in the protein expression levels of *PDX-1* in RIN-m5f cells treated by NMN (50, 100, and 200 μmol/L) for 36 or 48 h compared with the control group ( $P>0.05$ ). Conclusion NMN can stimulate insulin secretion and upregulate the mRNA expression of *PDX-1* in RIN-m5f cells.

Keywords: nicotinamide mononucleotide pancreatic and duodenal homeobox 1 forkhead box-containing protein O-1 RIN-m5f cell

收稿日期 2010-11-05 修回日期 网络版发布日期

DOI: 10.3969/j.issn.1672-7347.2011.10.005

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This work was supported by the National High-Tech R&D Program of China ("863" Program) (2009AA22704), the National Natural Science Foundation of China (30873089, 81173129), the Program for Changjiang Scholars and Innovative Research Team in University (IRT0946), the Open Foundation of Innovative Platform in University of Hunan Province of China (10K078), and the Science and Technology Plan Key Grant of Hunan Province of China (2009TP4068-2), the Fundamental Research Funds for the Central Universities (201023100001).

通讯作者: LIU Zhaoqian, E-mail: liuzhaoqian63@126.com

作者简介: SHENG Feifeng, master, assistant pharmacist, mainly engaged in the research of pharmacogenetics.

作者Email: liuzhaoqian63@126.com

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