



Protective effects of naringin against gp120-induced injury mediated by P2X₇ receptors in BV2 microglial cells

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Genet. Mol. Res. 15 (2): gmr.15028649

Received March 23, 2016

Accepted April 11, 2016

Published May 13, 2016

DOI <http://dx.doi.org/10.4238/gmr.15028649>

ABSTRACT. This study was aimed at exploring the effects of P2X₇ receptors on gp120-induced injury and naringin's protective effects against gp120-induced injury in BV2 microglia. BV2 microglia injury model was established by gp120 treatment and MTS assay was used to verify whether naringin has a cell-protective effect against gp120-induced injury. Changes in P2X₇ receptor expression were assayed using RT-PCR, qPCR, and western blot. Results showed that the ODs of the Ctrl, gp120, gp120+naringin, and gp120+BBG groups were 0.91 ± 0.10 , 0.71 ± 0.09 , 0.83 ± 0.10 , and 0.83 ± 0.10 , respectively. Compared to the control group, the gp120 group showed a significantly decreased cell survival rate. Cell survival rates of the gp120+naringin group increased significantly compared to those of the gp120 group, while no difference was observed when compared to the gp120+BBG group. The relative P2X₇ mRNA expression levels

in the Ctrl, gp120, gp120+naringin, and gp120+BBG groups were 0.73 ± 0.06 , 1.05 ± 0.06 , 0.78 ± 0.05 , and 0.81 ± 0.04 , respectively. The corresponding P2X₇ protein expression levels were 0.46 ± 0.04 , 0.79 ± 0.04 , 0.38 ± 0.07 , and 0.42 ± 0.06 . P2X₇ mRNA and protein expression in the gp120 group increased significantly compared to those in the control group, and declined in the gp120+naringin group compared to those in the gp120 group. Therefore, P2X₇ receptors might be involved in gp120-induced injury in BV2 microglia, and naringin might play a protective role by inhibiting the up-regulated expression of P2X₇ receptors.

Key words: AIDS-dementia complex; BV2 microglial cells; Naringin; Gp120; P2X₇ receptor