Potential Mechanism of Blood Vessels Protection by Resveratrol, a

Red Wine Component

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Abstract

Resveratrol-mediated <u>heme</u> oxgenase-1 (HO-1) induction has been shown to occur in primary <u>neuronal</u> cultures and has been implicated as having potential neuroprotective action. Further, <u>antioxidant</u> properties of <u>resveratrol</u> have been reported to protect against <u>coronary heart disease</u>. We attempted to examine the HO-1 inducing potency of <u>resveratrol</u> and the regulatory mechanism of its induction in rat <u>aortic smooth muscle cells</u> (RASMC). We showed that resveratrol-induced HO-1 expression was concentration- and time-dependent. The level of HO-1 expression and its <u>promoter</u> activity mediated by <u>resveratrol</u> was <u>attenuated</u> by nuclear factor-kappa B (<u>NF-kappaB</u>) inhibitors, but not by <u>mitogen-activated protein kinase</u> (<u>MAPK</u>) inhibitors. Deletion of <u>NF-kappaB</u> binding sites in the <u>promoter region</u> strongly reduced <u>luciferase</u> activity. Collectively, we suggest that resveratrol-mediated HO-1 expression occurs, at least in part, through the <u>NF-kappaB</u> pathway.