Anti-apoptotic mechanism of silkworm hemolymph in staurosporine-induced apoptosis

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There exist some common pathways in apoptosis despite a wide range of inducing signals, and mitochondria play a crucial role especially by releasing cytochrome c into cytosol, which forms complex with Apaf-1 to turn on the caspase cascade reaction1). Silkworm hemolymph (SH) has shown anti-apoptotic activities in mammalian2) and insect cell apoptosis3), and five 30kDa proteins in SH are the major inhibitors of apoptosis. Here we developed anti-apoptotic mechanism of SH in staurosporine-induced HeLa cell apoptosis. SH did not directly inhibit caspase-3 and caspase-9 activities in cell-free reaction, but rather increased caspase activities by improving caspase reaction condition. This supports the claim that anti-apoptotic effect of SH lies in further upstream events than caspase activation. Cytochrome c release and the translocation of Bax to mitochondria after staurosporine treatment were blocked by SH. This indicates that SH affects a step above Bax translocation such as Bax conformational change by Bid in staurosporine-induced HeLa cell apoptosis. SH effects on cytosolic calcium concentration, generation of reactive oxygen species, and mitochondrial membrane potential were also determined.

References

- Kirsten Lauber, Helga A. E. Appel, Stephan F. Schlosser, Michael Gregor, Klaus Schulze-Osthoff, and Sebastian Wesselborg, "The Adapter Protein Apoptotic Protease-activating Factor-1 Is Proteolytically Processed during Apoptosis" (2001), JBC, 276, 29772-29781
- 2. Shin Sik Choi, Won Jong Rhee, and Tai Hyun Park, "Inhibition of Human Cell Apoptosis by Silkworm Hemolymph" (2002), *Biotechnol. Prog.*, **18**, 874-878
- 3. Won Jong Rhee, Eun Jeong Kim, and Tai Hyun Park, "Silkworm hemolymph as a potent inhibitor of apoptosis in Sf9 cells" (2002), BBRC, 295, 779-783