

Ordering of ceramide formation, caspase activation, and mitochondrial changes during CD95- and DNA damage–induced apoptosis

Annemiek D. Tepper, ... , Wim J. van Blitterswijk, Jannie Borst

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Erratum

J. Clin. Invest. 103:971–978 (1999) During the production process, the word caspase was misspelled in the title; the correct title appears above. Also, in the legend for Table 1 the mu symbol (μ) was formatted incorrectly; the correct legend appears below. We regret the error. Table 1 Jurkat cells (J16) were preincubated for 2 h with zVAD-fmk (50 μ M), DEVD-CHO (100 μ M) or left untreated and then exposed to etoposide (10 μ g/ml) or IR (30 Gy). After 16 h incubation, Cer content, nuclear fragmentation, mitochondrial transmembrane potential and cell viability were determined in parallel samples as described in the Methods section. The results are representative of two independent experiments.

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Annemiek D. Tepper, Evert de Vries, Wim J. van Blitterswijk, and Jannie Borst

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Table 1

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Estrogen receptor α mediates the nongenomic activation of endothelial nitric oxide synthase by estrogen

Zhong Chen, Ivan S. Yuhanna, Zoya Galcheva-Gargova, Richard H. Karas, Michael E. Mendelsohn, and Philip W. Shaul

J. Clin. Invest. 103:401–406 (1999)

During the production process, panels *a* and *b* of Figure 2 were mistakenly repeated as panels *c* and *d*. The correct display of the figure and accompanying legend is reproduced here. We regret the error and have provided corrected reprints to the corresponding author: Philip W. Shaul, Department of Pediatrics, University of Texas Southwestern Medical Center, 5323 Harry Hines Boulevard, Dallas, Texas 75235-9063, USA. Phone: (214) 648-2015; Fax: (214) 648-2481; E-mail: pshaul@mednet.swmed.edu.

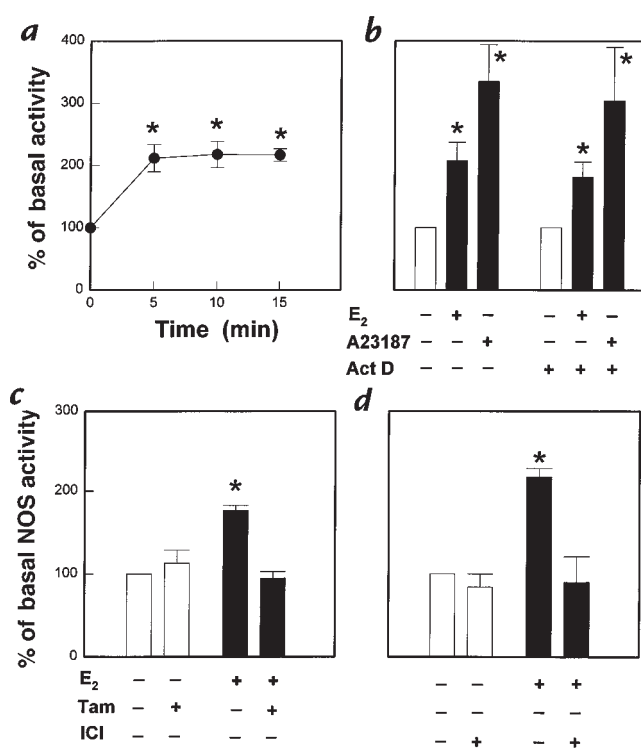


Figure 1

Rapid activation of eNOS in endothelial cells. (a) Effect of E₂ on eNOS activity in intact PAEC. [³H]L-arginine conversion to [³H]L-citrulline was measured over 5–15 min in the presence of 10⁻⁸ M E₂. (b) Effect of actinomycin D (Act D) on the rapid activation of eNOS. After 120 min preincubation in the absence or presence of 25 μ g/ml Act D, 15 min incubations were done with or without continued Act D and either 10⁻⁸ M E₂ or the calcium ionophore A23187 (10⁻⁵ M). (c) Effect of tamoxifen on E₂-stimulated eNOS activity. Fifteen-minute incubations were performed in the absence or presence of 10⁻⁸ M E₂, with or without 10⁻⁶ M tamoxifen (Tam) added simultaneously. Partial inhibition (50%–70%) was also noted with 10⁻⁸ M Tam (13). (d) Effect of ICI 182,780 on E₂-stimulated eNOS activity. Fifteen-minute incubations were performed in the absence or presence of 10⁻⁸ M E₂, with or without 10⁻⁵ M ICI 182,780 added simultaneously. Full inhibition was also observed with 10⁻⁶ M ICI 182,780 (13). Values are mean \pm SEM; *n* = 4–6. **P* < 0.05 vs. basal. E₂, estradiol-17 β ; eNOS, endothelial nitric oxide; PAEC, pulmonary artery endothelial cells.