

# always your partner



## M30 CytoDeath<sup>™</sup> ELISA (PEVIVA<sup>®</sup>)

Measurement of soluble caspase-cleaved fragments of the intermediate filament protein keratin 18 (K18) containing the M30 neo-epitope (K18-Asp396-NE). Released from human, monkey or bovine epithelial cells in cell culture.

Cat. No.:	10900
Tests:	96
Method:	ELISA
Range:	250 – 3000 U/L (units are defined against a synthetic peptide standard containing the M30 and M6 epitopes; 1 U/L = 1.24 pM)
LLOD:	60 U/L
Incubation time:	4.5 hours
Sample volume:	25 μl
Sample type:	cell cultures (epithelial cells)
Sample preparation:	Store samples at $2 - 8$ °C up to 4 hours. For longer periods, store samples frozen at -20 °C or lower.
Species:	Human, primates

### Intended Use:

Quantitative measurement of the apoptotic cell death biomarker K18-Asp396-NE in cell culture experiments. Can be used for cell lysates and/or culture supernatants. The assay only detects apoptosis in cells of epithelial origin that express K18. Cells should be of human, monkey or bovine origin. To be used to determine accumulation of caspase-cleaved K18 (ccK18) in cell cultures, providing an integrative measure of apoptosis. The K18-Asp396 neo-epitope is formed by caspase-3, -7 or -9 activation.

\*Note:

caspase-cleaved K18 = ccK18 previously Cytokeratin 18 (CK18/ccCK18)

### **References:**

- Fayad W, et al. (2009). Identification of a novel topoisomerase inhibitor effective in cells overexpressing drug efflux transporters. *PLoS One* 4(10):e7238.
- Hernlund E, et al. (2009). Ovarian carcinoma cells with low levels of b-F1-ATPase are sensitive to combined platinum and 2-deoxy-D-glucose treatment. *Mol Cancer Ther* 2009;8(7).
- Herrmann R, *et al.* (2008). Screening for Compounds that Induce Apoptosis of Cancer Cells Grown as Multicellular Spheroids. *J Biomol Screen*. 13(1):1-8.
- Lakshmikanthan V, et al. (2006). SAHA-sensitized prostate cancer cells to TNFalpha-related apoptosis-inducing ligand (TRAIL): mechanisms leading to synergistic apoptosis. Int J Cancer 119:221-8.
- Cummings J, et al. (2005) Validation of pharmacodynamic assays to evaluate the clinical efficacy of an antisense compound (AEG 35156) targeted to the X-linked inhibitor of apoptosis protein XIAP. Br J Cancer 92, 532-538.
- Erdal H, et al. (2005). Induction of lysosomal membrane permeabilization by compounds that activate p53-independent apoptosis. *Proc. Natl. Acad. Sci. USA* 102, 192-197.
- Schutte B, *et al.* (2004). Keratin 8/18 breakdown and reorganization during apoptosis. *Exp Cell Res.* 297, 11-26.
- Kramer G, *et al.* (2004). Differentiation between Cell Death Modes using Measurements of Different Soluble Forms of Extracellular Cytokeratin 18. *Cancer Research* 64, 1751-1756.
- Hägg, M. et al. (2002). A novel high-through-put assay for screening of pro-apoptotic drugs. *Invest. New Drugs*, 20: 253-259.
- Leers MP, *et al.* (1999). Immunocytochemical detection and mapping of a Cytokeratin 18 neoepitope exposed during early apoptosis. *J Pathol* 187, 567-572.

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