

Cytochrome c Apoptosis WB Antibody Cocktail ab110415

[9 References](#) [1 图像](#)

概述

产品名称	Cytochrome c Apoptosis WB抗体Cocktail
检测类型	Quantitative
种属反应性	与反应: Human
产品概述	<p>The permeabilization of mitochondrial outer membrane and the subsequent release of cytochrome c and other apoptogenic proteins from mitochondrial intermembrane space into the cytoplasm is considered a hallmark of many apoptotic pathways. Therefore assaying these proteins in mitochondrial and cytoplasmic fractions is of prime interest for many researchers.</p> <p>ab 110415 (MSA12) is a Western blot antibody cocktail that allows for the detection of cytochrome c in cytoplasmic and mitochondria-containing fractions for determining the proportion of released cytochrome c from mitochondria to the cytoplasm from apoptosis. The kit includes antibodies against a cytoplasmic protein marker, glyceraldehyde-3-phosphodehydrogenase (GAPDH), and 2 mitochondrial markers, pyruvate dehydrogenase subunit E1-alpha (a matrix marker), and ATP synthase subunit alpha (an inner membrane marker). This set of control markers allows for the monitoring and/or optimization of the permeabilization conditions.</p> <p>Cocktail Antibodies:</p> <p>Mouse anti Cyt. c monoclonal: Amount: 50 µg Working concentration: 1 µg/ml</p> <p>Mouse anti GAPDH monclonal: Amount: 5ug Working concentration: 0.1 µg/ml</p> <p>Mouse anti PDH-E1-alpha monoclonal: Amount: 100ug Working concentration: 2 µg/ml</p> <p>Mouse anti C-V-alpha monoclonal: Amount: 25ug Working concentration: 0.5 µg/ml</p>

说明 This product was previously called ApoTrack™ Cytochrome c Apoptosis WB Antibody Cocktail

Other apoptosis assays

For more apoptosis assays, review the [apoptosis assay and apoptosis marker guide](#).

经测试应用 适用于: WB

性能

存放说明 Please refer to protocols.

组件	180 µg
HeLa Lysate Control	1 x 50µg
Pre-mixed Solution of 4 monoclonal Antibodies	1 x 180µg

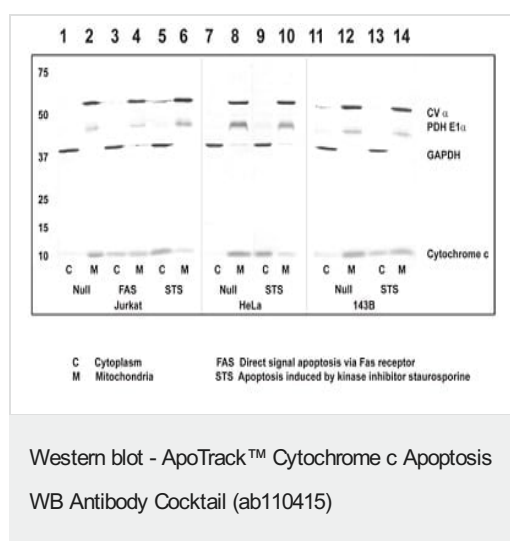
应用

The Abpromise guarantee **Abpromise™** 承诺保证使用 ab110415 于以下的经测试应用

“应用说明”部分 下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。

应用	Ab评论	说明
WB		Use a concentration of 3.6 µg/ml. The antibody cocktail (0.9 mg/ml) should be diluted 250x to a final working concentration of 3.6 µg/ml for Western blotting.

图片



In this experiment, apoptosis was induced in Jurkat and 143B osteosarcoma cells by FAS and also by treatment with staurosporine (HeLa cells were also treated, but only with STS). Mitochondrial and cytoplasmic fractions were isolated (using kit Cell Fractionation Kit [ab109719/MS861](#)) and probed using ab110415 (MSA12). As is clear from the gels, cytochrome c has translocated partially in FAS-induced cells and STS-treated osteosarcoma cells, and almost completely in STS-treated Jurkat and HeLa cells. The three control targets allow for verification of the "cleanness" of the cell fractionation.

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