## Cytochrome c (6H2.B4) Mouse mAb





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Applications: IP, IF-IC, FC-FP	<b>Reactivity:</b> H M R	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 14	Source/Isotype: Mouse IgG1	<b>UniProt ID:</b> #P999999	<b>Entrez-Gene Id:</b> 54205		
Information Im Im		Immunofluorescence	<b>Application</b> Immunoprecipitation Immunofluorescence (Immunocytochemistry) Flow Cytometry (Fixed/Permeabilized)			<b>Dilution</b> 1:100 1:300 1:200 - 1:400		
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.						
Specificity/Ser	nsitivity	Cytochrome c (6H2.B4) Mouse mAb recognizes endogenous levels of total cytochrome c protein by immunofluorescence. This antibody is not recommended for western blot.				ne c protein by		
Source / Purifi	cation	Monoclonal antibody is produced by immunizing animals with recombinant rat cytochrome c prote						
Background		Cytochrome c is a well conserved electron-transport protein and is part of the respiratory chain localized to mitochondrial intermembrane space (1). Upon apoptotic stimulation, cytochrome c released from mitochondria associates with procaspase-9 (47 kDa)/Apaf 1. This complex processes caspase-9 from inactive proenzyme to its active form (2). This event further triggers caspase-3 activation and eventually leads to apoptosis (3).						
Background R	eferences	1. Schagger H.H. et al. (2002) <i>Biochem. Biophys. Acta.</i> 1555, 154-159. 2. Li, P. et al. (1997) <i>Cell</i> 91, 479-489. 3. Liu, X. et al. (1996) <i>Cell</i> 86, 147-157.						
Species Reacti	vity	Species reactivity is determined by testing in at least one approved application (e.g., western blot).						
Applications K	ley	<b>IP:</b> Immunoprecipitation <b>IF-IC:</b> Immunofluorescence (Immunocytochemistry) <b>FC-FP:</b> Flow Cytometry (Fixed/Permeabilized)						
Cross-Reactivi	ty Key	H: Human M: Mouse R: Rat						
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