KAP-1 (4E1) Mouse mAb



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Applications: W, IF-IC	Reactivity: H	Sensitivity: Endogenous	MW (kDa): 100, 75	Source/Isotype: Mouse IgG1	UniProt ID: #Q13263	Entrez-Gene Id: 10155
Product Usage Information		Application Western Blotting Immunofluorescence	: (Immunocytochen	istry)		Dilution 1:1000 1:200
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/Sensitivity		KAP-1 (4E1) Mouse mAb recognizes endogenous levels of total KAP-1 protein.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with recombinant protein specific to the carboxy terminus of human KAP-1 protein.				
Background		KAP-1 is a member of the TIF1 (transcriptional intermediary factor 1) family, a group of transcriptional regulators that play key roles in development and differentiation. Members of this family are characterized by the presence of two conserved motifs – an N-terminal RING-B box-coiled-coil motif and a C-terminal PHD finger and bromodomain unit (1,2). KAP-1 is a corepressor for KRAB (Kruppel associated box) domain containing zinc finger proteins. The KRAB domain containing zinc finger proteins are a large group of transcription factors that are vertebrate-specific, varied in their expression patterns between species, and thought to regulate gene transcription programs that control speciation (3,4).KAP-1 has been shown to be essential for early embryonic development and spermatogenesis (6,5). It functions to either activate or repress transcription in response to environmental or developmental signals by chromatin remodeling and histone modification. The recruitment and association of KAP-1 with heterochromatin protein (HP1) is essential for transcriptional repression, and for progression through differentiation of F9 embryonic carcinoma cells (6,7). KAP-1 also plays a role in the DNA damage response. Phosphorylation of KAP-1 on Ser824 occurs in an ATM-dependent manner in response to genotoxic stress and is thought to be essential for chromatin relaxation, which is in turn required for the DNA damage response (8).				
Background References		1. Le Douarin, B. et al. (1995) <i>EMBO J.</i> 14, 2020-2033. 2. Le Douarin, B. et al. (1996) <i>EMBO J.</i> 15, 6701-6715. 3. Friedman, J.R. et al. (1996) <i>Genes Dev.</i> 10, 2067-2078. 4. Krebs, C.J. et al. (2005) <i>Genomics</i> 85, 752-761. 5. Weber, P. et al. (2002) <i>Development</i> 129, 2329-2337. 6. Cammas, F. et al. (2004) <i>Genes Dev.</i> 18, 2147-2160. 7. Cammas, F. et al. (2007) <i>Differentiation</i> 75, 627-37. 8. Ziv, Y. et al. (2006) <i>Nat. Cell Biol.</i> 8, 870-876.				
Species Reactivity		Species reactivity is determined by testing in at least one approved application (e.g., wes				western blot).
Western Blot Buffer		IMPORTANT: For western blots, incubate membrane with diluted primary antibody in dry milk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.				n 5% w/v nonfat
Applications Key		W: Western Blotting IF-IC: Immunofluorescence (Immunocytochemistry)				
Cross-Reactivity Key		H: Human				

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