

40758

Phospho-CASC5 (Thr943/Thr1155) (D8D4N) Rabbit mAb



Orders: 877-616-CELL (2355)

orders@cellsignal.com

Support: 877-678-TECH (8324)

Web: info@cellsignal.com

cellsignal.com

3 Trask Lane | Danvers | Massachusetts | 01923 | USA

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Applications: W, IF-IC	Reactivity: H	Sensitivity: Endogenous	MW (kDa): 265	Source/Isotype: Rabbit IgG	UniProt ID: #Q8NG31	Entrez-Gene Id: 57082
Product Usage Information	2	Application Western Blotting Immunofluorescence	e (Immunocytochen	nistry)		Dilution 1:1000 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.				
		For a carrier free (BSA and azide free) version of this product see product #72586.				
Specificity/Sensitivity		Phospho-CASC5 (Thr943/Thr1155) (D8D4N) Rabbit mAb recognizes endogenous levels of CASC5 protein only when phosphorylated at Thr943 or Thr1155. The protein sequences surrounding these two sites are identical.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Thr943 of human CASC5 protein.				
Background		Kinetochores are mitotic structures that form on centromeres and attach to mitotic spindle microtubules. Kinetochore attachment to microtubules regulates chromosome segregation and progression through mitosis. Unattached kinetochores signal to the spindle assembly checkpoint (SAC) machinery, arresting cells in mitosis (1). CASC5, also known as Knl1 or Blinkin, is the largest subunit of the Knl1–Mis12–Ndc80 complex (KMN) network, a structural component of kinetochores required for microtubule binding. CASC5 functions in the formation of kinetochore–microtubule attachments, chromosome segregation, and in activating the SAC. CASC5 has been implicated in human diseases, including leukemia and microcephaly (2). Activation of the SAC is regulated in part by mitotic phosphorylation of CASC5 at several sites, including Ser24, Ser60, Thr943, and Thr1155 (3,4). The sequences surrounding Thr943 and Thr1155 are identical.				
Background References		 Foley, E.A. and Kapoor, T.M. (2013) Nat Rev Mol Cell Biol 14, 25-37. Ghongane, P. et al. (2014) J Cell Sci 127, 3415-23. Nijenhuis, W. et al. (2014) Nat Cell Biol 16, 1257-64. Schleicher, K. et al. (2017) Open Biol 7, 170099. doi: 10.1098/rsob.170099. 				
Species Reactivity		Species reactivity is determined by testing in at least one approved application (e.g., western blot).				
Western Blot Buffer		IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.				
Applications Key		W: Western Blotting IF-IC: Immunofluorescence (Immunocytochemistry)				
Cross-Reactivity Key		H: Human				

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