Calpain 2 L Antibody	arge Sul	bunit (M-ty	vpe)			BISignaling CHNOLOGY® 877-616-CELL (2355) orders@cellsignal.com 877-678-TECH (8324) info@cellsignal.com	
#2					Lana Danvara Ma	cellsignal.com	
For Research Use Only. N	ot for Use in	Diagnostic Proc	edures.	3 Hask	Lane   Danvers   Ma	ssachusetts 01923 USA	
Applications: WB	Reactivity: H M R	Sensitivity: Endogenous	<b>MW (kDa):</b> 80	Source: Rabbit	UniProt ID: #P17655	Entrez-Gene Id: 824	
Product Usage Information		plication stern Blotting			Dilution 1:1000		
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.					
Specificity / Sensitiv	vity Calpain 2 Large Subunit (M-type) Antibody detects endogenous levels of total calpain 2 (large subunit) protein. The antibody detects full-length calpain 2 as well as calpain 2 autoproteolytically cleaved at serie 20. The antibody does not detect recombinant calpain 1.						
Source / Purification	Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to the human sequence of calpain 2 (large subunit). Antibodies are purified by protein A and peptide affinity chromatography.						
Background	sma calp millin stud calci sign nega influ and	Calpain is a calcium-dependent thiol proteinase that is functionally active as a heterodimer composed of a small regulatory subunit and one of at least two large catalytic subunits (calpain 1 or calpain 2). <i>In vitro</i> , calpain 1 (mu-calpain) requires micromolar levels of calcium, while calpain 2 (M-calpain) requires millimolar levels of calcium for activation. The regulation of calpain <i>in vivo</i> is the subject of many current studies, which suggest that proteolytic activity is regulated post-transcriptionally by mechanisms such as calcium requirements, subcellular localization of the heterodimer, phosphorylation via the EGFR-Erk signaling cascade, endogenous inhibitors (calpastatin), and autoproteolytic cleavage (1). Calpastatin negatively regulates autoproteolytic cleavage of calpain 1 between Gly27 and Leu28 (2). Calpain influences cell migration by modifying rather than degrading its substrates responsible for cell adhesion and cytoskeletal arrangement. Control of calpain activity has caught the attention of drug development since limiting its activity could mute invasiveness of tumors or chronic inflammation (1).					
Background Referer	und References1. Perrin, B.J. and Huttenlocher, A. (2002) Int. J. Biochem. Cell Biol. 34, 722-725.2. Melloni, E. et al. (1996) Biochem. Biophys. Res. Commun. 229, 193-197.						
Species Reactivity	Spec	ies reactivity is dete	ermined by testing i	n at least one appro	ved 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	WB:	Western Blotting					
Cross-Reactivity Ke	Χ: Χε	<ul> <li>H: human M: mouse R: rat Hm: hamster Mk: monkey Vir: virus Mi: mink C: chicken Dm: D. melanogaster</li> <li>X: Xenopus Z: zebrafish B: bovine Dg: dog Pg: pig Sc: S. cerevisiae Ce: C. elegans Hr: horse</li> <li>GP: Guinea Pig Rab: rabbit All: all species expected</li> </ul>					
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