

## **Mouse Kremen-1 Antibody**

Monoclonal Rat IgG<sub>2A</sub> Clone # 252525 Catalog Number: MAB1647

DESCRIPTION		
Species Reactivity	Mouse	
Specificity	Detects mouse Kremen-1 in direct ELISAs and Western blots. In Western blots, 100% cross-reactivity with recombinant human (rh) Kremen-is observed. In direct ELISAs, approximately 10% cross-reactivity with rhKremen-1 is observed. In direct ELISAs and Western blots, no cross-reactivity with recombinant mouse Kremen-2 is observed.	
Source	Monoclonal Rat IgG <sub>2A</sub> Clone # 252525	
Purification	Protein A or G purified from hybridoma culture supernatant	
Immunogen	Mouse myeloma cell line NS0-derived recombinant mouse Kremen-1 Ala20-Gly395 Accession # Q640Q6	
Formulation	Lyophilized from a 0.2 µm filtered solution in PBS with Trehalose. See Certificate of Analysis for details. *Small pack size (-SP) is supplied either lyophilized or as a 0.2 µm filtered solution in PBS.	
APPLICATIONS		
Please Note: Optimal dilut	ions should be determined by each laboratory for each application	n. General Protocols are available in the Technical Information section on our website.
	Recommended Concentration	Sample
Western Blot	1 ug/ml	Recombinant Mouse Kremen 1 (Catalog # 1647-KP)

PREPARATION AND STORAGE		
Reconstitution	Reconstitute at 0.5 mg/mL in sterile PBS.	
Shipping	The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below. *Small pack size (-SP) is shipped with polar packs. Upon receipt, store it immediately at -20 to -70 °C	
Stability & Storage	Use a manual defrost freezer and avoid repeated freeze-thaw cycles.  12 months from date of receipt, -20 to -70 °C as supplied.  1 month, 2 to 8 °C under sterile conditions after reconstitution.  6 months, -20 to -70 °C under sterile conditions after reconstitution.	

## BACKGROUND

Kremen (Kringle-containing protein marking the eye and the nose) proteins are type I transmembrane proteins that contain extracellular kringle, WSC and CUB domains and an intracellular region without any conserved motifs (1). Two related members, Kremen-1 and -2, have been identified. Kremens bind a subset of the secreted Dickkopf (Dkk) proteins (Dkk-1, -2, and -4) with high affinity to modulate the canonical Wnt signaling pathway that is transduced by the ternary receptor complex composed of Wnt, the seven-transmembrane domain receptor Frizzled, and the LDL-receptor-related protein 5/6 (LRP5/6) co-receptor (2, 3). Within the Dkk family, Dkk-1 and -4 bind directly to the LRP5/6 co-receptor to antagonize the canonical Wnt/β-catenin signaling pathway, but not the planar cell polarity (PCP) signaling pathway that does not involve LRP5/6 (4). In contrast, Dkk-3 has no effect on Wnt signaling and Dkk-2 can function either as an LRP agonist or antagonist, depending on whether the cell expresses Kremen (5). Kremen co-operates with Dkk to antagonize Wnt signaling via formation of a Kremen-Dkk-LRP ternary complex that triggers the internalization and clearance of the complex from the cell surface (3). All three extracellular domains but not the cytoplasmic region of a membrane anchored Kremen are needed for binding to the second cysteine-rich domain of Dkks (3). Mouse Kremen-1 cDNA encodes a 473 amino acid (aa) glycosylated protein with a putative 19 aa signal peptide, a 372 aa extracellular domain, a 21 aa transmembrane domain and a 60 aa cytoplasmic domain. In the extracellular domain, it shares 92% and 41% amino acid sequence identity with human Kremen-1 and mouse Kremen-2, respectively. Mouse Kremen-1 is widely expressed in diverse embryonic (apical ectodermal ridge of the developing fore- and hindlimb buds, telencephalon and the first brachial arch, myotome and sensory tissues) and adult (lung, heart, kidney, skeletal muscle and testis) tissues (1).

## References:

- 1. Nakamura, T. et al. (2001) Biochim Biophys Acta 1518:63.
- 2. Davidson G. et al. (2002) Development 129:5587.
- 3. Mao, B. et al. (2002) Nature 417:664.
- 4. Zorn, A.M. (2001) Curr. Biol. 11:R592.
- 5. Mao, B. and C. Niehrs (2003) Gene 302:179.

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