

**LRIG1 (E1U2B) Rabbit mAb**

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<b>Applications:</b> W, IP	<b>Reactivity:</b> H	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 140	<b>Source/Isotype:</b> Rabbit IgG	<b>UniProt ID:</b> #Q96JA1	<b>Entrez-Gene Id:</b> 26018
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**Product Usage Information****Application**

Western Blotting  
Immunoprecipitation

**Dilution**

1:1000  
1:50

**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**

LRIG1 (E1U2B) Rabbit mAb recognizes endogenous levels of total LRIG1 protein.

**Source / Purification**

Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues near the carboxy terminus of human LRIG1 protein.

**Background**

Leucine-rich immunoglobulin repeats 1 (LRIG1) is a type I transmembrane protein containing 15 leucine rich repeats and three immunoglobulin domains in the extracellular domain. Researchers characterize LRIG1 as a negative regulator of receptor tyrosine kinase signaling. In studies with ErbB family members and Met kinase, LRIG regulates signaling by increasing ubiquitination and lysosomal degradation of the receptors (1,2). Additional work indicates that LRIG1 plays a role in neurotropic signaling by negatively regulating Ret signaling (3,4). Expression profile studies demonstrate that LRIG1 is a marker in the quiescent population of stem cells in the intestine (5). Interestingly, the genetic ablation of one allele of LRIG1 in mice with an APC+/- background results in development of highly dysplastic adenomas, indicating a role for LRIG1 in tumor suppression (1). Indeed, down-regulation of LRIG1 is tentatively involved in tumor aggressiveness in several tumor types, including glioma (6), head and neck cancer (7), and cervical adenocarcinoma (8).

**Background References**

- Powell, A.E. et al. (2012) *Cell* 149, 146-58.
- Segatto, O. et al. (2011) *J Cell Sci* 124, 1785-93.
- Shattuck, D.L. et al. (2007) *Mol Cell Biol* 27, 1934-46.
- Ledda, F. et al. (2008) *J Neurosci* 28, 39-49.
- Muñoz, J. et al. (2012) *EMBO J* 31, 3079-91.
- Mao, F. et al. (2013) *Int J Oncol* 42, 1081-7.
- Sheu, J.J. et al. (2013) *Oncogene* 33, 1375-84.
- Muller, S. et al. (2013) *Int J Oncol* 42, 247-52.

**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 **IP:** Immunoprecipitation

**Cross-Reactivity Key**

**H:** Human

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