

Product datasheet for AP01369PU-N

OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

Smad Interacting Protein 1 (ZEB2) Rabbit Polyclonal Antibody

Product data:

Product Type: Primary Antibodies

Applications: IHC, WB

Recommended Dilution: Western Blot: 1/500-1/1000.

Immunohistochemistry on Paraffin Sections: 1/50-1/200.

Reactivity: Human, Mouse

Host: Rabbit

Clonality: Polyclonal

Immunogen: Synthetic peptide, corresponding to amino acids 71-120 of Human SIP1.

Specificity: This antibody detects endogenous levels of SIP1 protein. (region surrounding Glu101)

Formulation: Phosphate buffered saline (PBS), pH~7.2

State: Aff - Purified

State: Liquid purified Ig fraction (> 95% pure by SDS-PAGE)

Preservative: 0.05% Sodium Azide

Concentration: 1.0 mg/ml

Purification: Affinity Chromatography using epitope-specific immunogen

Conjugation: Unconjugated

Storage: Store undiluted at 2-8°C for one month or (in aliquots) at -20°C for longer.

Avoid repeated freezing and thawing.

Stability: Shelf life: one year from despatch.

Predicted Protein Size: ~157 kDa

Gene Name: zinc finger E-box binding homeobox 2

Database Link: Entrez Gene 24136 MouseEntrez Gene 9839 Human

O60315

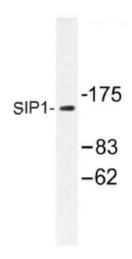


Background:

SMAD regulates gene expression by interacting with different classes of transcription factors including DNA-binding multi-zinc finger proteins. SIP1, for SMAD interacting protein 1, is a member of the delta-EF1/Zfh1 family of 2-handed zinc finger/homeodomain proteins. SIP1 contains a SMAD-binding domain, a homeodomain and two clusters of zinc fingers on the Nand C-termini. SIP1, also known as SMADIP1 and ZFHX1B, can be induced by TGF beta treatment. SIP1 plays a crucial role in normal embryonic development of neural structures and the neural crest. The human SIP1 gene maps to chromosome 2g22. Mutations in the SIP1 gene cause a form of Hirschsprung disease (HSCR). Patients with SIP1 mutations show mental retardation, delayed motor development, epilepsy, microcephaly, distinct facial features and/or congenital heart disease-all symptoms of HSCR.

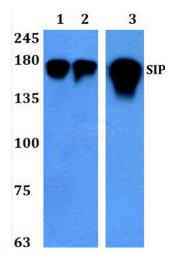
Synonyms: ZFX1B, ZEB2, SMADIP1, SIP1, KIAA0569 **Protein Families:** Druggable Genome, Transcription Factors

Product images:

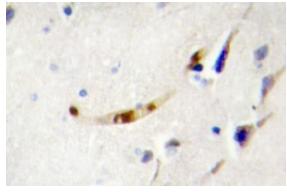


Western blot analysis of SIP1 Antibody at 1/500 dilution: Lane 1: Hela whole cell lysate. Lane 2: HEK293T whole cell lysate. Lane 3: NIH-3T3 whole cell lysate.

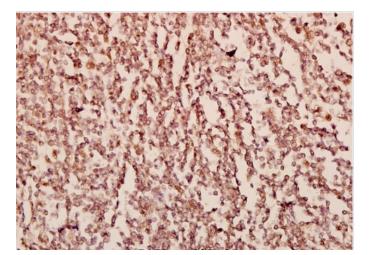




Western blot analysis of SIP1 antibody in extracts from HepG2 cells.



Immunohistochemistry analysis of SIP1 Antibody in paraffin-embedded human breast carcinoma tissue at 1/50.



Immunohistochemistry analysis of SIP1 antibody in paraffin-embedded human brain tissue