

SIP1 (E101) polyclonal antibody

Catalog: BS3009

Host: Rabbit

Reactivity: Human, Mouse

BackGround:

SIP1, also known as SMADIP1 and ZFH1B, can be induced by TGF β treatment. SIP1 plays a crucial role in normal embryonic development of neural structures and the neural crest. The human SIP1 gene maps to chromosome 2q22. 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.

Product:

Rabbit IgG, 1mg/ml in PBS with 0.02% sodium azide, 50% glycerol, pH7.2

Molecular Weight:

~ 157 kDa

Swiss-Prot:

O60315

Purification&Purity:

The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen and the purity is > 95% (by SDS-PAGE).

Applications:

WB: 1:500~1:1000

IHC: 1:50~1:200

Storage&Stability:

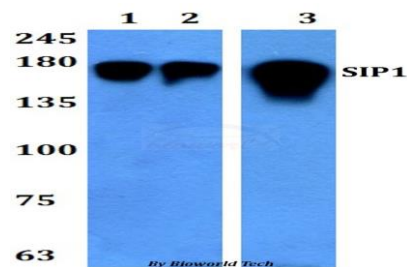
Store at 4 °C short term. Aliquot and store at -20 °C long term. Avoid freeze-thaw cycles.

Specificity:

SIP1 (E101)pAb detects endogenous levels of SIP1 pro-

tein.

DATA:

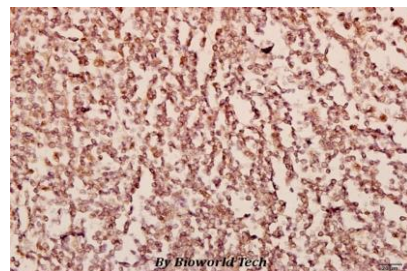


Western blot (WB) analysis of SIP1 (E101) polyclonal antibody at 1:500 dilution

Lane1:Hela whole cell lysate

Lane2:HEK293T whole cell lysate

Lane3:NIH-3T3 whole cell lysate



Immunohistochemistry (IHC) analyzes of SIP1 (E101) pAb in paraffin-embedded human breast carcinoma tissue at 1:50.

Note:

For research use only, not for use in diagnostic procedure.

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