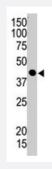
TSG101 polyclonal antibody

Catalog # PAB1758 Size 400 uL

Applications



Western Blot (Cell lysate)

The TSG101 polyclonal antibody (Cat # PAB1758) is used in Western blot to detect TSG101 in ZR-75-1 cell lysate

Specification	
Product Description	Rabbit polyclonal antibody raised against synthetic peptide of TSG101.
Immunogen	A synthetic peptide (conjugated with KLH) corresponding to C-terminus of human TSG101.
Host	Rabbit
Reactivity	Human
Form	Liquid
Purification	Protein G purification
Recommend Usage	Western Blot (1:1000) The optimal working dilution should be determined by the end user.
Storage Buffer	In PBS (0.09% sodium azide)
Storage Instruction	Store at 4°C. For long term storage store at -20°C. Aliquot to avoid repeated freezing and thawing.
Note	This product contains sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which shoul d be handled by trained staff only.

Applications

• Western Blot (Cell lysate)

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Gene Info — TSG101	
Entrez GenelD	<u>7251</u>
Protein Accession#	T101_HUMAN
Gene Name	TSG101
Gene Alias	TSG10, VPS23
Gene Description	tumor susceptibility gene 101
Omim ID	<u>601387</u>
Gene Ontology	Hyperlink
Gene Summary	The protein encoded by this gene belongs to a group of apparently inactive homologs of ubiquitin- conjugating enzymes. The gene product contains a coiled-coil domain that interacts with stathmin, a cytosolic phosphoprotein implicated in tumorigenesis. The protein may play a role in cell growth and differentiation and act as a negative growth regulator. In vitro steady-state expression of this t umor susceptibility gene appears to be important for maintenance of genomic stability and cell cy cle regulation. Mutations and alternative splicing in this gene occur in high frequency in breast can cer and suggest that defects occur during breast cancer tumorigenesis and/or progression. [provi ded by RefSeq
Other Designations	tumor susceptibility protein

Publication Reference

<u>Metabolic characterization of a woman homozygous for the Ser113Leu missense mutation in carnitine</u> palmitoyl transferase II.

Haap M, Thamer C, Machann J, Tschritter O, Loblein K, Kellerer M, Schick F, Jacob S, Haring HU, Stumvoll M. The Journal of Clinical Endocrinology and Metabolism 2002 May; 87(5):2139.

• <u>A novel nonsense mutation (515del4) in muscle carnitine palmitoyltransferase II deficiency.</u>

Deschauer M, Wieser T, Schroder R, Zierz S.

Molecular Genetics and Metabolism 2002 Feb; 75(2):181.

😵 Abnova

Product Information

 Human liver mitochondrial carnitine palmitoyltransferase I: characterization of its cDNA and chromosomal localization and partial analysis of the gene.
Britton CH, Schultz RA, Zhang B, Esser V, Foster DW, McGarry JD.

PNAS 1995 Mar; 92(6):1984.

Pathway

• Endocytosis

Disease

- <u>Genetic Predisposition to Disease</u>
- Hepatitis C
- Lung Neoplasms
- Pulmonary Disease
- Urinary Bladder Neoplasms
- Werner syndrome