

Product datasheet

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ARG65128 anti-SPARTIN antibody

Package: 100 μg Store at: -20°C

Summary

Product Description Goat Polyclonal antibody recognizes SPARTIN

Tested Reactivity Hu

Tested Application ICC/IF, WB

Host Goat

Clonality Polyclonal

Isotype IgG

Target Name SPARTIN

Species Human

Immunogen C-EASGTDVKQLDQGNK

Conjugation Un-conjugated

Alternate Names Spastic paraplegia 20 protein; SPARTIN; Trans-activated by hepatitis C virus core protein 1; Spartin;

TAHCCP1

Application Instructions

Application table	Application	Dilution
	ICC/IF	Assay - dependent
	WB	1 - 3 μg/ml
P.P. STATE	WB: Recommend incubate at RT for 1h. * The dilutions indicate recommended starting dilutions and the optimal dilutions or concentrations should be determined by the scientist.	

Properties

Form Liquid

Purification Purified from goat serum by antigen affinity chromatography.

Buffer Tris saline (pH 7.3), 0.02% Sodium azide and 0.5% BSA.

Preservative 0.02% Sodium azide

Stabilizer 0.5% BSA

Concentration 0.5 mg/ml

Storage instruction For continuous use, store undiluted antibody at 2-8°C for up to a week. For long-term storage, aliquot

and store at -20°C or below. Storage in frost free freezers is not recommended. Avoid repeated freeze/thaw cycles. Suggest spin the vial prior to opening. The antibody solution should be gently mixed

before use.

Note For laboratory research only, not for drug, diagnostic or other use.

Bioinformation

Database links <u>GeneID: 23111 Human</u>

Swiss-port # Q8N0X7 Human

Background This gene encodes a protein containing a MIT (Microtubule Interacting and Trafficking molecule)

domain, and is implicated in regulating endosomal trafficking and mitochondria function. The protein localizes to mitochondria and partially co-localizes with microtubules. Stimulation with epidermal growth factor (EGF) results in protein translocation to the plasma membrane, and the protein functions in the degradation and intracellular trafficking of EGF receptor. Multiple alternatively spliced variants, encoding the same protein, have been identified. Mutations associated with this gene cause autosomal

recessive spastic paraplegia 20 (Troyer syndrome). [provided by RefSeq, Nov 2008]

Research Area Cell Biology and Cellular Response antibody; Gene Regulation antibody

Calculated Mw 73 kDa

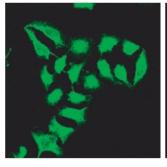
PTM Ubiquitinated; ubiquitination does not require ITCH and WWP1.

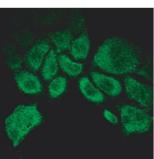
Images

98 kDa 64 kDa 50 kDa 36 kDa

ARG65128 anti-SPARTIN antibody WB image

Western Blot: HeLa lysate (35 μ g protein in RIPA buffer) stained with ARG65128 anti-SPARTIN antibody at 2 μ g/ml dilution.





ARG65128 anti-SPARTIN antibody ICC/IF image

Immunofluorescence: methanol-fixed HeLa stained with ARG65128 anti-SPARTIN antibody ($2\mu g/ml$) before (left) and after (right) introduction of siRNA.