

## ARG58147 anti-Stathmin 1 antibody

Package: 100 µl  
Store at: -20°C

### Summary

Product Description	Rabbit Polyclonal antibody recognizes Stathmin 1
Tested Reactivity	Hu, Ms, Rat
Tested Application	FACS, ICC/IF, IHC-P, IP, WB
Host	Rabbit
Clonality	Polyclonal
Isotype	IgG
Target Name	Stathmin 1
Species	Human
Immunogen	Synthetic peptide derived from Human Stathmin 1.
Conjugation	Un-conjugated
Alternate Names	EC 1.1.1.n12; Multifunctional protein 2; Peroxisomal multifunctional enzyme type 2; SDR8C1; EC 4.2.1.107; 17-beta-HSD 4; 3R; MFE-2; PRLTS1; 3-alpha,7-alpha,12-alpha-trihydroxy-5-beta-cholest-24-enoyl-CoA hydratase; Short chain dehydrogenase/reductase family 8C member 1; 17-beta-hydroxysteroid dehydrogenase 4; DBP; MPF-2; EC 4.2.1.119; D-bifunctional protein

### Application Instructions

Application table	Application	Dilution
	FACS	
	ICC/IF	1:50 - 1:100
	IHC-P	1:50 - 1:100
	IP	1:50
	WB	1:500 - 1:2000
Application Note	* The dilutions indicate recommended starting dilutions and the optimal dilutions or concentrations should be determined by the scientist.	
Positive Control	PC-12	

### Properties

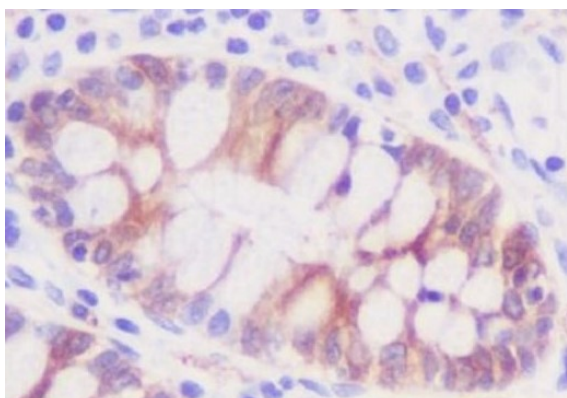
Form	Liquid
Purification	Affinity purified.
Buffer	PBS (pH 7.4), 0.02% Sodium azide and 50% Glycerol.
Preservative	0.02% Sodium azide
Stabilizer	50% Glycerol

<b>Storage instruction</b>	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. 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.
<b>Note</b>	For laboratory research only, not for drug, diagnostic or other use.

## Bioinformation

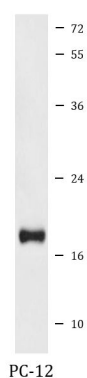
<b>Gene Symbol</b>	HSD17B4
<b>Gene Full Name</b>	hydroxysteroid (17-beta) dehydrogenase 4
<b>Background</b>	The protein encoded by this gene is a bifunctional enzyme that is involved in the peroxisomal beta-oxidation pathway for fatty acids. It also acts as a catalyst for the formation of 3-ketoacyl-CoA intermediates from both straight-chain and 2-methyl-branched-chain fatty acids. Defects in this gene that affect the peroxisomal fatty acid beta-oxidation activity are a cause of D-bifunctional protein deficiency (DBPD). An apparent pseudogene of this gene is present on chromosome 8. Multiple alternatively spliced transcript variants encoding distinct isoforms have been found for this gene. [provided by RefSeq, May 2014]
<b>Function</b>	Bifunctional enzyme acting on the peroxisomal beta-oxidation pathway for fatty acids. Catalyzes the formation of 3-ketoacyl-CoA intermediates from both straight-chain and 2-methyl-branched-chain fatty acids. [UniProt]
<b>Calculated Mw</b>	80 kDa
<b>Cellular Localization</b>	Cytoplasm > Cytoskeleton. [UniProt]

## Images



ARG58147 anti-Stathmin 1 antibody IHC-P image

Immunohistochemistry: Paraffin-embedded Human colon stained with ARG58147 anti-Stathmin 1 antibody.



ARG58147 anti-Stathmin 1 antibody WB image

Western blot: PC-12 cell lysate stained with ARG58147 anti-Stathmin 1 antibody.