

Product datasheet

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ARG58811 anti-17 beta HSD 4 antibody

Package: 50 μg Store at: -20°C

Summary

Product Description Rabbit Polyclonal antibody recognizes 17 beta HSD 4

Tested Reactivity Hu

Predict Reactivity Ms, Rat, Hm

Tested Application IHC-P

Host Rabbit

Clonality Polyclonal

Isotype IgG

Target Name 17 beta HSD 4

Species Human

Immunogen Synthetic peptide corresponding to aa. 744-761 of Human 17 beta HSD 4 (NIMLSQKLQMILKDYAKL).

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
	IHC-P	0.5 - 1 μg/ml
Application Note	IHC-P: Antigen Retrieval: By heat mediation. * The dilutions indicate recommended starting dilutions and the optimal dilutions or concentrations should be determined by the scientist.	

Properties

Form Liquid

Purification Affinity purification with immunogen.

Buffer 0.9% NaCl, 0.2% Na2HPO4, 0.05% Thimerosal, 0.05% Sodium azide and 5% BSA.

Preservative 0.05% Thimerosal and 0.05% Sodium azide

Stabilizer 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.

Bioinformation

Gene Symbol HSD17B4

Gene Full Name hydroxysteroid (17-beta) dehydrogenase 4

Background 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]

Function Bifunctional enzyme acting on the peroxisomal beta-oxidation pathway for fatty acids. Catalyzes the

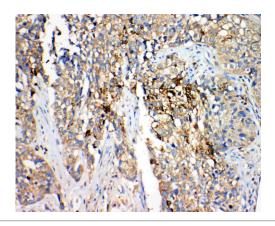
formation of 3-ketoacyl-CoA intermediates from both straight-chain and 2-methyl-branched-chain fatty

acids. [UniProt]

Calculated Mw 80 kDa

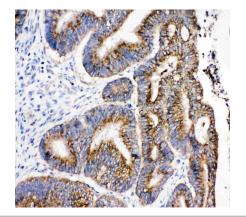
Cellular Localization Peroxisome. [UniProt]

Images



ARG58811 anti-17 beta HSD 4 antibody IHC-P image

Immunohistochemistry: Paraffin-embedded Human lung cancer stained with ARG58811 anti-17 beta HSD 4 antibody.



ARG58811 anti-17 beta HSD 4 antibody IHC-P image

Immunohistochemistry: Paraffin-embedded Human intestinal cancer stained with ARG58811 anti-17 beta HSD 4 antibody.