

ARG58604 anti-EHHADH antibody

Package: 100 μl Store at: -20°C

Summary

| Product Description | Rabbit Polyclonal antibody recognizes EHHADH |
|---------------------|---|
| Tested Reactivity | Hu, Ms, Rat |
| Tested Application | FACS, IHC-P, WB |
| Host | Rabbit |
| Clonality | Polyclonal |
| Isotype | lgG |
| Target Name | EHHADH |
| Species | Human |
| Immunogen | KLH-conjugated synthetic peptide between aa. 662-690 of Human EHHADH. |
| Conjugation | Un-conjugated |
| Alternate Names | LBP; ECHD; LBFP; L-PBE; PBE; PBFE; EC 1.1.1.35; EC 4.2.1.17; EC 5.3.3.8; FRTS3; Peroxisomal bifunctional enzyme |

Application Instructions

| Application table | Application | Dilution |
|-------------------|--|-------------|
| | FACS | 1:10 - 1:50 |
| | IHC-P | 1:10 - 1:50 |
| | WB | 1:1000 |
| Application Note | * The dilutions indicate recommended starting dilutions and the optimal dilutions or concentrations should be determined by the scientist. | |
| Positive Control | Rat liver | |

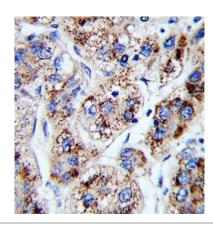
Properties

| Form | Liquid |
|---------------------|---|
| Purification | Purification with Protein A and immunogen peptide. |
| Buffer | PBS and 0.09% (W/V) Sodium azide. |
| Preservative | 0.09% (W/V) Sodium azide |
| 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

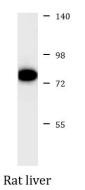
| Gene Symbol | EHHADH |
|-----------------------|---|
| Gene Full Name | enoyl-CoA, hydratase/3-hydroxyacyl CoA dehydrogenase |
| Background | The protein encoded by this gene is a bifunctional enzyme and is one of the four enzymes of the peroxisomal beta-oxidation pathway. The N-terminal region of the encoded protein contains enoyl-CoA hydratase activity while the C-terminal region contains 3-hydroxyacyl-CoA dehydrogenase activity. Defects in this gene are a cause of peroxisomal disorders such as Zellweger syndrome. Two transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, Oct 2009] |
| Calculated Mw | 79 kDa |
| PTM | Acetylated, leading to enhanced enzyme activity. Acetylation is enhanced by up to 80% after treatment either with trichostin A (TSA) or with nicotinamide (NAM) with highest increase on Lys-346. Acetylation and enzyme activity increased by about 1.5% on addition of fatty acids. [UniProt] |
| Cellular Localization | Peroxisome. [UniProt] |

Images



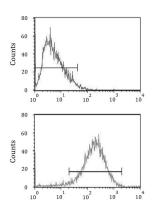
ARG58604 anti-EHHADH antibody IHC-P image

Immunohistochemistry: Formalin-fixed and paraffin-embedded Human hepatocarcinoma tissue stained with ARG58604 anti-EHHADH antibody.



ARG58604 anti-EHHADH antibody WB image

Western blot: 35 μg of Rat liver lysate stained with ARG58604 anti-EHHADH antibody at 1:1000 dilution.



ARG58604 anti-EHHADH antibody FACS image

Flow Cytometry: HepG2 cells stained with ARG58604 anti-EHHADH antibody (bottom histogram) or without primary antibody as control (top histogram), followed by incubation with FITC labelled secondary antibody.