

## **Product datasheet for TP300388**

## OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com

CN: techsupport@origene.cn

## Fumarylacetoacetate hydrolase (FAH) (NM\_000137) Human Recombinant Protein

**Product data:** 

**Product Type:** Recombinant Proteins

**Description:** Recombinant protein of human fumarylacetoacetate hydrolase (fumarylacetoacetase) (FAH), 20

με

Species: Human
Expression Host: HEK293T

**Expression cDNA** >RC200388 protein sequence Clone or AA Sequence: Red=Cloning site Green=Tags(s)

MSFIPVAEDSDFPIHNLPYGVFSTRGDPRPRIGVAIGDQILDLSIIKHLFTGPVLSKHQDVFNQPTLNSF MGLGQAAWKEARVFLQNLLSVSQARLRDDTELRKCAFISQASATMHLPATIGDYTDFYSSRQHATNVGIM FRDKENALMPNWLHLPVGYHGRASSVVVSGTPIRRPMGQMKPDDSKPPVYGACKLLDMELEMAFFVGPGN RLGEPIPISKAHEHIFGMVLMNDWSARDIQKWEYVPLGPFLGKSFGTTVSPWVVPMDALMPFAVPNPKQD PRPLPYLCHDEPYTFDINLSVNLKGEGMSQAATICKSNFKYMYWTMLQQLTHHSVNGCNLRPGDLLASGT ISGPEPENFGSMLELSWKGTKPIDLGNGQTRKFLLDGDEVIITGYCQGDGYRIGFGQCAGKVLPALLPS

**TRTRPLEQKLISEEDLAANDILDYKDDDDKV** 

Tag: C-Myc/DDK
Predicted MW: 46.2 kDa

**Concentration:** >0.05 μg/μL as determined by microplate BCA method

**Purity:** > 80% as determined by SDS-PAGE and Coomassie blue staining

**Buffer:** 25 mM Tris-HCl, 100 mM glycine, pH 7.3, 10% glycerol

**Preparation:** Recombinant protein was captured through anti-DDK affinity column followed by conventional

chromatography steps.

**Note:** For testing in cell culture applications, please filter before use. Note that you may experience

some loss of protein during the filtration process.

Storage: Store at -80°C.

Stability: Stable for 12 months from the date of receipt of the product under proper storage and handling

conditions. Avoid repeated freeze-thaw cycles.

**RefSeq:** NP 000128





**Locus ID:** 2184

UniProt ID: <u>P16930</u>, <u>A0A384P5L6</u>

RefSeq Size: 1810 Cytogenetics: 15q25.1 RefSeq ORF: 1257

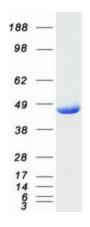
**Summary:** This gene encodes the last enzyme in the tyrosine catabolism pathway. FAH deficiency is

associated with Type 1 hereditary tyrosinemia (HT). [provided by RefSeq, Jul 2008]

**Protein Families:** Druggable Genome

**Protein Pathways:** Metabolic pathways, Tyrosine metabolism

## **Product images:**



Coomassie blue staining of purified FAH protein (Cat# TP300388). The protein was produced from HEK293T cells transfected with FAH cDNA clone (Cat# [RC200388]) using MegaTran 2.0 (Cat# [TT210002]).