**Proteins** 

**Product** Data Sheet



# Fumarylacetoacetase/FAH Protein, Human (HEK293, His)

Cat. No.: HY-P70307

Synonyms: rHuFumarylacetoacetase/FAH, His; Fumarylacetoacetase; FAA; Beta-Diketonase;

Fumarylacetoacetate Hydrolase; FAH

Species: Human Source: HEK293

Accession: P16930 (S2-S419)

Gene ID: 2184

Molecular Weight: Approximately 43.0 kDa

## **PROPERTIES**

AA Sequence					
	SFIPVAEDSD	FPIHNLPYGV	FSTRGDPRPR	IGVAIGDQIL	
	DLSIIKHLFT	GPVLSKHQDV	FNQPTLNSFM	GLGQAAWKEA	
	RVFLQNLLSV	SQARLRDDTE	LRKCAFISQA	SATMHLPATI	
	GDYTDFYSSR	QHATNVGIMF	RDKENALMPN	WLHLPVGYHG	
	RASSVVVSGT	PIRRPMGQMK	PDDSKPPVYG	ACKLLDMELE	
	MAFFVGPGNR	LGEPIPISKA	HEHIFGMVLM	NDWSARDIQK	
	WEYVPLGPFL	GKSFGTTVSP	WVVPMDALMP	FAVPNPKQDP	
	RPLPYLCHDE	PYTFDINLSV	NLKGEGMSQA	ATICKSNFKY	
	MYWTMLQQLT	HHSVNGCNLR	PGDLLASGTI	SGPEPENFGS	
	MLELSWKGTK	PIDLGNGQTR	KFLLDGDEVI	ITGYCQGDGY	
	RIGFGQCAGK	VLPALLPS			
Biological Activity	The enzyme activity of this recombinant protein is testing in progress, we cannot offer a guarantee yet.				
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Appearance	Lyophilized powder.				
Formulation	Lyophilized from a 0.2 μm filtered solution of 20 mM Tris-HCl, 150 mM NaCl, pH 8.5.				
Endotoxin Level	<1 FII/ug_determined by IAI method				
Endotoxin Level	<1 EU/μg, determined by LAL method.				
Reconsititution	tonsititution It is not recommended to reconstitute to a concentration less than 100 μg/mL in ddH <sub>2</sub> O. For long term storage it is				
Reconstitution	recommended to add a carrier protein (0.1% BSA, 5% HSA, 10% FBS or 5% Trehalose).				
		p. e.e (0.12 / 0 20/ 1, 0 / 0	, 2070. 20 0. 070		
Storage & Stability	Stored at -20°C for 2 years.	After reconstitution, it is sta	able at 4°C for 1 week or -20°	°C for longer (with carrier protein). It is	
,	recommended to freeze aliquots at -20°C or -80°C for extended storage.				
		1			
Shipping	Room temperature in conti	nental US; may vary elsewl	here.		

## **DESCRIPTION**

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#### Background

Fumarylacetoacetase is an enzyme encoded by the FAH gene on human chromosome 15. The FAH gene is believed to be involved in the catabolism of phenylalanine in humans and is a key enzyme in the metabolism of phenylalanine and tyrosine, which is mainly expressed in the liver. FAH is a protein homodimer that splits the carbon-carbon bond of fumarylacetic acid in a hydrolysis reaction. The active site of FAH contained  $Ca^{2+}$ . Deficiency of FAH causes hereditary tyrosinemia type  $1^{[1][2][3]}$ .

Caution: Product has not been fully validated for medical applications. For research use only.

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