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CYP2J2 Protein (AA 1-502) (Strep Tag)



Overview

Quantity:	1 mg
Target:	CYP2J2
Protein Characteristics:	AA 1-502
Origin:	Human
Source:	Tobacco (Nicotiana tabacum)
Protein Type:	Recombinant
Purification tag / Conjugate:	This CYP2J2 protein is labelled with Strep Tag.
Application:	ELISA, Western Blotting (WB), SDS-PAGE (SDS)

Product Details

Sequence:

MLAAMGSLAA ALWAVVHPRT LLLGTVAFLL AADFLKRRP KNYPPGPWRL PFLGNFFLVD FEQSHLEVQL FVKKYGNLFS LELGDISAVL ITGLPLIKEA LIHMDQNFGN RPVTPMREHI FKKNGLIMSS GQAWKEQRRF TLTALRNFGL GKKSLEERIQ EEAQHLTEAI KEENGQPFDP HFKINNAVSN IICSITFGER FEYQDSWFQQ LLKLLDEVTY LEASKTCQLY NVFPWIMKFL PGPHQTLFSN WKKLKLFVSH MIDKHRKDWN PAETRDFIDA YLKEMSKHTG NPTSSFHEEN LICSTLDLFF AGTETTSTTL RWALLYMALY PEIQEKVQAE IDRVIGQGQQ PSTAARESMP YTNAVIHEVQ RMGNIIPLNV PREVTVDTTL AGYHLPKGTM ILTNLTALHR DPTEWATPDT FNPDHFLENG QFKKREAFMP FSIGKRACLG EQLARTELFI FFTSLMQKFT FRPPNNEKLS LKFRMGITIS PVSHRLCAVP QV

Sequence without tag. The proposed Strep-Tag is based on experience s with the expression system, a different complexity of the protein could make another tag necessary. In case you have a special request, please contact us.

Characteristics:

Key Benefits:

- Made in Germany from design to production by highly experienced protein experts.
- Protein expressed with ALiCE® and purified by multi-step, protein-specific process to ensure correct folding and modification.
- These proteins are normally active (enzymatically functional) as our customers have reported (not tested by us and not guaranteed).
- State-of-the-art algorithm used for plasmid design (Gene synthesis).

This protein is a **made-to-order protein** and will be made for the first time for your order. Our experts in the lab will ensure that you receive a correctly folded protein.

The big advantage of ordering our **made-to-order proteins** in comparison to ordering custom made proteins from other companies is that there is no financial obligation in case the protein cannot be expressed or purified.

Expression System:

- ALiCE®, our Almost Living Cell-Free Expression System is based on a lysate obtained from Nicotiana tabacum c.v.. This contains all the protein expression machinery needed to produce even the most difficult-to-express proteins, including those that require posttranslational modifications.
- During lysate production, the cell wall and other cellular components that are not required for
 protein production are removed, leaving only the protein production machinery and the
 mitochondria to drive the reaction. During our lysate completion steps, the additional
 components needed for protein production (amino acids, cofactors, etc.) are added to
 produce something that functions like a cell, but without the constraints of a living system all that's needed is the DNA that codes for the desired protein!

Concentration:

- The concentration of our recombinant proteins is measured using the absorbance at 280nm.
- The protein's absorbance will be measured in several dilutions and is measured against its specific reference buffer.
- We use the Expasy's ProtParam tool to determine the absorption coefficient of each protein.

Purification:

Two step purification of proteins expressed in Almost Living Cell-Free Expression System (ALiCE®):

- 1. In a first purification step, the protein is purified from the cleared cell lysate using StrepTag capture material. Eluate fractions are analyzed by SDS-PAGE.
- 2. Protein containing fractions of the best purification are subjected to second purification step through size exclusion chromatography. Eluate fractions are analyzed by SDS-PAGE and

Product Details		
	Western blot.	
Purity:	>80 % as determined by SDS PAGE, Size Exclusion Chromatography and Western Blot.	
Endotoxin Level:	Low Endotoxin less than 1 EU/mg (< 0.1 ng/mg)	
Target Details		
Target:	CYP2J2	
Alternative Name:	CYP2J2 (CYP2J2 Products)	
Background:	Cytochrome P450 2J2 (EC 1.14.14) (Albendazole monooxygenase (hydroxylating)) (EC	
	1.14.14.74) (Albendazole monooxygenase (sulfoxide-forming)) (EC 1.14.14.73) (Arachidonic	
	acid epoxygenase) (CYPIIJ2) (Hydroperoxy icosatetraenoate isomerase) (EC	
	5.4.4.7), FUNCTION: A cytochrome P450 monooxygenase involved in the metabolism of	
	polyunsaturated fatty acids (PUFA) in the cardiovascular system (PubMed:8631948,	
	PubMed:19965576). Mechanistically, uses molecular oxygen inserting one oxygen atom into a	
	substrate, and reducing the second into a water molecule, with two electrons provided by	
	NADPH via cytochrome P450 reductase (NADPHhemoprotein reductase) (PubMed:8631948,	
	PubMed:19965576). Catalyzes the epoxidation of double bonds of PUFA (PubMed:8631948,	
	PubMed:19965576). Converts arachidonic acid to four regioisomeric epoxyeicosatrienoic acids	
	(EpETrE), likely playing a major role in the epoxidation of endogenous cardiac arachidonic acid	
	pools (PubMed:8631948). In endothelial cells, participates in eicosanoids metabolism by	
	converting hydroperoxide species into hydroxy epoxy metabolites. In combination with 15-	
	lipoxygenase metabolizes arachidonic acid and converts hydroperoxyicosatetraenoates	
	(HpETEs) into hydroxy epoxy eicosatrienoates (HEETs), which are precursors of vasodilatory	
	trihydroxyicosatrienoic acids (THETAs). This hydroperoxide isomerase activity is NADPH- and	
	O2-independent (PubMed:19737933). Catalyzes the monooxygenation of a various xenobiotics,	
	such as danazol, amiodarone, terfenadine, astemizole, thioridazine, tamoxifen, cyclosporin A	
	and nabumetone (PubMed:19923256). Catalyzes hydroxylation of the anthelmintics	
	albendazole and fenbendazole (PubMed:23959307). Catalyzes the sulfoxidation of fenbedazole	
	(PubMed:19923256). {ECO:0000269 PubMed:19737933, ECO:0000269 PubMed:19923256,	
	ECO:0000269 PubMed:19965576, ECO:0000269 PubMed:23959307,	
	ECO:0000269 PubMed:8631948}.	
Molecular Weight:	57.6 kDa	

UniProt:

P51589

Application Details

Application Notes:	In addition to the applications listed above we expect the protein to work for functional studies as well. As the protein has not been tested for functional studies yet we cannot offer a guarantee though.
Comment:	ALiCE®, our Almost Living Cell-Free Expression System is based on a lysate obtained from Nicotiana tabacum c.v This contains all the protein expression machinery needed to produce even the most difficult-to-express proteins, including those that require post-translational modifications.
	During lysate production, the cell wall and other cellular components that are not required for protein production are removed, leaving only the protein production machinery and the mitochondria to drive the reaction. During our lysate completion steps, the additional components needed for protein production (amino acids, cofactors, etc.) are added to produce something that functions like a cell, but without the constraints of a living system - all that's needed is the DNA that codes for the desired protein!
Restrictions:	For Research Use only
Handling	
Format:	Liquid
Buffer:	The buffer composition is at the discretion of the manufacturer. If you have a special request, please contact us.
Handling Advice:	Avoid repeated freeze-thaw cycles.
Storage:	-80 °C
Storage Comment:	Store at -80°C.
Expiry Date:	Unlimited (if stored properly)