

Recombinant Human AACs/Acetoacetyl-CoA Synthetase Protein (His Tag)



Catalog Number:PKSH031250

Note: Centrifuge before opening to ensure complete recovery of vial contents.

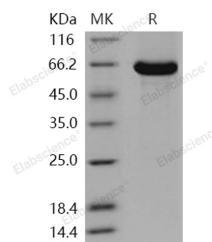
Description

Synonyms	ACSF1;SUR-5
Species	Human
Expression Host	Baculovirus-Insect Cells
Sequence	Met 1-Phe 672
Accession	NP_076417.2
Calculated Molecular Weight	77.0 kDa
Observed molecular weight	60 kDa
Tag	N-His

Properties

Purity	> 96 % as determined by reducing SDS-PAGE.
Endotoxin	< 1.0 EU per μ g of the protein as determined by the LAL method.
Storage	Generally, lyophilized proteins are stable for up to 12 months when stored at -20 to -80°C. Reconstituted protein solution can be stored at 4-8°C for 2-7 days. Aliquots of reconstituted samples are stable at < -20°C for 3 months.
Shipping	This product is provided as lyophilized powder which is shipped with ice packs.
Formulation	Lyophilized from sterile 20mM Tris, 500mM NaCl, pH 7.4 Normally 5 % - 8 % trehalose, mannitol and 0.01% Tween80 are added as protectants before lyophilization. Please refer to the specific buffer information in the printed manual.
Reconstitution	Please refer to the printed manual for detailed information.

Data



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Background

Acetoacetyl-CoA Synthetase (AACs) is a novel cytosolic ketone body (acetoacetate)-specific ligase. The AACs in adipose tissue plays an important role in utilizing ketone body for the fatty acid-synthesis during adipose tissue development. It had been improved that Acetoacetyl-CoA Synthetase is an essential enzyme for the synthesis of fatty acid and cholesterol from ketone bodies, was found to be highly expressed in mouse adipose tissue, and GC box and C/EBPs motif were crucial for AACs promoter activity in 3T3-L1 adipocytes. Moreover, AACs promoter activity was controlled mainly by C/EBPalpha during adipogenesis. AACs gene expression is particularly abundant in white adipose tissue, as it is induced during adipocyte differentiation. The human AACs promoter is a PPARgamma target gene and that this nuclear receptor is recruited to the AACs promoter by direct interaction with Sp1 (stimulating protein-1). The

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Acetoacetyl-CoA Synthetase has important roles in the regulation of ketone body utilization in rat liver and that these hypocholesterolemic agents have the ability to remedy the impaired utilization of ketone bodies under the diabetic condition.

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