

Product datasheet

Recombinant Human HGFA Inhibitor 2 protein
ab153796

1 Image

Description

Product name	Recombinant Human HGFA Inhibitor 2 protein	
Purity	> 95 % SDS-PAGE. Greater than 95% as determined by SEC-HPLC and reducing SDS-PAGE. Lyophilized from a 0.2 µM filtered solution.	
Endotoxin level	< 1.000 Eu/µg	
Expression system	HEK 293 cells	
Accession	O43291	
Protein length	Protein fragment	
Animal free	No	
Nature	Recombinant	
Species	Human	
Sequence	ADRERSIHDFCLVSKVVGRCRASMPRWWYNVTDGSCQL FVYGGCDGNSNN YLTKEECLKKCATVTENATGDLATSRNAADSSVPSAPRR QDSEDHSSDMF NYEEYCTANAVTGPCRASFPRWYFDVERNSCNNFIYGGC RGNKNSYRSEE ACMLRCFRQQENPPLPLGSKVDHHHHHH	
Predicted molecular weight	20 kDa including tags	
Amino acids	28 to 197	
Tags	His tag C-Terminus	

Specifications

Our [Abpromise guarantee](#) covers the use of **ab153796** in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Applications	SDS-PAGE HPLC
Form	Lyophilized

Additional notes

Reconstituted protein solution can be stored at 4-7°C for 2-7 days. Aliquots of reconstituted samples are stable at < -20°C for 3 months.

Preparation and Storage

Stability and Storage

Shipped at Room Temperature. The lyophilized protein is stable for a few weeks at room temperature. Store at -20°C. Please see notes section.

pH: 7.40

Constituents: 99% Phosphate Buffer, 0.88% Sodium chloride

Reconstitution

Always centrifuge tubes before opening. Do not mix by vortex or pipetting. It is not recommended to reconstitute to a concentration less than 100 µg/ml. Dissolve the lyophilized protein in 1X PBS. Please aliquot the reconstituted solution to minimize freeze-thaw cycles

General Info

Function

Inhibitor of HGF activator. Also inhibits plasmin, plasma and tissue kallikrein, and factor XIa.

Tissue specificity

Expressed in placenta, kidney, pancreas, prostate, testis, thymus, and trachea.

Involvement in disease

Defects in SPINT2 are the cause of diarrhea type 3 (DIAR3) [MIM:270420]; also known as congenital sodium diarrhea (CSD). DIAR3 is a rare, inherited diarrhea of infancy. A diagnosis of DIAR3 is made on the findings of a life-threatening secretory diarrhea, severe metabolic acidosis, and hyponatremia secondary to extraordinarily high fecal losses of sodium, with low or normal excretion of urinary sodium, in the absence of infectious, autoimmune, and endocrine causes.

Sequence similarities

Contains 2 BPTI/Kunitz inhibitor domains.

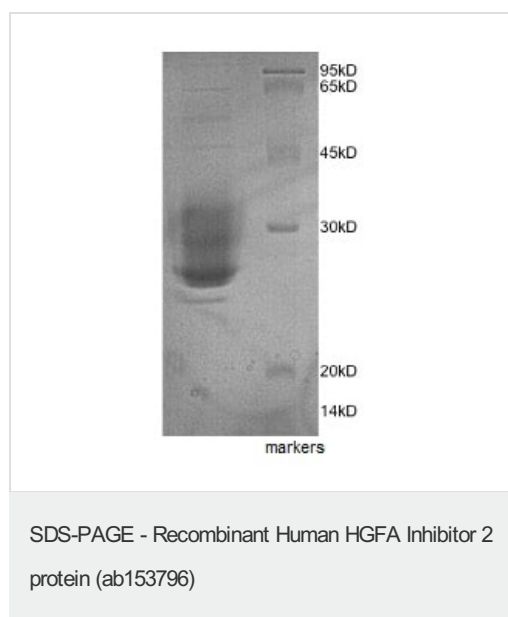
Domain

This inhibitor contains two inhibitory domains.

Cellular localization

Membrane.

Images



SDS-PAGE analysis using ab153796.

Predicted MW 20 kDa.

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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