

Product datasheet

Recombinant human Cripto1/CRIPTO protein ab84064

1 Image

Description	
Product name	Recombinant human Cripto1/CRIPTO protein
Biological activity	ab84064 has been shown to stimulate MAPK phosphorylation in HUVEC cells. 200ng/ml is sufficient to stimulate phosphorylation.
Purity	> 95 % SDS-PAGE.
Expression system	HEK 293 cells
Protein length	Protein fragment
Animal free	No
Nature	Recombinant
Species	Human
Sequence	Theoretical sequence: LGHQEFARPSRGYLAFRDDSWPQEEPAIRPRSSQRVPP MGIQHSKELN RTCCLNGGTCMLGSFCACPPSFYGRNCEHDVRKENCGS VPHDTWLPPKKC SLCKCWHGQLRCFPQAFLPGCDGLVMDEHLVASRTPE LPPS
Amino acids	31 to 169

Specifications	
Our Abpromise guarantee covers the use of ab84064 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
Form	Lyophilized
Additional notes	ab84064 has been shown to stimulate MAPK phosphorylation in HUVEC cells. 200ng/ml is sufficient to stimulate phosphorylation. This product was previously labelled as Cripto1 This product was previously labelled as Cripto1

Preparation and Storage

Stability and Storage

Shipped at 4°C. After reconstitution store at -20°C. Avoid freeze / thaw cycles.

Constituents: 1% Human serum albumin, 10% Trehalose

This product is an active protein and may elicit a biological response in vivo, handle with caution.

Reconstitution

It is recommended that 0.5 ml of sterile phosphate-buffered saline be added to the vial. Following reconstitution short-term storage at 4°C is recommended, with longer-term storage in aliquots at -18 to -20°C. Repeated freeze thawing is not recommended.

General Info

Function

Could play a role in the determination of the epiblastic cells that subsequently give rise to the mesoderm.

Tissue specificity

Preferentially expressed in gastric and colorectal carcinomas than in their normal counterparts.

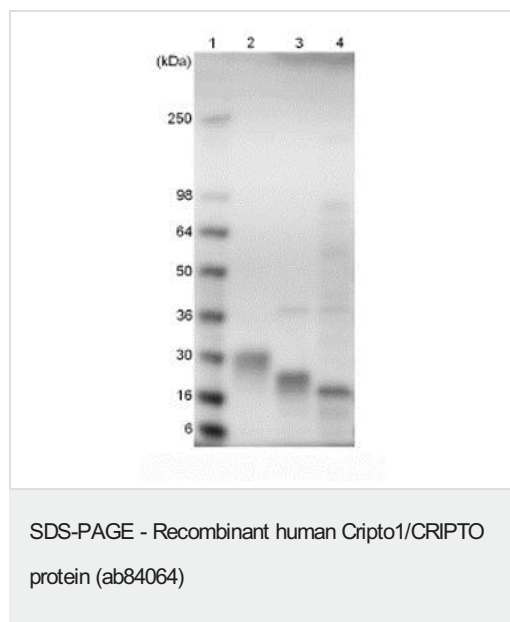
Sequence similarities

Contains 1 EGF-like domain.

Cellular localization

Cell membrane.

Images



Lane 1- MW markers; Lane 2- ab84064 ; Lane 3- ab84064 treated with PNGase F to remove potential N-linked glycans; Lane 4- ab84064 treated with a glycosidase cocktail to remove potential N- and O-linked glycans. Approximately 5 µg of protein was loaded per lane; Gel was stained using Coomassie.

A drop in MW after treatment with PNGase F indicates presence of N-linked glycans. A further drop in MW after treatment with the glycosidase cocktail indicates the presence of O-linked glycans. Additional bands in lane 3 and lane 4 are glycosidase enzymes.

O-fucosylation at Thr-88 has been confirmed.

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

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