

Recombinant human Cripto1/CRIPTO protein ab84064

画像数 1

製品の詳細

製品名	Recombinant human Cripto1/CRIPTO protein
生理活性	ab84064 has been shown to stimulate MAPK phosphorylation in HUVEC cells. 200ng/ml is sufficient to stimulate phosphorylation.
精製度	> 95 % SDS-PAGE.
発現系	HEK 293 cells
タンパク質長	Protein fragment
Animal free	No
由来	Recombinant
生物種	Human
配列	<p>Theoretical sequence:</p> <p>LGHQEFARPSRGYLAFRDDSIWPQEEPAIRPRSSQRVPP</p> <p>MGIQHSKELN</p> <p>RTCCNLNGGTCMLGSFCACPPSFYGRNCEHDVRKENC</p> <p>GSVPHDTWLPKKC</p> <p>SLCKCWHGQLRCFPQAFLPGCDGLVMDEHLVASRTPE LPPS</p>
領域	31 to 169

特性

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.

アプリケーション	SDS-PAGE
製品の状態	Lyophilized
備考	<p>ab84064 has been shown to stimulate MAPK phosphorylation in HUVEC cells. 200ng/ml is sufficient to stimulate phosphorylation.</p> <p>This product was previously labelled as Cripto1</p> <p>This product was previously labelled as Cripto1</p>

前処理および保存

保存方法および安定性

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.

再構成

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.

関連情報

機能

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

組織特異性

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

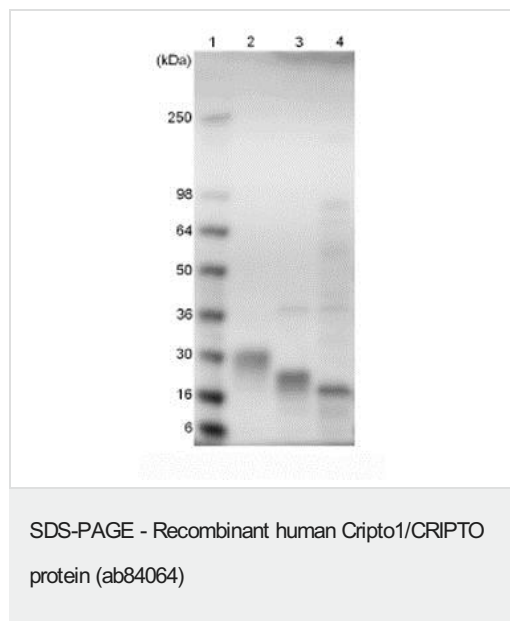
配列類似性

Contains 1 EGF-like domain.

細胞内局在

Cell membrane.

画像



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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