abcam

Product datasheet

Recombinant Human Cytokeratin 5 protein ab132931

1 图像

描述

产品名称 重组人Cytokeratin 5蛋白

纯**度** >= 80 % Purified via GST Tag.

Glutathione Sepharose

表达系统 Wheat germ

Accession P13647

蛋白长度 Full length protein

无动物成分 No

性质 Recombinant

种属 Human

序列 MSRQSSVSFRSGGSRSFSTASAITPSVSRTSFTSVSRSGGGG

GGGFGRVS

LAGACGVGGYGSRSLYNLGGSKRISISTSGGSFRNRFGAGAG

GGYGFGGG

AGSGFGFGGAGGGFGLGGGAGFGGGFGGPGFPVCPPGGIQE

VTVNQSLL

 ${\tt TPLNLQIDPSIQRVRTEEREQIKTLNNKFASFIDKVRFLEQQ}$

NKVLDTKW

TLLQEQGTKTVRQNLEPLFEQYINNLRRQLDSIVGERGRLDS

ELRNMQDL

VEDFKNKYEDEINKRTTAENEFVMLKKDVDAAYMNKVELEAK

VDALMDEI

NFMKMFFDAELSQMQTHVSDTSVVLSMDNNRNLDLDSIIAEV

KAQYEEIA

NRSRTEAESWYQTKYEELQQTAGRHGDDLRNTKHEISEMNRM

IQRLRAEI

DNVKKQCANLQNAIADAEQRGELALKDARNKLAELEEALQKA

KQDMARLL

REYQELMNTKLALDVEIATYRKLLEGEECRLSGEGVGPVNIS

VVTGSVSS

GYGSGSGYGGGLGGGLGGGLAGGGSGSYYSSSSGGVGL

SGGLSVGG

SGFSASSGRGLGVGFGSGGGSSSSVKFVSTTSSSRKSFKS

预测分子量 91 kDa including tags

氨基酸 1 to 590

技术指标

Our Abpromise quarantee covers the use of ab132931 in the following tested applications.

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

应**用** ELISA

Western blot SDS-PAGE

形式 Liquid

补充说明

制备和贮存

稳**定性和存储** Shipped on dry ice. Upon delivery aliquot and store at -80℃. Avoid freeze / thaw cycles.

pH: 8.00

Constituents: 0.31% Glutathione, 0.79% Tris HCI

常规信息

疾病相关

Defects in KRT5 are a cause of epidermolysis bullosa simplex Dowling-Meara type (DM-EBS) [MIM:131760]. DM-EBS is a severe form of intraepidermal epidermolysis bullosa characterized by generalized herpetiform blistering, milia formation, dystrophic nails, and mucous membrane involvement.

Defects in KRT5 are the cause of epidermolysis bullosa simplex with migratory circinate erythema (EBSMCE) [MIM:609352]. EBSMCE is a form of intraepidermal epidermolysis bullosa characterized by unusual migratory circinate erythema. Skin lesions appear from birth primarily on the hands, feet, and legs but spare nails, ocular epithelia and mucosae. Lesions heal with brown pigmentation but no scarring. Electron microscopy findings are distinct from those seen in the DM-EBS, with no evidence of tonofilament clumping.

Defects in KRT5 are a cause of epidermolysis bullosa simplex Weber-Cockayne type (WC-EBS) [MIM:131800]. WC-EBS is a form of intraepidermal epidermolysis bullosa characterized by blistering limited to palmar and plantar areas of the skin.

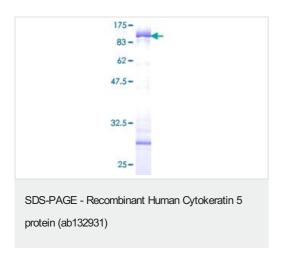
Defects in KRT5 are a cause of epidermolysis bullosa simplex Koebner type (K-EBS) [MIM:131900]. K-EBS is a form of intraepidermal epidermolysis bullosa characterized by generalized skin blistering. The phenotype is not fundamentally distinct from the Dowling-Meara type, althought it is less severe.

Defects in KRT5 are the cause of epidermolysis bullosa simplex with mottled pigmentation (MP-EBS) [MIM:131960]. MP-EBS is a form of intraepidermal epidermolysis bullosa characterized by blistering at acral sites and 'mottled' pigmentation of the trunk and proximal extremities with hyperand hypopigmentation macules.

Defects in KRT5 are the cause of Dowling-Degos disease (DDD) [MIM:179850]; also known as Dowling-Degos-Kitamura disease or reticulate acropigmentation of Kitamura. DDD is an autosomal dominant genodermatosis. Affected individuals develop a postpubertal reticulate hyperpigmentation that is progressive and disfiguring, and small hyperkeratotic dark brown

Belongs to the intermediate filament family.

图片



12.5% SDS-PAGE analysis of ab132931 stained with Coomassie Blue

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