abcam

Product datasheet

Anti-XPB antibody [EPR14161] ab190698

יולצעבע RabMAb

★★★★★ 1 Abreviews 1 References 画像数3

製品の概要

製品名 Anti-XPB antibody [EPR14161]

製品の詳細 Rabbit monoclonal [EPR14161] to XPB

由来種 Rabbit

アプリケーション 適用あり: ICC/IF. WB

種交差性 交差種: Human

交差が予測される動物種: Cow, Cynomolgus monkey, Orangutan 4

免疫原 Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

ポジティブ・コントロール HeLa, 293 and Jurkat whole cell lysate (ab7899), A549 cells

特記事項 This product is a recombinant monoclonal antibody, which offers several advantages including:

- High batch-to-batch consistency and reproducibility

- Improved sensitivity and specificity

- Long-term security of supply

- Animal-free production

For more information see here.

Our RabMAb® technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb® patents**.

製品の特性

製品の状態 Liquid

保存方法 Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C long

term. Avoid freeze / thaw cycle.

バッファー pH: 7.2

Preservative: 0.01% Sodium azide

Constituents: 40% Glycerol (glycerin, glycerine), 0.05% BSA, 59% PBS

精製度 Protein A purified

モノクローナル ポリモノ クローン名 EPR14161

アイソタイプ ΙgG

The Abpromise guarantee <u>Abpromise保証は、</u>次のテスト済みアプリケーションにおけるab190698の使用に適用されます アプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご検討ください。

アプリケーション	Abreviews	特記事項
ICC/IF		1/500.
WB	*** <u>*</u> (1)	1/1000 - 1/10000. Predicted molecular weight: 89 kDa.

ターゲット情報

機能

ATP-dependent 3'-5' DNA helicase, component of the core-TFIIH basal transcription factor, involved in nucleotide excision repair (NER) of DNA and, when complexed to CAK, in RNA transcription by RNA polymerase II. Acts by opening DNA either around the RNA transcription start site or the DNA damage.

関連疾患

Defects in ERCC3 are the cause of xeroderma pigmentosum complementation group B (XP-B) [MIM:610651]; also known as xeroderma pigmentosum II (XP2) or XP group B (XPB) or xeroderma pigmentosum group B combined with Cockayne syndrome (XP-B/CS). Xeroderma pigmentosum is an autosomal recessive pigmentary skin disorder characterized by solar hypersensitivity of the skin, high predisposition for developing cancers on areas exposed to sunlight and, in some cases, neurological abnormalities. Some XP-B patients present features of Cockayne syndrome, including dwarfism, sensorineural deafness, microcephaly, mental retardation, pigmentary retinopathy, ataxia, decreased nerve conduction velocities.

Defects in ERCC3 are a cause of trichothiodystrophy photosensitive (TTDP) [MIM:601675].

TTDP is an autosomal recessive disease characterized by sulfur-deficient brittle hair and nails, ichthyosis, mental retardation, impaired sexual development, abnormal facies and cutaneous photosensitivity correlated with a nucleotide excision repair (NER) defect. Neonates with trichothiodystrophy and ichthyosis are usually born with a collodion membrane. The severity of the ichthyosis after the membrane is shed is variable, ranging from a mild to severe lamellar ichthyotic phenotype. There are no reports of skin cancer associated with TTDP.

配列類似性

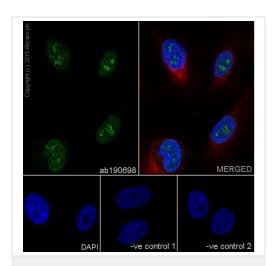
Belongs to the helicase family. RAD25/XPB subfamily. Contains 1 helicase ATP-binding domain.

Contains 1 helicase C-terminal domain.

細胞内局在

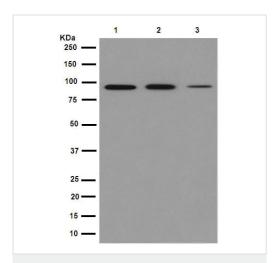
Nucleus.

画像



Immunocytochemistry/ Immunofluorescence - Anti-XPB antibody [EPR14161] (ab190698)

Immunocytochemical analysis of 4% Paraformaldehyde fixed A549 cells, labeling XPB using ab190698 at a 1/500 dilution and Goat anti rabbit lgG (Alexa Fluor® 555) secondary antibody at a 1/200 dilution and counter-stained with Dapi.



Western blot - Anti-XPB antibody [EPR14161] (ab190698)

All lanes : Anti-XPB antibody [EPR14161] (ab190698) at 1/10000 dilution

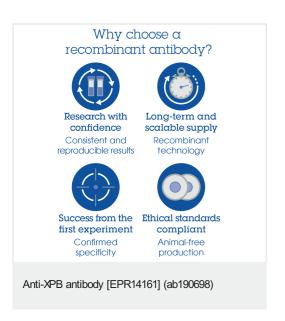
Lane 1 : HeLa cell lysate
Lane 2 : 293 cell lysate
Lane 3 : Jurkat cell lysate

Lysates/proteins at 10 µg per lane.

Secondary

All lanes : Goat Anti-Rabbit lgG, (H+L), Peroxidase conjugated at 1/1000 dilution

Predicted band size: 89 kDa



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