

Anti-XPB antibody [2327C4a] ab70596

1 图像

概述

产品名称	Anti-XPB抗体[2327C4a]
描述	小鼠单克隆抗体[2327C4a] to XPB
宿主	Mouse
经测试应用	适用于: WB, Dot blot
种属反应性	与反应: Human
免疫原	Recombinant fragment (Human) from an N terminal region of XPB.
阳性对照	Recombinant XPB protein.

性能

形式	Liquid
存放说明	Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.
存储溶液	Preservative: 0.05% Sodium Azide Constituents: 1% BSA, PBS, pH 7.4
纯度	Protein G purified
纯化说明	This antibody was purified using protein G column chromatography from culture supernatant of hybridoma cultured in a medium containing bovine IgG-depleted (approximately 95%) fetal bovine serum. It was filtered through a 0.22 µm membrane.
克隆	单克隆
克隆编号	2327C4a
同种型	IgG1

应用

The Abpromise guarantee **Abpromise™** 承诺保证使用ab70596于以下的经测试应用

“应用说明”部分 下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。

应用	Ab评论	说明

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WB		Use at an assay dependent dilution. Predicted molecular weight: 87 kDa.
Dot blot		Use at an assay dependent dilution.

靶标

功能

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.

疾病相关

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.

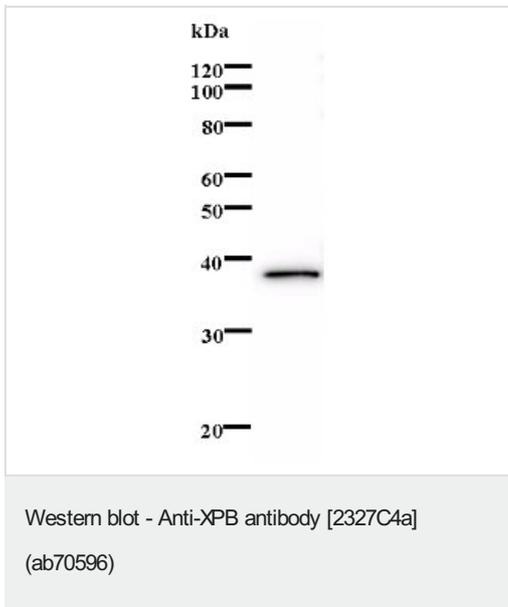
序列相似性

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

细胞定位

Nucleus.

图片



Anti-XPB antibody [2327C4a] (ab70596) + immunising recombinant fragment

Predicted band size: 87 kDa

Observed band size: 38 kDa

The molecular weight of the band on the western blot does not correspond to the molecular weight of the natural protein because only a fragment of the protein was used.

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

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