abcam

Product datasheet

Anti-DLX3 antibody [NCI-R183-14] ab211492



★★★★ 1 Abreviews 2 Images

Overview

Product name Anti-DLX3 antibody [NCI-R183-14]

Description Rabbit monoclonal [NCI-R183-14] to DLX3

Host species Rabbit

Tested applications Suitable for: WB

Species reactivity Reacts with: Human

Predicted to work with: Mouse

Synthetic peptide. This information is proprietary to Abcam and/or its suppliers. **Immunogen**

Positive control WB: Saos-2 cell transfected with mouse TetOFFv5DLX3/flag TDO untreated with

DOX(Doxorubicin 2µg/ml); Saos-2 cell transfected with human TetOFFv5DLX3/flag TDO

untreated with DOX(Doxorubicin 2µg/ml).

General notes 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.

Properties

Form

Storage instructions 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.

Storage buffer pH: 7.2

Preservative: 0.01% Sodium azide

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

Purity Protein A purified

Clonality Monoclonal

Clone number NCI-R183-14

Isotype IgG

Applications

The Abpromise guarantee Our Abpromise guarantee covers the use of ab211492 in the following tested applications.

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

Application	Abreviews	Notes
WB		1/1000. Detects a band of approximately 45, 38 kDa (predicted molecular weight: 32 kDa).

Target

Function Likely to play a regulatory role in the development of the ventral forebrain. May play a role in

craniofacial patterning and morphogenesis.

Involvement in disease Defects in DLX3 are a cause of trichodentoosseous syndrome (TDO) [MIM:190320]. TDO is an

autosomal dominant syndrome characterized by enamel hypoplasia and hypocalcification with

associated strikingly curly hair.

Defects in DLX3 are the cause of amelogenesis imperfecta type 4 (Al4) [MIM:104510]; also known as amelogenesis imperfecta hypomaturation-hypoplastic type with taurodontism. Al4 is an

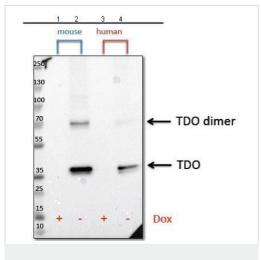
autosomal dominant defect of enamel formation associated with enlarged pulp chambers.

Sequence similarities Belongs to the distal-less homeobox family.

Contains 1 homeobox DNA-binding domain.

Cellular localization Nucleus.

Images



Western blot - Anti-DLX3 antibody [NCI-R183-14] (ab211492)

The data was kindly provided by Dr. Maria Morasso (NIH/NIAMS).

All lanes : Anti-DLX3 antibody [NCI-R183-14] (ab211492) at 1/1000 dilution

Lane 1: Saos-2 (Human osteosarcoma cell line) cell transfected

with mouse TetOFFv5DLX3/flag TDO treated with

DOX(Doxorubicin 2µg/ml)

Lane 2: Saos-2 (Human osteosarcoma cell line) cell transfected

with mouse TetOFFv5DLX3/flag TDO untreated with

DOX(Doxorubicin 2µg/ml)

Lane 3: Saos-2 (Human osteosarcoma cell line) cell transfected

with human TetOFFv5DLX3/flag TDO treated with

DOX(Doxorubicin 2µg/ml)

Lane 4: Saos-2 (Human osteosarcoma cell line) cell transfected

with human TetOFFv5DLX3/flag TDO untreated with

DOX(Doxorubicin 2µg/ml)

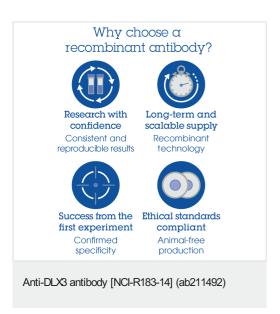
Predicted band size: 32 kDa

Observed band size: 38,45 kDa

Exposure time: 30 seconds

Blocking/Dilution buffer: 5% NFDM/TBST.

This product only recognizes DLX3 carrying the TDO (Tricho-Dento-Osseous syndrome) frameshift mutation.



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

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