SZABO SCANDIC

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Produktinformation

- Forschungsprodukte & Biochemikalien
- Zellkultur & Verbrauchsmaterial
- Diagnostik & molekulare Diagnostik
- Laborgeräte & Service

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See the following pages for more information!

Lieferung & Zahlungsart

siehe unsere Liefer- und Versandbedingungen

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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DLX1 (Human) Matched Antibody Pair

Catalog # : H00001745-AP61 **規格 :** [1 Set]

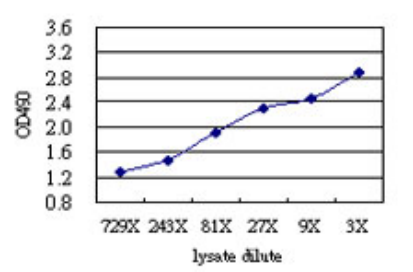
List All

Specification

Product Description: This antibody pair set comes with matched antibody pair to detect and quantify protein level of human DLX1.

Reactivity: Human

Quality Control Testing: Standard curve using DLX1 293T overexpression lysate (non-denatured) as an analyte.



Sandwich ELISA detection sensitivity ranging from approximately 729x to 3x dilution of the DLX1 293T overexpression lysate (non-denatured).

Supplied Product: Antibody pair set content:
1. Capture antibody: mouse monoclonal anti-DLX1 (100 ug)
2. Detection antibody: rabbit MaxPab® affinity purified polyclonal anti-DLX1 (50 ug)
*Reagents are sufficient for at least 3-5 x 96 well plates using recommended protocols.

Storage Instruction: Store reagents of the antibody pair set at -20°C or lower. Please aliquot to avoid repeated freeze thaw cycle. Reagents should be returned to -20°C storage immediately after use.

MSDS: Download

Applications

ELISA Pair (Transfected lysate)

Protocol Download

Gene Information

Entrez GeneID: 1745

Gene Name: DLX1

Gene Alias: -

Gene Description: distal-less homeobox 1

Omim ID: 600029

Gene Ontology: Hyperlink

Application Image

ELISA Pair (Transfected lysate)

Gene Summary: This gene encodes a member of a homeobox transcription factor gene family similiar to the Drosophila distal-less gene. The encoded protein is localized to the nucleus where it may function as a transcriptional regulator of signals from multiple TGF-{beta} superfamily members. The encoded protein may play a role in the control of craniofacial patterning and the differentiation and survival of inhibitory neurons in the forebrain. This gene is located in a tail-to-tail configuration with another member of the family on the long arm of chromosome 2. Alternatively spliced transcript variants encoding different isoforms have been described. [provided by RefSeq

Other Designations: OTTHUMP00000082494,OTTHUMP00000082497,distal-less homeo box 1

Related Disease

Attention Deficit Disorder with Hyperactivity Autistic Disorder Cleft Lip Cleft Palate Genetic Predisposition to Disease NARP Schizophrenia Schizophrenia