SZABO SCANDIC

Part of Europa Biosite

Produktinformation

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

Weitere Information auf den folgenden Seiten!
See the following pages for more information!

Lieferung & Zahlungsart

siehe unsere Liefer- und Versandbedingungen

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

SZABO-SCANDIC HandelsgmbH

Quellenstraße 110, A-1100 Wien

T. +43(0)1 489 3961-0

F. +43(0)1 489 3961-7

mail@szabo-scandic.com

www.szabo-scandic.com

linkedin.com/company/szaboscandic

MSX1 (Human) Matched Antibody Pair

Catalog # : H00004487-AP41 **規格 :** [1 Set]

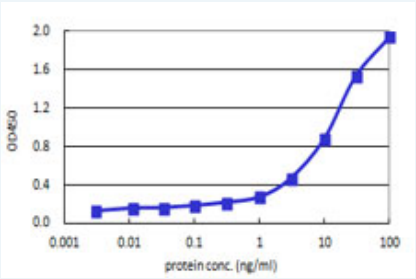
List All

Specification

Product Description: This antibody pair set comes with matched antibody pair to detect and quantify protein level of human MSX1.

Reactivity: Human

Quality Control Testing: Standard curve using recombinant protein (H00004487-Q01) as an analyte.



Sandwich ELISA detection sensitivity ranging from 0.1 ng/ml to 100 ng/ml.

Supplied Product: Antibody pair set content:
1. Capture antibody: mouse monoclonal anti-MSX1, IgG2a Kappa (100 ug)
2. Detection antibody: biotinylated mouse monoclonal anti-MSX1, IgG2a Kappa (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.

Applications

ELISA Pair (Recombinant protein)

Protocol Download

Gene Information

Entrez GeneID: 4487

Gene Name: MSX1

Gene Alias: HOX7,HYD1

Gene Description: msh homeobox 1

Omim ID: 106600, 142983, 189500, 608874

Gene Ontology: Hyperlink

Gene Summary: This gene encodes a member of the muscle segment homeobox gene

Application Image

ELISA Pair (Recombinant protein)

family. The encoded protein functions as a transcriptional repressor during embryogenesis through interactions with components of the core transcription complex and other homeoproteins. It may also have roles in limb-pattern formation, craniofacial development, particularly odontogenesis, and tumor growth inhibition. Mutations in this gene, which was once known as homeobox 7, have been associated with nonsyndromic cleft lip with or without cleft palate 5, Witkop syndrome, Wolf-Hirschom syndrome, and autosomoal dominant hypodontia. [provided by RefSeq

Other Designations: OTTHUMP00000115387,homeobox 7,msh homeo box 1,msh homeobox homolog 1

Related Disease

Breast Neoplasms Cleft Lip Cleft Palate Disease Models, Animal Genetic Predisposition to Disease Multiple System Atrophy Parkinson Disease Parkinson disease Sleep Apnea, Obstructive