SZABO SCANDIC

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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

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MSX1 (Human) IP-WB Antibody Pair

Catalog # : H00004487-PW2 **規格 :** [1 Set]

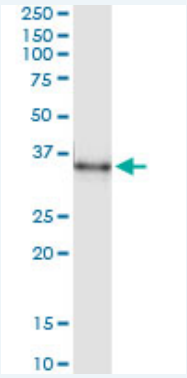
List All

Specification

Product Description: This IP-WB antibody pair set comes with one antibody for immunoprecipitation and another to detect the precipitated protein in western blot.

Reactivity: Human

Quality Control Testing: Immunoprecipitation-Western Blot (IP-WB)



Immunoprecipitation of MSX1 transfected lysate using mouse monoclonal anti-MSX1 and Protein A Magnetic Bead (U0007), and immunoblotted with rabbit polyclonal anti-MSX1.

Supplied Product: Antibody pair set content:
1. Antibody pair for IP: mouse monoclonal anti-MSX1 (300 ug)
2. Antibody pair for WB: rabbit polyclonal anti-MSX1 (50 ul)

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

Immunoprecipitation-Western Blot

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

Application Image

Immunoprecipitation-Western Blot

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