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

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AHI1 (Human) Matched Antibody Pair

Catalog # : H00054806-AP21 **規格 :** [1 Set]

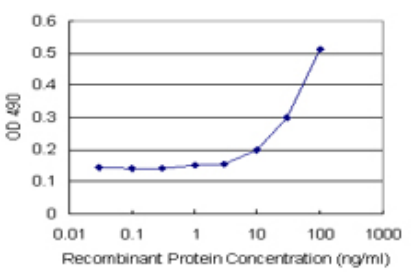
List All

Specification

Product Description: This antibody pair set comes with matched antibody pair to detect and quantify protein level of human AHI1.

Reactivity: Human

Quality Control Testing: Standard curve using recombinant protein (H00054806-P01) as an analyte.



Sandwich ELISA detection sensitivity ranging from 10 ng/ml to 100 ng/ml.

Supplied Product: Antibody pair set content:
1. Capture antibody: rabbit MaxPab® affinity purified polyclonal anti-AHI1 (100 ug)
2. Detection antibody: mouse purified polyclonal anti-AHI1 (20 ug)
*Reagents are sufficient for at least 1-2 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 (Recombinant protein)

Protocol Download

Gene Information

Entrez GeneID: 54806

Gene Name: AHI1

Gene Alias: AHI-1,DKFZp686J1653,FLJ14023,FLJ20069,JBTS3,ORF1,dJ71N10.1

Gene Description: Abelson helper integration site 1

Omim ID: 608629, 608894

Gene Ontology: Hyperlink

Application Image

ELISA Pair (Recombinant protein)

Gene Summary: This gene is apparently required for both cerebellar and cortical development in humans. This gene mutations cause specific forms of Joubert syndrome-related disorders. Joubert syndrome (JS) is a recessively inherited developmental brain disorder with several identified causative chromosomal loci. Alternatively spliced transcript variants encoding different isoforms have been identified. [provided by RefSeq

Other Designations: OTTHUMP00000017263,OTTHUMP00000017264,OTTHUMP00000017265,contatins SH3 and WD40 domains,jouberin

Related Disease

Abnormalities, Multiple Autistic Disorder Cerebellar Ataxia Diabetes Mellitus, Type 2 Genetic Predisposition to Disease Kidney Diseases, Cystic Mental Retardation Schizophrenia Schizophrenia Syndrome Tobacco Use Disorder