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Produktinformation



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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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NNMT (Human) Matched Antibody Pair

Catalog # : H00004837-AP11

規格 : [1 Set]

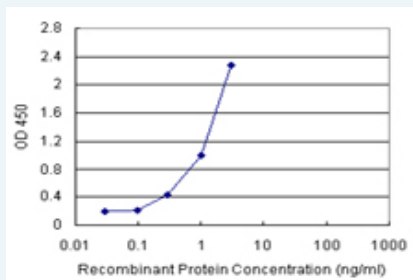
[List All](#)

Specification

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

Reactivity: Human

Quality Control Testing: Standard curve using recombinant protein (H00004837-P01) 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: rabbit MaxPab® affinity purified polyclonal anti-NNMT (100 ug)
 2. Detection antibody: mouse monoclonal anti-NNMT, IgG1 Kappa (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: [4837](#)

Gene Name: NNMT

Gene Alias: -

Gene Description: nicotinamide N-methyltransferase

Omim ID: [600008](#)

Gene Ontology: [Hyperlink](#)

Application Image

ELISA Pair (Recombinant protein)

Gene Summary: N-methylation is one method by which drug and other xenobiotic compounds are metabolized by the liver. This gene encodes the protein responsible for this enzymatic activity which uses S-adenosyl methionine as the methyl donor. [provided by RefSeq]

Other Designations: -

Gene Pathway

Metabolic pathways Nicotinate and nicotinamide metabolism

Related Disease

Aortic Aneurysm, Abdominal Atherosclerosis Brain Ischemia Cleft Lip Cleft Palate Ductus Arteriosus, Patent Genetic Predisposition to Disease Heart Defects, Congenital Hyperhomocysteinemia Infant, Premature, Diseases Migraine Disorders Migraine with Aura Prenatal Exposure Delayed Effects Spinal Dysraphism Stroke

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