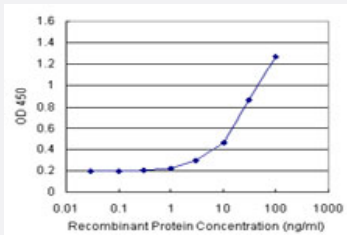


EDN3 (Human) Matched Antibody Pair

Catalog # H00001908-AP11

Size 1 Set

Applications



Sandwich ELISA detection sensitivity ranging from 1 ng/ml to 100 ng/ml.

Specification

Product Description	This antibody pair set comes with a matched antibody pair to detect and quantify the protein level of human EDN3.
Reactivity	Human
Quality Control Testing	Standard curve using recombinant protein (H00001908-P01) as an analyte. Sandwich ELISA detection sensitivity ranging from 1 ng/ml to 100 ng/ml.
Supplied Product	Antibody pair set content: 1. Capture antibody: rabbit MaxPab® affinity purified polyclonal anti-EDN3 (100 ug) 2. Detection antibody: mouse monoclonal anti-EDN3, 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.

Applications

- ELISA Pair (Recombinant protein)

[Protocol Download](#)

Gene Info — EDN3

Entrez GeneID [1908](#)

Gene Name EDN3

Gene Alias ET3, MGC15067, MGC61498

Gene Description endothelin 3

Omim ID [131242](#) [142623](#) [209880](#) [277580](#)

Gene Ontology [Hyperlink](#)

Gene Summary

The protein encoded by this gene is a member of the endothelin family. Endothelins are endothelium-derived vasoactive peptides involved in a variety of biological functions. The active form of this protein is a 21 amino acid peptide processed from the precursor protein. The active peptide is a ligand for endothelin receptor type B (EDNRB). The interaction of this endothelin with EDNRB is essential for development of neural crest-derived cell lineages, such as melanocytes and enteric neurons. Mutations in this gene and EDNRB have been associated with Hirschsprung disease (HSCR) and Waardenburg syndrome (WS), which are congenital disorders involving neural crest-derived cells. Four alternatively spliced transcript variants encoding three distinct isoforms have been observed. [provided by RefSeq]

Other Designations OTTHUMP00000031420|truncated endothelin 3

Pathway

- [Hypertrophic cardiomyopathy \(HCM\)](#)
- [Neuroactive ligand-receptor interaction](#)
- [Vascular smooth muscle contraction](#)

Disease

- [Cardiovascular Diseases](#)
- [Cystic Fibrosis](#)
- [Diabetes Mellitus](#)
- [Down Syndrome](#)
- [Edema](#)

- [Genetic Predisposition to Disease](#)
- [Hirschsprung Disease](#)
- [Sleep Apnea](#)