

Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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Lieferung & Zahlungsart

siehe unsere Liefer- und Versandbedingungen

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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EDN3 (Human) IP-WB Antibody Pair

Catalog #: H00001908-PW1 規格:[1 Set]

List All

Specification		Application Image
Product Description:	This IP-WB antibody pair set comes with one antibody for immunoprecipitation and another to detect the precipitated protein in western blot.	Immunoprecipitation-Western Blot
Reactivity:	Human	
Quality Control Testing:	Immunoprecipitation-Western Blot (IP-WB)	
	75 -	
	50 – 37 –	
	25	
	20-	
	15-	
	Immunoprecipitation of EDN3 transfected lysate using mouse monoclonal anti-EDN3 and Protein A Magnetic Bead (<u>U0007</u>), and immunoblotted with rabbit polyclonal anti-EDN3.	
Supplied Product:	Antibody pair set content: 1. Antibody pair for IP: mouse monoclonal anti-EDN3 (300 ug) 2. Antibody pair for WB: rabbit polyclonal anti-EDN3 (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		
mmunoprecipit	ation-Western Blot	
Gene Informatio	on	
Entrez GeneID:	1908	
Gene Name:	EDN3	
Gene Alias:	ET3,MGC15067,MGC61498	
Gene Description:	endothelin 3	
Omim ID:	<u>131242, 142623, 209880, 277580</u>	

Page 1 of 2 2016/5/20

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

Gene Pathway

Hypertrophic cardiomyopathy (HCM) Neuroactive ligand-receptor interaction Vascular smooth muscle contraction

Related Disease

Cardiovascular Diseases Cystic Fibrosis Diabetes Mellitus, Type 2 Down Syndrome Edema Genetic Predisposition to Disease Hirschsprung Disease Sleep Apnea, Obstructive

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Page 2 of 2 2016/5/20