

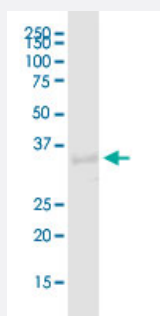
MaxPab®

PHOX2A MaxPab rabbit polyclonal antibody (D01)

Catalog # H00000401-D01

Size 100 uL

Applications



Immunoprecipitation

Immunoprecipitation of PHOX2A transfected lysate using anti-PHOX2A MaxPab rabbit polyclonal antibody and Protein A Magnetic Bead, and immunoblotted with ([H00000401-B01P](#)).

Specification

Product Description	Rabbit polyclonal antibody raised against a full-length human PHOX2A protein.
Immunogen	PHOX2A (NP_005160.2, 1 a.a. ~ 284 a.a) full-length human protein.
Sequence	MDYSLNSYDSCVAAMEASAYGDFGACSQPGGFQYSPLRPAFPAAGPPCPALGSSNCALGALR DHQPAPYSAPYKFFPEPSGLHEKRRKQRRIRTTFTSAQLKELERVFAETHYPDIYTREELALKIDLT EARVQVWFQNRRAKFRKQERAASAKGAAGAAGAKKGEARCSSEDDDSKESTCSPTPDSTASL PPPPAPGLASPRLSPLPVALGSGPGPGPGPQPLKGALWAGVAGGGGGPGAGAAELLKAW QPAESGPGPFSGVLSSFHRKPGPALKTNLF
Host	Rabbit
Reactivity	Human
Quality Control Testing	Antibody reactive against mammalian transfected lysate.
Storage Buffer	No additive
Storage Instruction	Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

Applications

- Immunoprecipitation

Immunoprecipitation of PHOX2A transfected lysate using anti-PHOX2A MaxPab rabbit polyclonal antibody and Protein A Magnetic Bead, and immunoblotted with ([H00000401-B01P](#)).

[Protocol Download](#)

Gene Info — PHOX2A

Entrez GeneID	401
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GeneBank Accession#	NM_005169.2
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Protein Accession#	NP_005160.2
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Gene Name	PHOX2A
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Gene Alias	ARIX, CFEOM2, FEOM2, MGC52227, NCAM2, PMX2A
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Gene Description	paired-like homeobox 2a
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Omim ID	602078 602753
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Gene Ontology	Hyperlink
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Gene Summary	The protein encoded by this gene contains a paired-like homeodomain most similar to that of the Drosophila aristaless gene product. The encoded protein plays a central role in development of the autonomic nervous system. It regulates the expression of tyrosine hydroxylase and dopamine beta-hydroxylase, two catecholaminergic biosynthetic enzymes essential for the differentiation and maintenance of the noradrenergic neurotransmitter phenotype. The encoded protein has also been shown to regulate transcription of the alpha3 nicotinic acetylcholine receptor gene. Mutations in this gene have been associated with autosomal recessive congenital fibrosis of the extraocular muscles. [provided by RefSeq]
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Other Designations	aristaless homeobox homolog arix homeodomain protein
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Disease

- [Sudden Infant Death](#)