

Datasheet

PHOX2A MaxPab rabbit polyclonal antibody (D01)

Catalog Number: H00000401-D01

Regulation Status: For research use only (RUO)

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:

MDYSYLSYDSCVAAMEASAYGDFGACSQPGGFQYS
PLRPAFPAAGPPCPALGSSNCALGALRDHQPAPYS
PYKFFPEPSGLHEKRKQRRIRTTFTSAQLKELERVFAE
THYPDIYTREELALKIDLTEARVQVWFQNRRAKFRKQE
RAASAKGAAGAAGAKKGEARCSSEDDDSKESTCSPT
PDSTASLPPPPAPGLASPRLSPLPVALGSGPGPGP
GPQPLKGALWAGVAGGGGGGPGAGAAELLKAWQPA
ESGPGPFSGVLSSFHRKPGPALKTNLF

Host: Rabbit

Reactivity: Human

Applications: IP

(See our web site product page for detailed applications information)

Protocols: See our web site at

<http://www.abnova.com/support/protocols.asp> or product page for detailed protocols

Storage Buffer: No additive

Storage Instruction: Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

Entrez GeneID: 401

Gene Symbol: PHOX2A

Gene Alias: ARIX, CFEOM2, FEOM2, MGC52227, NCAM2, PMX2A

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]