DNAxPAb | Hard-to-Find Antibody

SLC26A4 DNAxPab

Catalog # H00005172-W01P Size 200 ug

Specification

Product Description	Rabbit polyclonal antibody raised against a partial-length human SLC26A4 DNA using DNAx™ Immune technology.
Technology	DNAx™ Immune
Immunogen	Extracellular membrane domain (ECD) human DNA
Host	Rabbit
Reactivity	Human
Purification	Protein A
Quality Control Testing	Antibody reactive against mammalian transfected lysate.
Storage Buffer	In 1x PBS, pH 7.4
Storage Instruction	Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

Applications

- Western Blot (Transfected lysate)

  Protocol Download

- Immunofluorescence (Transfected cell)
- Flow Cytometry (Transfected cell)

Gene Info — SLC26A4

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Entrez GeneID	5172
GeneBank Accession#	BC153002
Protein Accession#	AAI53003.1
Gene Name	SLC26A4
Gene Alias	DFNB4, EVA, PDS
Gene Description	solute carrier family 26, member 4
Omim ID	274600 600791 605646
Gene Ontology	Hyperlink
Gene Summary	Mutations in this gene are associated with Pendred syndrome, the most common form of syndro mic deafness, an autosomal-recessive disease. It is highly homologous to the SLC26A3 gene; th ey have similar genomic structures and this gene is located 3' of the SLC26A3 gene. The encode d protein has homology to sulfate transporters. [provided by RefSeq
Other Designations	pendrin

Disease

- Auditory Threshold
- Autoimmune Diseases
- Bone Conduction
- Deafness
- Functional Laterality
- Genetic Predisposition to Disease
- Goiter
- Hearing Loss
- HIV Infections
- Pendred syndrome
- Presbycusis
- Speech Perception

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- Syndrome
- Thyroid Diseases
- Thyroid Neoplasms
- Vertigo

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