

DNAxPAb

Hard-to-Find
Antibody

WBSCR22 DNAxPab

Catalog # H00114049-W01P Size 200 ug

Specification

Product Description	Rabbit polyclonal antibody raised against a full-length human WBSCR22 DNA using DNAx™ Immune technology.
Technology	DNAx™ Immune
Immunogen	Full-length human DNA
Sequence	MASRGRRPEHGGPPELFYDETEARKYVRNSRMDIQTRMAGRALELLYPENKPCYLLDIGCGTGL SGSYLSDEGHYVWGLDISPAMLDEAVDREIEGDLGLGDMGQGIPFKPGTFDGCISISAVQWLCNA NKKSENPAKRLYCFASFSLVLRGSRVQLQYPENSEQLELITQATKAGFSGGMVVDYPNSAK AKKFYLCFLSGPSTFIPEGLSENQDEVEPRESVFTNERFPLRMSRRGMVRKSRAWVLEKKERHR RQGREVRPDTQYTGRKRKPRF
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 — WBSCR22

Entrez GeneID	114049
GeneBank Accession#	NM_017528.2
Protein Accession#	NP_059998.2
Gene Name	WBSCR22
Gene Alias	HASJ4442, HUSSY-3, MGC19709, MGC2022, MGC5140, PP3381, WBMT
Gene Description	Williams Beuren syndrome chromosome region 22
Gene Ontology	Hyperlink
Gene Summary	This gene encodes a protein containing a nuclear localization signal and an S-adenosyl-L-methionine binding motif typical of methyltransferases, suggesting that the encoded protein may act on DNA methylation. This gene is deleted in Williams syndrome, a multisystem developmental disorder caused by the deletion of contiguous genes at 7q11.23. [provided by RefSeq]
Other Designations	Williams Beuren syndrome chromosome region 22 protein Williams-Beuren candidate region putative methyltransferase

Pathway

- [Aminophosphonate metabolism](#)
- [Androgen and estrogen metabolism](#)
- [Histidine metabolism](#)
- [Selenoamino acid metabolism](#)
- [Tyrosine metabolism](#)