DNAxPAb | Hard-to-Find Antibody

WBSCR16 DNAxPab

Catalog # H00081554-W01P Size 200 ug

Specification

Product Description	Rabbit polyclonal antibody raised against a full-length human WBSCR16 DNA using DNAx™ Immune technology.
Technology	DNAx™ Immune
Immunogen	Full-length human DNA
Sequence	MALVALVAGARLGRRLSGPGLGRGHWTAAGRSRSRREAAEAEAEVPVVQYVGERAARADRVF VWGFSFSGALGVPSFVVPSSGPGPRAGARPRRRIQPVPYRLELDQKISSAACGYGFTLLSSKTA DVTKVWGMGLNKDSQLGFHRSRKDKTRGYEYVLEPSPVSLPLDRPQETRVLQVSCGRAHSLVL TDREGVFSMGNNSYGQCGRKVVENEIYSESHRVHRMQDFDGQVVQVACGQDHSLFLTDKGEVY SCGWGADGQTGLGHYNITSSPTKLGGDLAGVNVIQVATYGDCCLAVSADGGLFGWGNSEYLQLA SVTDSTQVNVPRCLHFSGVGKVRQAACGGTGCAVLNGEGHVFVWGYGILGKGPNLVESAVPEM IPPTLFGLTEFNPEIQVSRIRCGLSHFAALTNKGELFVWGKNIRGCLGIGRLEDQYFPWRVTMPGEP VDVACGVDHMVTLAKSFI
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 — WBSCR16

Entrez GeneID	81554
GeneBank Accession#	BC007823.2
Protein Accession#	AAH07823.1
Gene Name	WBSCR16
Gene Alias	DKFZp434D0421, MGC189739, MGC44931
Gene Description	Williams-Beuren syndrome chromosome region 16
Gene Ontology	Hyperlink
Gene Summary	This gene encodes an RCC1-like G-exchanging factor. It is deleted in Williams syndrome, a multi system developmental disorder caused by the deletion of contiguous genes at 7q11.23. [provided by RefSeq
Other Designations	RCC1-like G exchanging factor-like