

DNAxPAb

Hard-to-Find
Antibody

BBS2 DNAxPab

Catalog # H00000583-W01P

Size 200 ug

Specification

Product Description	Rabbit polyclonal antibody raised against a full-length human BBS2 DNA using DNAx™ Immune technology.
Technology	DNAx™ Immune
Immunogen	Full-length human DNA
Sequence	MLLPVFTLKL RHKISPRMVAIGRYDGHPCLA AATQTGKVFIHNP HTRNQHV SASRVFQSPLESDV SLLNINQAVSCLTAGVLNPELG YDALLVGTQTNLLAYDVYNN SDFYREVADGANVVVLGTLGDIS SPLAIGGN CALQGFNHEGS DLFWTVTGDNVNSLALCDFDGDGKKELLVGSEDFDIRVFKED EIV AEMTETEIVTSLCPMYGSRFGYALSNGTVGVYDKTSRYWRIKSKNHAMSIHAFDLNSDGVNELITG WSNGKVDARS DRTGEVIFKDNFSSA IAGVVEGDYRMDGHIQLICCSVDGEIRGYLPGTAEMRGNL MDTSAEQDLIRELSQKKQNL LLELRNYEENAKAELASPLNEADGHRGIIPANTRLHTT LSVSLGNET QTAHTELRISTSN DTTIRAVLIFAEGIFTGESHVHPSIHNLS SSIPIVPPKDV PVDLHLKAFVGYRSS TQFHVFESTRQLPRFS MYALTSLDPASEPISYVNF TIAERAQRV VVWL GQNFLLPEDTHIQNAPFQ VCFTSLRNGGHLHIKILSGEITINTDDIDL AGDIQSMASFFAIEDLQVEADFPVYFEELRKVLVKVD EYHSVHQKLSAD MADHSNLIRSL LVGAEDARLMRDMKTMKSR YMELYDLNRDLLNGYKIRCNNHT ELLGNLKAVNQAIQRAGRLRVGKPKNQVITACRDAIRSN NINTL FKIMRVGTASS
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 — BBS2

Entrez GeneID [583](#)

GeneBank Accession# [BC014140.2](#)

Protein Accession# [AAH14140.1](#)

Gene Name BBS2

Gene Alias BBS, MGC20703

Gene Description Bardet-Biedl syndrome 2

Omim ID [209900 606151](#)

Gene Ontology [Hyperlink](#)

Gene Summary This gene is a member of the Bardet-Biedl syndrome (BBS) gene family. Bardet-Biedl syndrome is an autosomal recessive disorder characterized by severe pigmentary retinopathy, obesity, polydactyly, renal malformation and mental retardation. The proteins encoded by BBS gene family members are structurally diverse and the similar phenotypes exhibited by mutations in BBS gene family members is likely due to their shared roles in cilia formation and function. Many BBS proteins localize to the basal bodies, ciliary axonemes, and pericentriolar regions of cells. BBS proteins may also be involved in intracellular trafficking via microtubule-related transport. The protein encoded by this gene forms a multiprotein BBSome complex with six other BBS proteins

Other Designations Bardet-Biedl syndrome 2 protein

Disease

- [Bardet-Biedl Syndrome](#)
- [Cardiovascular Diseases](#)
- [Diabetes Mellitus](#)
- [Edema](#)

- [Obesity](#)
- [Retinal Diseases](#)