

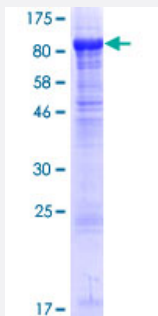
Full-Length

BBS2 (Human) Recombinant Protein (P01)

Catalog # H00000583-P01

Size 25 ug, 10 ug

Applications



Specification

Product Description

Human BBS2 full-length ORF (AAH14140.1, 1 a.a. - 721 a.a.) recombinant protein with GST-tag at N-terminal.

Sequence

MLLPVFTLKL RHKISPRMVAIGRYDGHPC LAAATQTGKVFIHNP HTRNQHV SASRVFQSPLESDV
SLLNINQAVSCLTAGVLNPELGYDALLVGTQTNLLAYDVYNNSDLFYREVADGANVVVLGTLGDIS
SPLAII GGNCALQGFNHEGSDLFWTVTGDNVNSLALCDFDGDGKKELLVGSEDFDIRVFKEDIV
AEMTETEIVTSLCPMYGSRFGYALSNGTVGVYDKTSRYWRIKSKNHAMSIHAFDLNSDGVNELITG
WSNGKVDARS DRTGEVIFKDNFSSA IAGVVEGDYRMDGHIQLICCSVDGEIRGYLPGTAEMRGNL
MDTSAEQDLIRELSQKKQNLLLELRNYEENAKAELASPLNEADGHRGIIPANTRLHTLSVSLGNET
QTAHTEL RISTSDNTIIRAVLIFAEGIFTGESHVVHPSIHNLSSSICIPVPPKDV PVDLHLKAFVGYRSS
TQFHFVFESTRQLPRFSMYALTSLDPASEPISYNFTIAERAQRVVVWL GQNFLLPEDTHIQNAPFQ
VCFTSLRNGGHLHIKILSGEITINTDDIDL AGDIQSMASFFAIEDLQVEADFPVYFEELRKVLVKVD
EYHSVHQKLSADMADHSNLIRSLLVGAEDARLMRDMKTMKSRYMELYDLNRDLLNGYKIRCNNHT
ELLGNLKA VNQAIQRAGRLRVGKPKNQVITACRDAIRSNNTLFLKIMRVGTASS

Host

Wheat Germ (in vitro)

Theoretical MW (kDa)

106.3

Interspecies Antigen Sequence

Mouse (90); Rat (90)

Preparation Method

[in vitro wheat germ expression system](#)

Purification	Glutathione Sepharose 4 Fast Flow
Quality Control Testing	12.5% SDS-PAGE Stained with Coomassie Blue.
Storage Buffer	50 mM Tris-HCl, 10 mM reduced Glutathione, pH=8.0 in the elution buffer.
Storage Instruction	Store at -80°C. Aliquot to avoid repeated freezing and thawing.
Note	Best use within three months from the date of receipt of this protein.

Applications

- Enzyme-linked Immunoabsorbent Assay
- Western Blot (Recombinant protein)
- Antibody Production
- Protein Array

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](#)