

Rabbit Anti-Human BBS10 Polyclonal antibody (DPABH-15616)

This product is for research use only and is not intended for diagnostic use.

PRODUCT INFORMATION

Immunogen	BBS10 fusion protein, sequence: MLPVSCKLPNMGTSQSYLSSSMPAGCVLPVGGNFEILLHYYLLNYAKKCHQSEETMVSMI IANALLGIPKVLYKSKTGKYSFPHTYIRAVHALQTNQPLVSSQTGLESVMGKYQLLTSVL QCLTKILTIDMVITVKRHPQKVHNQDSEDEL (C-term-151aa encoded by BC013795)
Isotype	IgG
Source/Host	Rabbit
Species Reactivity	Human, Zebrafish, Mouse
Purification	Antigen affinity purification
Conjugate	Unconjugated
Applications	WB, IF, ELISA
Positive Control	HepG2 cells, A549 cells, MCF-7 cells
Format	Liquid
Size	50 μl, 100 μl
Buffer	PBS with 0.02% sodium azide and 50% glycerol pH 7.3.
Preservative	0.02% Sodium Azide
Storage	Store at -20°C. Aliquoting is unnecessary for -20°C storage.

BACKGROUND

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Introduction	This gene is a member of the Bardet-Biedl syndrome (BBS) gene family. Bardet-Biedl
	syndrome is an autosomal recessive disorder characterized by progressive retinal
	degeneration, 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 is likely not a ciliary protein but
	rather has distant sequence homology to type II chaperonins. As a molecular chaperone, this
	protein may affect the folding or stability of other ciliary or basal body proteins. Inhibition of this
	proteins expression impairs ciliogenesis in preadipocytes. Mutations in this gene cause Bardet-
	Biedl syndrome type 10.

Keywords

BBS10; Bardet-Biedl syndrome 10; C12orf58; Bardet-Biedl syndrome 10 protein;

GENE INFORMATION

Entrez Gene ID	79738
UniProt ID	Q8TAM1

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