



# Human TTC8 blocking peptide (CDBP3098)

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

## PRODUCT INFORMATION

<b>Product Overview</b>	Blocking/Immunizing peptide for anti-TTC8 antibody
<b>Antigen Description</b>	This gene encodes a protein that has been directly linked to Bardet-Biedl syndrome. The primary features of this syndrome include retinal dystrophy, obesity, polydactyly, renal abnormalities and learning disabilities. Experimentation in non-human eukaryotes suggests that this gene is expressed in ciliated cells and that it is involved in the formation of cilia. A mutation in this gene has also been implicated in nonsyndromic retinitis pigmentosa. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Jan 2014]
<b>Species</b>	Human
<b>Conjugate</b>	Unconjugated
<b>Applications</b>	Apuri, BL, ELISA
<b>Format</b>	Lyophilized powder
<b>Size</b>	100 µg
<b>Preservative</b>	None
<b>Storage</b>	Shipped at ambient temperature, store at -20°C.

## GENE INFORMATION

<b>Gene Name</b>	<a href="#">TTC8 tetratricopeptide repeat domain 8 [ Homo sapiens ]</a>
<b>Official Symbol</b>	TTC8
<b>Synonyms</b>	TTC8; tetratricopeptide repeat domain 8; tetratricopeptide repeat protein 8; BBS8; TPR repeat protein 8; Bardet-Biedl syndrome type 8; bardet-Biedl syndrome 8 protein; RP51;

<b>Entrez Gene ID</b>	<a href="#">123016</a>
<b>mRNA Refseq</b>	<a href="#">NM_144596</a>
<b>Protein Refseq</b>	<a href="#">NP_653197</a>
<b>UniProt ID</b>	Q8TAM2
<b>Chromosome Location</b>	14q31.3
<b>Function</b>	protein binding;