



# Anti-BSND (aa 1-320) polyclonal antibody (CPBT-27041MH)

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

## PRODUCT INFORMATION

<b>Product Overview</b>	Mouse Polyclonal antibody to Human BSND.
<b>Antigen Description</b>	This gene encodes an essential beta subunit for CLC chloride channels. These heteromeric channels localize to basolateral membranes of renal tubules and of potassium-secreting epithelia of the inner ear. Mutations in this gene have been associated with Bartter syndrome with sensorineural deafness.
<b>Immunogen</b>	Full length protein, corresponding to amino acids 1-320 of Human BSND
<b>Isotype</b>	IgG
<b>Source/Host</b>	Mouse
<b>Species Reactivity</b>	Human
<b>Purification</b>	Protein A purified
<b>Conjugate</b>	Unconjugated
<b>Applications</b>	WB, IHC-P
<b>Cellular Localization</b>	Cell membrane; Multipass membrane protein. Cytoplasm.
<b>Format</b>	Liquid
<b>Size</b>	50 µg
<b>Buffer</b>	Preservative: None Constituents: 1X PBS, pH 7.2
<b>Preservative</b>	None

<b>Storage</b>	Shipped at 4°C. Upon delivery aliquot and store at -20°C or -80°C. Avoid repeated freeze / thaw cycles.
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## GENE INFORMATION

<b>Gene Name</b>	<a href="#">BSND Bartter syndrome, infantile, with sensorineural deafness (Barttin) [ Homo sapiens ]</a>
<b>Official Symbol</b>	BSND
<b>Synonyms</b>	BSND; Bartter syndrome, infantile, with sensorineural deafness (Barttin); deafness, autosomal recessive 73 , DFNB73; barttin; BART; BART; Bartter syndrome infantile with sensorineural deafness; Bartter syndrome, infantile, with sensorineural deafness (Barttin); Barttin; deafness, autosomal recessive 73; DFNB 73; DFNB73; deafness, autosomal recessive 73; DFNB73;
<b>Entrez Gene ID</b>	<a href="#">7809</a>
<b>Protein Refseq</b>	<a href="#">NP_476517</a>
<b>UniProt ID</b>	<a href="#">Q5VU50</a>
<b>Chromosome Location</b>	1p32.3
<b>Function</b>	contributes_to chloride channel activity; chloride channel regulator activity;