

DNAxPAb

Hard-to-Find  
Antibody

# CENPJ DNAxPab

Catalog # H00055835-W01P

Size 200 ug

## Specification

Product Description	Rabbit polyclonal antibody raised against a full-length human CENPJ DNA using DNAx™ Immune technology.
Technology	<a href="#">DNAx™ Immune</a>
Immunogen	Full-length human DNA
Sequence	MRKLQKERKVFKEYTTAARTFPDKKEREEIQLTKQIADLREDLKRKETKWSSTHSRLRSQIQMLV RENTDLREEIKVMERFRLDAWKRAEAISSLEVEKKDKLANTSVRFQNSQISSGTQVEKYKKNYL PMQGNPPRRSKSAPPRDLGNLDKGQAASPREPLEPLNFPDPEYKEEEEDQDIQGEISHPDGKV EKVYKNGCRVILFPNGTRKEVSADGKTITVTFFNGDVKQVMPDQQRVIYAAAQTTHTTTYPEGLEVL HFSSGQIEKHYPDGRKEITFPDQTVKNLFPDGGQEESIFPDGTVRVQRDGNKLIEFNNGQRELHTA QFKRREYPDGTVKTVYANGHQETKYRSGRIRVKDKEGNVLMDEL
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 — CENPJ

Entrez GeneID	<a href="#">55835</a>
GeneBank Accession#	<a href="#">BC024209.2</a>
Protein Accession#	<a href="#">AAH24209.3</a>
Gene Name	CENPJ
Gene Alias	BM032, CPAP, LAP, LIP1, MCPH6, MGC131581, MGC131582, MGC142222, MGC142224
Gene Description	centromere protein J
Omim ID	<a href="#">608393</a> <a href="#">609279</a>
Gene Ontology	<a href="#">Hyperlink</a>
Gene Summary	This gene encodes a protein that belongs to the centromere protein family. During cell division, this protein plays a structural role in the maintenance of centrosome integrity and normal spindle morphology, and it is involved in microtubule disassembly at the centrosome. This protein can function as a transcriptional coactivator in the Stat5 signaling pathway, and also as a coactivator of NF-kappaB-mediated transcription, likely via its interaction with the coactivator p300/CREB-binding protein. Mutations in this gene are associated with primary autosomal recessive microcephaly, a disorder characterized by severely reduced brain size and mental retardation. [provided by RefSeq]
Other Designations	LAG-3-associated protein LYST-interacting protein LIP1 LYST-interacting protein LIP7 OTTHUM P00000018137 centrosomal P4.1-associated protein

## Disease

- [Genetic Predisposition to Disease](#)
- [Microcephaly](#)