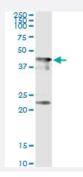
# CENPJ (Human) IP-WB Antibody Pair

Catalog # H00055835-PW2 Size 1 Set

## Applications



Immunoprecipitation of CENPJ transfected lysate using mouse monoclonal anti-CENPJ and Protein A Magnetic Bead (<u>U0007</u>), and immunoblotted with rabbit polyclonal anti-CENPJ.

Specification	
Product Description	This IP-WB antibody pair set comes with one antibody for immunoprecipitation and another to detect the precipitated protein in western blot.
Reactivity	Human
Interspecies Antigen Sequence	Mouse (83); Rat (81)
Quality Control Testing	Immunoprecipitation-Western Blot (IP-WB) Immunoprecipitation of CENPJ transfected lysate using mouse monoclonal anti-CENPJ and Protein A Magnetic Bead ( <u>U0007</u> ), and immunoblotted with rabbit polyclonal anti-CENPJ.
Supplied Product	Antibody pair set content: 1. Antibody pair for IP: mouse monoclonal anti-CENPJ (300 ug) 2. Antibody pair for WB: rabbit polyclonal anti-CENPJ (50 ul)
Storage Instruction	Store reagents of the antibody pair set at -20°C or lower. Please aliquot to avoid repeated freeze tha w cycle. Reagents should be returned to -20°C storage immediately after use.

#### Applications

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• Immunoprecipitation-Western Blot

Protocol Download

### Gene Info — CENPJ

Entrez GenelD	<u>55835</u>
Gene Name	CENPJ
Gene Alias	BM032, CPAP, LAP, LIP1, MCPH6, MGC131581, MGC131582, MGC142222, MGC142224
Gene Description	centromere protein J
Omim ID	<u>608393</u> <u>609279</u>
Gene Ontology	Hyperlink
Gene Summary	This gene encodes a protein that belongs to the centromere protein family. During cell division, thi s protein plays a structural role in the maintenance of centrosome integrity and normal spindle mor phology, 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-ka ppaB-mediated transcription, likely via its interaction with the coactivator p300/CREB-binding pro tein. Mutations in this gene are associated with primary autosomal recessive microcephaly, a dis order 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 P0000018137 centrosomal P4.1-associated protein

#### Disease

- Genetic Predisposition to Disease
- Microcephaly