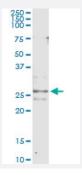


CFC1 (Human) IP-WB Antibody Pair

Catalog # H00055997-PW2 Size 1 Set

Applications



Immunoprecipitation of CFC1 transfected lysate using mouse monoclonal anti-CFC1 and Protein A Magnetic Bead (<u>U0007</u>), and immunoblotted with rabbit polyclonal anti-CFC1.

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 (54); Rat (55)
Quality Control Testing	Immunoprecipitation-Western Blot (IP-WB) Immunoprecipitation of CFC1 transfected lysate using mouse monoclonal anti-CFC1 and Protein A Magnetic Bead (U0007), and immunoblotted with rabbit polyclonal anti-CFC1.
Supplied Product	Antibody pair set content: 1. Antibody pair for IP: mouse monoclonal anti-CFC1 (300 ug) 2. Antibody pair for WB: rabbit polyclonal anti-CFC1 (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



Immunoprecipitation-Western Blot

Protocol Download

Gene Info — CFC1	
Entrez GeneID	<u>55997</u>
Gene Name	CFC1
Gene Alias	CRYPTIC, FLJ77897, HTX2, MGC133213
Gene Description	cripto, FRL-1, cryptic family 1
Omim ID	<u>217095</u> <u>605194</u> <u>605376</u>
Gene Ontology	<u>Hyperlink</u>
Gene Summary	This gene encodes a member of the epidermal growth factor (EGF)- Cripto, Frl-1, and Cryptic (CFC) family. EGF-CFC family member proteins share a variant EGF-like motif, a conserved cysteine-rich domain, and a C-terminal hydrophobic region. These proteins play key roles in intercellular signaling pathways during vertebrate embryogenesis. Mutations in this gene can cause autosoma I visceral heterotaxy. This protein is involved in left-right asymmetric morphogenesis during organ development. [provided by RefSeq
Other Designations	cryptic

Disease

- Cleft Lip
- Cleft Palate
- Heart Defects
- Heart Septal Defects