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Produktinformation

- Forschungsprodukte & Biochemikalien
- Zellkultur & Verbrauchsmaterial
- Diagnostik & molekulare Diagnostik
- Laborgeräte & Service

Weitere Information auf den folgenden Seiten!
See the following pages for more information!

Lieferung & Zahlungsart

siehe unsere Liefer- und Versandbedingungen

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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GTF2IRD1 (Human) IP-WB Antibody Pair

Catalog # : H00009569-PW2 **規格 :** [1 Set]

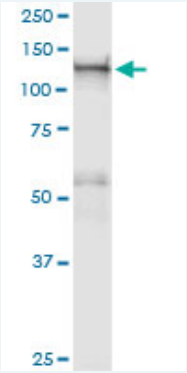
List All

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

Quality Control Testing: Immunoprecipitation-Western Blot (IP-WB)



Immunoprecipitation of GTF2IRD1 transfected lysate using rabbit polyclonal anti-GTF2IRD1 and Protein A Magnetic Bead (U0007), and immunoblotted with mouse purified polyclonal anti-GTF2IRD1.

Supplied Product: Antibody pair set content:
1. Antibody pair for IP: rabbit polyclonal anti-GTF2IRD1 (300 ul)
2. Antibody pair for WB: mouse purified polyclonal anti-GTF2IRD1 (50 ug)

Storage Instruction: Store reagents of the antibody pair set at -20°C or lower. Please aliquot to avoid repeated freeze thaw cycle. Reagents should be returned to -20°C storage immediately after use.

Applications

Immunoprecipitation-Western Blot

Protocol Download

Gene Information

Entrez GeneID: 9569

Gene Name: GTF2IRD1

Gene Alias: BEN,CREAM1,GTF3,MUSTRD1,RBAP2,WBS,WBSCR11,WBSCR12,hMusTRD1alpha1

Gene Description: GTF2I repeat domain containing 1

Omim ID: 194050, 604318

Gene Ontology: Hyperlink

Application Image

Immunoprecipitation-Western Blot

Gene Summary:	The protein encoded by this gene contains five GTF2I-like repeats and each repeat possesses a potential helix-loop-helix (HLH) motif. It may have the ability to interact with other HLH-proteins and function as a transcription factor or as a positive transcriptional regulator under the control of Retinoblastoma protein. This gene is deleted in Williams-Beuren syndrome, a multisystem developmental disorder caused by deletion of multiple genes at 7q11.23. Alternative splicing of this gene generates at least 2 transcript variants. [provided by RefSeq
Other Designations:	GTF2I repeat domain-containing 1,Williams-Beuren syndrome chromosome region 11,binding factor for early enhancer,general transcription factor 3,muscle TFII-I repeat domain-containing protein 1 alpha 1

Gene Pathway

Basal transcription factors

Related Disease

Celiac Disease Genetic Predisposition to Disease