



# GTD2A rabbit pAb

<b>Catalog No</b>	BYab-08087
<b>Isotype</b>	IgG
<b>Reactivity</b>	Human;Rat;Mouse;
<b>Applications</b>	WB
<b>Gene Name</b>	GTF2IRD2 GTF2IRD2A
<b>Protein Name</b>	GTD2A
<b>Immunogen</b>	Synthesized peptide derived from human GTD2A AA range: 529-579
<b>Specificity</b>	This antibody detects endogenous levels of GTD2A at Human
<b>Formulation</b>	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.202% sodium azide.
<b>Source</b>	Polyclonal, Rabbit,IgG
<b>Purification</b>	The antibody was affinity-purified from rabbit serum by affinity-chromatography using specific immunogen.
<b>Dilution</b>	WB 1:500-2000
<b>Concentration</b>	1 mg/ml
<b>Purity</b>	≥90%
<b>Storage Stability</b>	-20°C/1 year
<b>Synonyms</b>	General transcription factor II-I repeat domain-containing protein 2A (GTF2I repeat domain-containing protein 2A) (Transcription factor GTF2IRD2-alpha)
<b>Observed Band</b>	105kD
<b>Cell Pathway</b>	Nucleus.
<b>Tissue Specificity</b>	Ubiquitous.
<b>Function</b>	miscellaneous:GTF2IRD2 is a gene located in the Williams-Beuren syndrome (WBS) critical region. WBS is the result of a chromosomal microdeletion 7q11.23 thought to arise as a consequence of unequal crossing over between highly homologous low-copy repeat sequences flanking the deleted region. Since GTF2IRD2 maps within this duplicated region, more than one copy of the gene was identified. WBS is a rare developmental disorder characterized by distinctive dysmorphic face, mild growth retardation, supraaortic stenosis and infantile hypercalcemia.,similarity:Belongs to the TFII-I family.,similarity:Contains 2 GTF2I-like repeats.,tissue specificity:Ubiquitous.,

Nanjing BYabscience technology Co.,Ltd

**Background**

This gene is one of several closely related genes on chromosome 7 encoding proteins containing helix-loop-helix motifs. These proteins may function as regulators of transcription. The encoded protein is unique in that its C-terminus is derived from CHARLIE8 transposable element sequence. This gene is located in a region of chromosome 7 that is deleted in Williams-Beuren syndrome, and loss of this locus may contribute to the cognitive phenotypes observed in this disease. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Jul 2013],

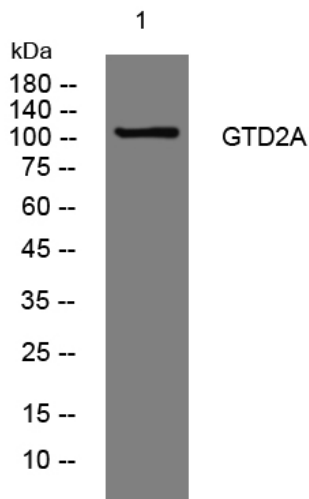
**matters needing attention**

Avoid repeated freezing and thawing!

**Usage suggestions**

This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.

## Products Images



Western blot analysis of lysates from HpeG2 cells, primary antibody was diluted at 1:1000, 4° over night