

GTF2IRD1 Rabbit pAb

货号: **AYP16661**

产品信息

反应	Human,Mouse,Rat
宿主	Rabbit
克隆性	Polyclonal
预测反应	
应用	WB
推荐浓度	WB: 1:500 - 1:2000
理论分子量	104kDa/106kDa/107kDa
实测分子量	106kDa
形式	Liquid
保存条件	Store at -20°C. Avoid freeze / thaw cycles. Buffer: PBS with 0.02% sodium azide,50% glycerol,pH7.3.
偶联物	Unconjugated
阳性对照	293T,HeLa,BT-474,SW480,Mouse ovary,Mouse thymus,Rat kidney,Rat liver
细胞定位	Nucleus
纯化	Affinity purification

抗原信息

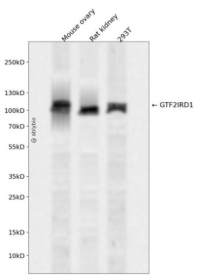
抗原信息	Recombinant fusion protein containing a sequence corresponding to amino acids 660-959 of human GTF2IRD1 (NP_057412.1).
序列	SLGFSPPALPPERDSGDPLVDES LKRQGFQENYDARLSRIDIAN TLREQVDLFNK KYGEALGIKYPVQVPYKRIKSNPGSV IIEGLPPGIPFRK PCTFGSQNLERILAVADKIKFTVTRPFQGLIPKPEDDANRLGEKVILREQVKELFNEKYGEALGLNRPVLV PYKLIRDSPDAVEVTGLPDDIPFRNPNTYDIHRLEKILKAREHVRMVIINQLQPFAEICNDAKVPAKDSSIPKRKRKRKRVSEGNS VSSSSSSSSSSSNPDSVASANQISLVQWPMYMYVDYAGLNVQLPGPLNY

靶点信息

研究背景	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 plays a role in craniofacial and cognitive development and mutations have been associated with Williams-Beuren syndrome, a multisystem developmental disorder caused by deletion of multiple genes at 7q11.23. Alternative splicing results in multiple transcript variants.
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基因ID	9569
基因名	GTF2IRD1
Swiss	Q9UHL9
别名	GTF2IRD1;BEN;CREAM1;GTF3;MUSTRD1;RBAP2;WBS;WBSER11;WBSER12

产品验证



Western blot analysis of GTF2IRD1 expressed in Mouse ovary, Rat kidney, 293T using GTF2IRD1 Rabbit pAb at 1:1000. Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) at 1:5000. Lysates/proteins: 30ug per lane. Blocking buffer: 5% non-fat dry milk in TBST. Detection: ECL Enhanced Kit. Exposure time: 12 0s.

实验步骤

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