

# RBM10 Rabbit pAb

货号: **AYP17221**

## 产品信息

|       |  |
|-------|--|
| 反应    | Human  |
| 宿主    | Rabbit   |
| 克隆性   | Polyclonal   |
| 预测反应  |  |
| 应用    | WB   |
| 推荐浓度  | <b>WB:</b> 1:500 - 1:2000  |
| 理论分子量 | 94kDa/103kDa   |
| 实测分子量 | 105kDa   |
| 形式    | Liquid   |
| 保存条件  | Store at -20°C. Avoid freeze / thaw cycles.<br>Buffer: PBS with 0.02% sodium azide,50% glycerol,pH7.3. |
| 偶联物   | Unconjugated   |
| 阳性对照  | Jurkat,MCF7,LO2,293T   |
| 细胞定位  | Nucleus  |
| 纯化    | Affinity purification  |

## 抗原信息

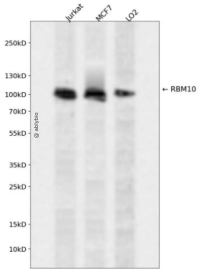
|      |   |
|------|---|
| 抗原信息 | Recombinant fusion protein containing a sequence corresponding to amino acids 1-180 of human RBM10 (NP_005667.2).   |
| 序列   | MEYERRGGRGDRTGRYGATDRSQDDGGENRSRDHDYRDMDYRSYPREYGSQEGKHDYDDSSSEEQSAEDSYEASPGS<br>ETQRRRRRRHRHSPGPPGFPRDGDYRDQDYRTEQEEEEEEEEEEEEKASNIVMLRMLPQAATEDDIRGQLQSHGVQ<br>AREVRLMRNKSSGQSRGFVFEFSH |

## 靶点信息

|      |   |
|------|---|
| 研究背景 | This gene encodes a nuclear protein that belongs to a family proteins that contain an RNA-binding motif. The encoded protein associates with hnRNP proteins and may be involved in regulating alternative splicing. Defects in this gene are the cause of the X-linked recessive disorder, TARP syndrome. Alternate splicing results in multiple transcript variants. |
| 基因ID | 8241  |

|       |   |
|-------|---|
| 基因名   | RBM10   |
| Swiss | P98175  |
| 别名    | RBM10;DXS8237E;GPATC9;GPATCH9;S1-1;TARPS;ZRANB5 |

## 产品验证



Western blot analysis of RBM10 expressed in Jurkat, MCF7, LO2 using RBM10 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: 120s.

## 实验步骤

访问官网浏览详情: [www.ablybio.cn](http://www.ablybio.cn)