

RUNX2 Rabbit mAb

货号: B28989

产品信息

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| 反应 | Human,Mouse,Rat |
| 宿主 | Rabbit |
| 克隆性 | Monoclonal |
| 预测反应 | |
| 应用 | IHC IF/ICC |
| 推荐浓度 | IHC: 1:50 - 1:200 IF/ICC: 1:50 - 1:200 |
| 理论分子量 | 54kDa/55kDa/56kDa |
| 实测分子量 | 60KDa |
| 形式 | Liquid |
| 保存条件 | Store at -20°C. Avoid freeze / thaw cycles. Buffer: PBS with 0.75% BSA,50% glycerol,pH7.3. |
| 偶联物 | Unconjugated |
| 阳性对照 | MCF7,HeLa,Mouse testis |
| 细胞定位 | Nucleus |
| 纯化 | Affinity purification |

抗原信息

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| 抗原信息 | Recombinant fusion protein corresponding to Human RUNX2. |
| 序列 | TITVFTNPPQVATYHRAIKVTVDGPREPRRHRQKLDDSKPSLFSDRLSDLGRIPHPSMRVGVPPQNPRPSLNSAPSPFNPQ GQSQITDPRQAQSSPPWSYD |

靶点信息

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| 研究背景 | This gene is a member of the RUNX family of transcription factors and encodes a nuclear protein with an Runt DNA-binding domain. This protein is essential for osteoblastic differentiation and skeletal morphogenesis and acts as a scaffold for nucleic acids and regulatory factors involved in skeletal gene expression. The protein can bind DNA both as a monomer or, with more affinity, as a subunit of a heterodimeric complex. Two regions of potential trinucleotide repeat expansions are present in the N-terminal region of the encoded protein, and these and other mutations in this gene have been associated with the bone development disorder cleidocranial dysplasia (CCD). Transcript variants that encode different protein isoforms result from the use of alternate promoters as well as alternate splicing. |
| 基因ID | 860 |
| 基因名 | RUNX2 |
| Swiss | Q13950 |
| 别名 | RUNX2;AML3;CBF-alpha-1;CBFA1;CCD;CCD1;CLCD;OSF-2;OSF2;PEA2aA;PEBP2aA |

产品验证

实验步骤

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