

Product datasheet

Anti-Raf1 antibody [410] ab656

1 References

概述

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|-------|---|
| 产品名称 | Anti-Raf1抗体[410] |
| 描述 | 小鼠单克隆抗体[410] to Raf1 |
| 宿主 | Mouse |
| 特异性 | This antibody is specifically reactive to Raf-1 N-terminal (1-240 a.a.) |
| 经测试应用 | 适用于: WB |
| 种属反应性 | 与反应: Human |
| 免疫原 | Purified recombinant human Raf-1 (mw:75 kDa). |

性能

| | |
|------|---|
| 形式 | Liquid |
| 存放说明 | Shipped at 4°C. Upon delivery aliquot and store at -20°C or -80°C. Avoid repeated freeze / thaw cycles. |
| 纯化说明 | Affinity purified from ascitic fluid. |
| 克隆 | 单克隆 |
| 克隆编号 | 410 |
| 骨髓瘤 | unknown |
| 同种型 | IgG1 |
| 轻链类型 | kappa |

应用

Our [Abpromise guarantee](#) covers the use of **ab656** in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

| 应用 | Ab评论 | 说明 |
|----|------|--|
| WB | | Use a concentration of 1 µg/ml. Will allow visualization of 0.2 µg/lane of human Raf-1 |

图标

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|--------------|--|
| 功能 | Involved in the transduction of mitogenic signals from the cell membrane to the nucleus. Part of the Ras-dependent signaling pathway from receptors to the nucleus. Protects cells from apoptosis mediated by STK3. |
| 组织特异性 | In skeletal muscle, isoform 1 is more abundant than isoform 2. |
| 疾病相关 | Defects in RAF1 are the cause of Noonan syndrome type 5 (NS5) [MIM:611553]. Noonan syndrome (NS) is a disorder characterized by dysmorphic facial features, short stature, hypertelorism, cardiac anomalies, deafness, motor delay, and a bleeding diathesis. It is a genetically heterogeneous and relatively common syndrome, with an estimated incidence of 1 in 1000-2500 live births. Defects in RAF1 are the cause of LEOPARD syndrome type 2 (LEOPARD2) [MIM:611554]. LEOPARD syndrome is an autosomal dominant disorder allelic with Noonan syndrome. The acronym LEOPARD stands for lentigines, electrocardiographic conduction abnormalities, ocular hypertelorism, pulmonic stenosis, abnormalities of genitalia, retardation of growth, and deafness. |
| 序列相似性 | Belongs to the protein kinase superfamily. TKL Ser/Thr protein kinase family. RAF subfamily. Contains 1 phorbol-ester/DAG-type zinc finger. Contains 1 protein kinase domain. Contains 1 RBD (Ras-binding) domain. |
| 翻译后修饰 | Phosphorylated upon DNA damage, probably by ATM or ATR. Phosphorylation at Thr-269 increases its kinase activity. Phosphorylation at Ser-259 induces the interaction with YWHAZ and inactivates kinase activity. Dephosphorylation of Ser-259 by the complex containing protein phosphatase 1, SHOC2 and M-Ras/MRAS relieves inactivation, leading to stimulate RAF1 activity. |
| 细胞定位 | Cytoplasm. Cell membrane. Colocalizes with RGS14 and BRAF in both the cytoplasm and membranes. |

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