

Anti-FGF8 antibody [MM0291-8D24] ab89550

★★★★★ [1 Abreviews](#) [1 图像](#)

概述

产品名称	Anti-FGF8抗体[MM0291-8D24]
描述	小鼠单克隆抗体[MM0291-8D24] to FGF8
宿主	Mouse
经测试应用	适用于: IHC-P, WB, Neutralising
种属反应性	与反应: Human
免疫原	Recombinant full length Human FGF8 protein.
阳性对照	WB: Human placenta tissue lysate.
常规说明	<p>The Life Science industry has been in the grips of a reproducibility crisis for a number of years. Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets your needs before purchasing.</p> <p>If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be found below, along with publications, customer reviews and Q&As</p>

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.
存储溶液	Constituent: PBS
纯度	Protein G purified
纯化说明	The IgG fraction of culture supernatant was purified by Protein G affinity chromatography and filtered through a 0.2 µm filter.
克隆	单克隆
克隆编号	MM0291-8D24
同种型	IgG1

应用

The Abpromise guarantee

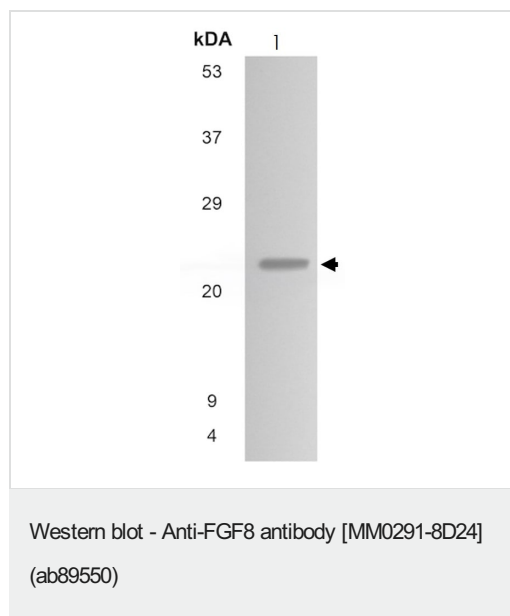
Abpromise™承诺保证使用ab89550于以下的经测试应用

“应用说明”部分 下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。

应用	Ab评论	说明
IHC-P	★★★★★ (1)	Use at an assay dependent concentration.
WB		1/500 - 1/1000. Predicted molecular weight: 27 kDa.
Neutralising		Use at an assay dependent concentration.

靶标	
功能	Stimulates growth of the cells in an autocrine manner. Mediates hormonal action on the growth of cancer cells.
疾病相关	<p>Defects in FGF8 are the cause of Kallmann syndrome type 6 (KAL6) [MIM:612702]. Kallmann syndrome is a disorder that associates hypogonadotropic hypogonadism and anosmia. Anosmia or hyposmia is related to the absence or hypoplasia of the olfactory bulbs and tracts. Hypogonadism is due to deficiency in gonadotropin-releasing hormone and probably results from a failure of embryonic migration of gonadotropin-releasing hormone-synthesizing neurons. In some patients other developmental anomalies can be present, which include renal agenesis, cleft lip and/or palate, selective tooth agenesis, and bimanual synkinesis. In some cases anosmia may be absent or inconspicuous.</p> <p>Defects in FGF8 are a cause of idiopathic hypogonadotropic hypogonadism (IHH) [MIM:146110]. IHH is defined as a deficiency of the pituitary secretion of follicle-stimulating hormone and luteinizing hormone, which results in the impairment of pubertal maturation and of reproductive function.</p>
序列相似性	Belongs to the heparin-binding growth factors family.
发展阶段	In adults expression is restricted to the gonads.
细胞定位	Secreted.

图片



Anti-FGF8 antibody [MM0291-8D24] (ab89550) at 1/500 dilution +
Human placenta tissue lysate

Predicted band size: 27 kDa

Observed band size: 23 kDa

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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- Response to your inquiry within 24 hours
- We provide support in Chinese, English, French, German, Japanese and Spanish
- Extensive multi-media technical resources to help you
- We investigate all quality concerns to ensure our products perform to the highest standards

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