# abcam

## Product datasheet

## Anti-EMX2 antibody ab110112

★★★★★ 2 Abreviews 1 References 画像数1

#### 製品の概要

製品名 Anti-EMX2 antibody

製品の詳細 Rabbit polyclonal to EMX2

由来種 Rabbit

アプリケーション 適用あり: IHC-P 種交差性 交差種: Human

交差が予測される動物種: Mouse 🔷

免疫原 Synthetic peptide derived from an internal sequence of Human EMX2.

ポジティブ・コントロール Human brain tissue

特記事項 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.

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

#### 製品の特性

製品の状態 Liquid

保存方法 Shipped at 4°C. Store at -20°C. Stable for 12 months at -20°C.

バッファー pH: 7.40

Preservative: 0.02% Sodium azide

Constituents: PBS, 0.88% Sodium chloride, 50% Glycerol (glycerin, glycerine)

精製度 Immunogen affinity purified

特記事項(精製) ab110112 was affinity purified from rabbit antiserum by affinity chromatography using epitope

specific immunogen.

ポリモノ ポリクローナル

アイソタイプ lgG **The Abpromise guarantee** <u>Abpromise保証は、</u>次のテスト済みアプリケーションにおけるab110112の使用に適用されます アプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご検討ください。

アプリケーション	Abreviews	特記事項
IHC-P	<b>★★★★☆ (1)</b>	1/50 - 1/100. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.

#### ターゲット情報

機能 Transcription factor, which in cooperation with EMX2, acts to generate the boundary between the

roof and archipallium in the developing brain. May function in combinations with OTX1/2 to specify

cell fates in the developing central nervous system.

組織特異性 Cerebral cortex.

**関連疾患** Defects in EMX2 are the cause of schizencephaly (SCHZC) [MIM:269160]. Schizencephaly is an

extremely rare human congenital disorder characterized by a full-thickness cleft within the cerebral hemispheres. These clefts are lined with gray matter and most commonly involve the parasylvian regions. Large portions of the cerebral hemispheres may be absent and replaced by cerebro-

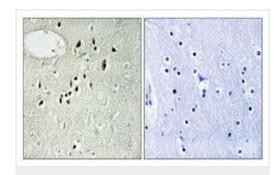
spinal fluid.

配列類似性 Belongs to the EMX homeobox family.

Contains 1 homeobox DNA-binding domain.

細胞内局在 Nucleus.

## 画像



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-EMX2 antibody (ab110112)

ab110112, at 1/50 dilution, staining EMX2 in paraffin embedded Human brain tissue by Immunohistochemistry in the presence (right panel) or absence (left panel) of immunizing peptide.

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

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