# abcam

# Product datasheet

# Anti-Junctional Adhesion Molecule 1/JAM-A antibody ab125886

★★★★★ 2 Abreviews 4 References 画像数 1

製品の概要

製品名 Anti-Junctional Adhesion Molecule 1/JAM-A antibody

製品の詳細 Rabbit polyclonal to Junctional Adhesion Molecule 1/JAM-A

由来種 Rabbit

アプリケーション **適用あり**: WB

**種交差性 交差種:** Mouse, Rat, Human

交差が予測される動物種: Chinese hamster 🔷

免疫原 Synthetic peptide corresponding to Mouse Junctional Adhesion Molecule 1/JAM-A aa 50-150

conjugated to keyhole limpet haemocyanin.

(Peptide available as ab156888)

ポジティブ・コントロール This antibody gave a positive signal in the following tissue lysates: Mouse Brain; Rat Brain;

Mouse Cortex; Rat Cortex; Mouse Cerebellum; Mouse Spinal Cord.

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 +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C or -

80°C. Avoid freeze / thaw cycle.

**バッファー** pH: 7.40

Preservative: 0.02% Sodium azide

Constituent: PBS

Batches of this product that have a concentration < 1mg/ml may have BSA added as a stabilising

agent. If you would like information about the formulation of a specific lot, please contact our

scientific support team who will be happy to help.

1

精製度 Immunogen affinity purified

**ポリ/モノ** ポリクローナル

アイソタイプ IgG

#### アプリケーション

**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ab125886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
WB	<b>★★★★☆ (1)</b>	Use a concentration of 1 µg/ml. Detects a band of approximately 42 kDa (predicted molecular weight: 32 kDa).

#### ターゲット情報

機能 Seems to plays a role in epithelial tight junction formation. Appears early in primordial forms of

cell junctions and recruits PARD3. The association of the PARD6-PARD3 complex may prevent the interaction of PARD3 with JAM1, thereby preventing tight junction assembly (By similarity). Plays a role in regulating monocyte transmigration involved in integrity of epithelial barrier. Involved in platelet activation. In case of orthoreovirus infection, serves as receptor for the virus.

**配列類似性** Belongs to the immunoglobulin superfamily.

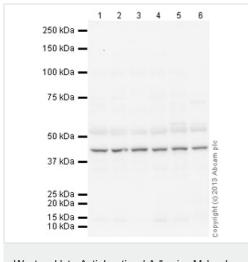
Contains 2 lg-like V-type (immunoglobulin-like) domains.

**翻訳後修飾** N-glycosylated.

細胞内局在 Cell junction > tight junction. Cell membrane. Localized at tight junctions of both epithelial and

endothelial cells.

#### 画像



Western blot - Anti-Junctional Adhesion Molecule
1/JAM-A antibody (ab125886)

**All lanes :** Anti-Junctional Adhesion Molecule 1/JAM-A antibody (ab125886) at 1 µg/ml

Lane 1: Brain (Mouse) Tissue Lysate

Lane 2: Brain (Rat) Tissue Lysate

Lane 3 : Mouse Cortex Tissue Lysate

Lane 4: Rat Cortex Tissue Lysate

Lane 5 : Cerebellum Mouse Tissue Lysate

Lane 6 : Spinal Cord (Mouse) Tissue Lysate

Lysates/proteins at 10 µg per lane.

### **Secondary**

All lanes: Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/10000

dilution

Developed using the ECL technique.

Performed under reducing conditions.

**Predicted band size:** 32 kDa **Observed band size:** 42 kDa

Additional bands at: 53 kDa, 99 kDa. We are unsure as to the

identity of these extra bands.

Exposure time: 3 minutes

Junctional Adhesion Molecule 1/JAM-A contains a number of potential glycosylation sites (SwissProt) which may explain its migration at a higher molecular weight than predicted. The predicted molecular weight of Junctional Adhesion Molecule 1/JAM-A is 32 kDa (SwissProt), however we expect to observe a banding pattern around 42 kDa. This blot was produced using a 4-12% Bistris gel under the MOPS buffer system. The gel was run at 200V for 50 minutes before being transferred onto a Nitrocellulose membrane at 30V for 70 minutes. The membrane was then blocked for an hour using 5% Bovine Serum Albumin before being incubated with ab125886 overnight at 4°C. Antibody binding was detected using an anti-rabbit antibody conjugated to HRP, and visualised using ECL development solution.

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