# abcam

### Product datasheet

# Anti-PRPF8/Prp8 antibody [2834C1a] ab51366

2 References 画像数 4

製品の概要

製品名 Anti-PRPF8/Prp8 antibody [2834C1a]

製品の詳細 Mouse monoclonal [2834C1a] to PRPF8/Prp8

由来種 Mouse

アプリケーション 適用あり: Flow Cyt, WB, IHC-P

種交差性 交差種: Mouse, Human

免疫原 Recombinant fragment corresponding to Human PRPF8/Prp8.

ポジティブ・コントロール NIH3T3 whole cell lysate, HeLa whole cell lysate This antibody gave a positive result in IHC in the

following FFPE tissue: Human normal cervix.

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 long

term.

**バッファー** pH: 7.40

Preservative: 0.05% Sodium azide Constituents: 1% BSA, PBS

精製度 Protein G purified

特記事項(精製) Filtered through a 0.22 μm membrane.

**ポリ/モノ** モノクローナル **クローン名** 2834C1a

アイソタイプ lgG1

1

**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ab51366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   |
|----------|-----------|--|
| Flow Cyt |           | Use 2µg for 10 <sup>6</sup> cells.  ab170190 - Mouse monoclonal lgG1, is suitable for use as an isotype control with this antibody.            |
| WB       |           | 1/100. Predicted molecular weight: 273 kDa.  |
| IHC-P    |           | Use a concentration of 5 µg/ml. 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    | Central component of the spliceosome, which may play a role in aligning the pre-mRNA 5'- and 3'- exons for ligation. Interacts with U5 snRNA, and with pre-mRNA 5'-splice sites in B spliceosomes and 3'-splice sites in C spliceosomes.  |  |
|-------|---|--|
| 組織特異性 | Widely expressed.   |  |
| 関連疾患  | Defects in PRPF8 are the cause of retinitis pigmentosa type 13 (RP13) [MIM:600059]. RP leads to degeneration of retinal photoreceptor cells. Patients typically have night vision blindness and loss of midperipheral visual field. As their condition progresses, they lose their far peripheral visual field and eventually central vision as well. RP13 inheritance is autosomal dominant. |  |
| 配列類似性 | Contains 1 MPN (JAB/Mov34) domain.  |  |

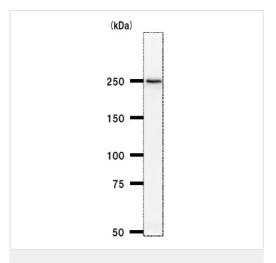
ドメイン The MPN domain has structural similarity with viral ribonucleases and RNase H, but unlike

RNases, it does not bind any metal ions.

**翻訳後修飾** Phosphorylated upon DNA damage, probably by ATM or ATR.

細胞内局在 Nucleus speckle.

#### 画像



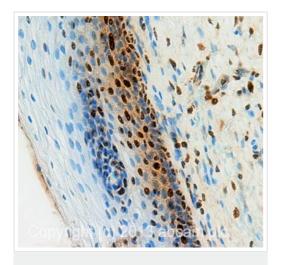
Western blot - Anti-PRPF8/Prp8 antibody [2834C1a] (ab51366)

Anti-PRPF8/Prp8 antibody [2834C1a] (ab51366) at 1/100 dilution  $\pm$  HeLa whole cell lysate at 25  $\mu g$ 

#### Secondary

Anti mouse IgG antibody at 1/2500 dilution

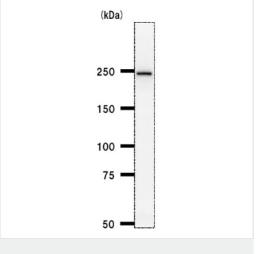
**Predicted band size:** 273 kDa **Observed band size:** 250 kDa



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-PRPF8/Prp8 antibody [2834C1a] (ab51366)

IHC image of PRPF8/Prp8 staining in Human Normal Cervix formalin fixed paraffin embedded tissue section, performed on a Leica Bond™ system using the standard protocol F. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH6, epitope retrieval solution 1) for 20 mins. The section was then incubated with ab51366, 5µg/ml, for 15 mins at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

For other IHC staining systems (automated and non-automated) customers should optimize variable parameters such as antigen retrieval conditions, primary antibody concentration and antibody incubation times.



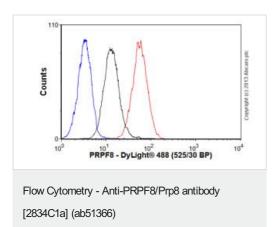
Anti-PRPF8/Prp8 antibody [2834C1a] (ab51366) at 1/100 dilution + NIH3T3 whole cell lysate at 25 µg

#### Secondary

Anti mouse IgG antibody at 1/2500 dilution

Predicted band size: 273 kDa Observed band size: 250 kDa

Western blot - Anti-PRPF8/Prp8 antibody [2834C1a] (ab51366)



Overlay histogram showing HeLa cells stained with ab51366 (red line). The cells were fixed with 80% methanol (5 min) and then permeabilized with 0.1% PBS-Tween for 20 min. The cells were then incubated in 1x PBS / 10% normal goat serum / 0.3M glycine to block non-specific protein-protein interactions followed by the antibody (ab51366, 2µg/1x10<sup>6</sup> cells) for 30 min at 22°C. The secondary antibody used was DyLight® 488 goat anti-mouse lgG (H+L) (ab96879) at 1/500 dilution for 30 min at 22°C. Isotype control antibody (black line) was mouse lgG1 [ICIGG1] (ab91353, 2µg/1x10<sup>6</sup> cells) used under the same conditions. Unlabelled sample (blue line). Acquisition of >5,000 events were collected using a 20mW Argon ion laser (488nm) and 525/30 bandpass filter.

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

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