

FITC Anti-Human Serum Albumin antibody ab19182

★★★★☆ [1 Abreviews](#) [3 References](#)

製品の概要

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| 製品名 | FITC Anti-Human Serum Albumin antibody |
| 製品の詳細 | FITC Goat polyclonal to Human Serum Albumin |
| 由来種 | Goat |
| 標識 | FITC. Ex: 493nm, Em: 528nm |
| アプリケーション | 適用あり: IHC-P, ICC |
| 種交差性 | 交差種: Human |
| 免疫原 | Full length protein. This information is proprietary to Abcam and/or its suppliers. |
| 特記事項 | <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> |

製品の特性

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| 製品の状態 | Liquid |
| 保存方法 | Shipped at 4°C. Store at +4°C. |
| バッファー | pH: 6.8 Preservative: 0.1% Sodium azide Constituents: PBS, 0.2% BSA |
| 精製度 | Immunogen affinity purified |
| 特記事項(精製) | The antibody was isolated by affinity chromatography using antigen coupled to agarose beads and conjugated to fluorescein isothiocyanate (FITC). |
| 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 **Abpromise保証は、次のテスト済みアプリケーションにおけるab19182の使用に適用されず**
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 | | Use at an assay dependent concentration. |
| ICC | | Use at an assay dependent concentration. |

ターゲット情報

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|-------|--|
| 機能 | Serum albumin, the main protein of plasma, has a good binding capacity for water, Ca(2+), Na(+), K(+), fatty acids, hormones, bilirubin and drugs. Its main function is the regulation of the colloidal osmotic pressure of blood. Major zinc transporter in plasma, typically binds about 80% of all plasma zinc. |
| 組織特異性 | Plasma. |
| 関連疾患 | Defects in ALB are a cause of familial dysalbuminemic hyperthyroxinemia (FDH) [MIM:103600]. FDH is a form of euthyroid hyperthyroxinemia that is due to increased affinity of ALB for T(4). It is the most common cause of inherited euthyroid hyperthyroxinemia in Caucasian population. |
| 配列類似性 | Belongs to the ALB/AFP/VDB family. Contains 3 albumin domains. |
| 翻訳後修飾 | Kenitra variant is partially O-glycosylated at Thr-620. It has two new disulfide bonds Cys-600 to Cys-602 and Cys-601 to Cys-606. Glycated in diabetic patients. Phosphorylation sites are present in the extracellular medium. Acetylated on Lys-223 by acetylsalicylic acid. |
| 細胞内局在 | Secreted. |

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

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