## abcam

### Product datasheet

# Native Human Serum Albumin protein (Rhodamine) ab8031

#### 1 References

**Description** 

Product name Native Human Serum Albumin protein (Rhodamine)

Expression system Native

Protein length Full length protein

Animal free No

Nature Native

Species Human
Predicted molecular weight 69 kDa

**Conjugation** Rhodamine. Ex: 550nm, Em: 570nm

**Description** Native Human Human Serum Albumin protein (Rhodamine)

#### **Specifications**

Our **Abpromise guarantee** covers the use of **ab8031** in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

**Applications** Dot blot

Flow Cytometry

Immunomicroscopy

Form Lyophilized

#### **Preparation and Storage**

Stability and Storage Shipped at 4°C. Add glycerol to a final volume of 50% for extra stability and aliquot. Store at -

 $20\,^{\circ}\text{C}\,.$  Avoid freeze / thaw cycle.

Preservative: 0.01% Sodium azide

Constituents: 0.42% Potassium phosphate, 0.87% Sodium chloride, 1% BSA

**Reconstitution** Store vial at 4° C prior to restoration. Restore with 1.0 mL of deionized water.

#### **General Info**

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**Function** 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.

**Tissue specificity** Plasma.

Involvement in disease Defects in ALB are a cause of familial dysalbuminemic hyperthyroxinemia (FDH) [MIM:103600].

 $\label{eq:fdh} FDH \ is \ a \ form \ of \ euthyroid \ hyperthyroxinemia \ that \ is \ due \ to \ increased \ affinity \ of \ ALB \ for \ T(4). \ It \ is$ 

the most common cause of inherited euthyroid hyperthyroxinemia in Caucasian population.

**Sequence similarities** Belongs to the ALB/AFP/VDB family.

Contains 3 albumin domains.

Post-translational

modifications

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 extracelllular medium.

Acetylated on Lys-223 by acetylsalicylic acid.

Cellular localization

Secreted.

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