

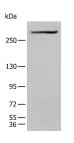
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# **SPTA1 Polyclonal Antibody**

E-AB-19527 Catalog No. Reactivity H,M Store at -20°C. Avoid freeze / thaw cycles. Rabbit **Storage** Host **Applications** WB,IHC,ELISA **Isotype IgG** 

**Note:** Centrifuge before opening to ensure complete recovery of vial contents.

## **Images**



Western blot analysis of Mouse heart tissue lysate using SPTA1 Polyclonal Antibody at dilution of 1:550



Immunohistochemistry of paraffinembedded Human gastric cancer tissue using SPTA1 Polyclonal Antibody at dilution of  $1:80(\times 200)$ 

## **Immunogen Information**

Synthetic peptide of human SPTA1 **Immunogen** 

NP003117 **Gene Accession** P02549 **Swissprot** 

**Synonyms** Alpha I spectrin, EL 2, EL2, Elliptocytosis

2,HPP,HS3,SPH3,SPTA 1,SPTA,SPTA1,SPTA1

#### **Product Information**

Calculated MW 280 kDa

**Observed MW** Refer to figures

**Buffer** PBS with 0.05% NaN3 and 40% Glycerol,pH7.4

**Purify** Antigen affinity purification

Dilution WB 1:500-1:2000, IHC 1:60-1:450, ELISA

1:5000-1:10000

#### **Background**

Spectrin is an actin crosslinking and molecular scaffold protein that links the plasma membrane to the actin cytoskeleton, and functions in the determination of cell shape, arrangement of transmembrane proteins, and organization of organelles. It is a tetramer made up of alpha-beta dimers linked in a head-to-head arrangement. This gene is one member of a family of alpha-spectrin genes. The encoded protein is primarily composed of 22 spectrin repeats which are involved in dimer formation. It forms weaker tetramer interactions than non-erythrocytic alpha spectrin, which may increase the plasma membrane elasticity and deformability of red blood cells. Mutations in this gene result in a variety of hereditary red blood cell disorders, including elliptocytosis type 2, pyropoikilocytosis, and spherocytic hemolytic anemia.