

# FTL, Biotinylated

Peptide-affinity purified goat antibody Catalog # AF1447b

#### **Product Information**

**Application** WB, IHC, Pep-ELISA

Primary Accession P02792

Other Accession NP 000137, 2512, 14325 (mouse), 29292 (rat)

Reactivity Human

**Predicted** Mouse, Rat, Dog

Host Goat
Clonality Polyclonal
Isotype IgG
Calculated MW 20020

### **Additional Information**

**Gene ID** 2512

Other Names Ferritin light chain, Ferritin L subunit, FTL

**Dilution** WB~~1:1000 IHC~~1:100~500 Pep-ELISA~~N/A

Format 0.5 mg IgG/ml in Tris saline (20mM Tris pH7.3, 150mM NaCl), 0.02% sodium

azide, with 0.5% bovine serum albumin

**Storage** Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store

at -20°C in small aliquots to prevent freeze-thaw cycles.

**Precautions** FTL, Biotinylated is for research use only and not for use in diagnostic or

therapeutic procedures.

## **Protein Information**

Name FTL

**Function** Stores iron in a soluble, non-toxic, readily available form. Important for iron

homeostasis. Iron is taken up in the ferrous form and deposited as ferric hydroxides after oxidation. Also plays a role in delivery of iron to cells. Mediates iron uptake in capsule cells of the developing kidney (By similarity).

Delivery to lysosomes by the cargo receptor NCOA4 for autophagic

degradation and release or iron (PubMed:24695223).

**Cellular Location** Cytoplasmic vesicle, autophagosome. Cytoplasm

{ECO:0000250 | UniProtKB:P29391}. Autolysosome

## **Background**

This gene encodes the light subunit of the ferritin protein. Ferritin is the major intracellular iron storage protein in prokaryotes and eukaryotes. It is composed of 24 subunits of the heavy and light ferritin chains. Variation in ferritin subunit composition may affect the rates of iron uptake and release in different tissues. A major function of ferritin is the storage of iron in a soluble and nontoxic state. Defects in this light chain ferritin gene are associated with several neurodegenerative diseases and hyperferritinemia-cataract syndrome. This gene has multiple pseudogenes.

#### References

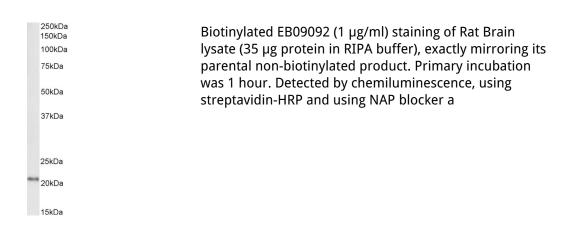
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## **Images**



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