abcam

Product datasheet

Anti-Ceruloplasmin antibody [EPSISR6-42] ab157452

Recombinant RabMAb

1 References 3 Images

Overview

Product name Anti-Ceruloplasmin antibody [EPSISR6-42]

Description Rabbit monoclonal [EPSISR6-42] to Ceruloplasmin

Host species Rabbit

Tested applications Suitable for: ICC/IF, WB

Unsuitable for: IHC-P or IP

Reacts with: Human Species reactivity

Immunogen Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

Positive control Human plasma and serum; HepG2 cells.

General notes This product is a recombinant monoclonal antibody, which offers several advantages including:

- High batch-to-batch consistency and reproducibility

- Improved sensitivity and specificity

- Long-term security of supply

- Animal-free production

For more information see here.

Our RabMAb® technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**® **patents**.

Mouse, Rat: We have preliminary internal testing data to indicate this antibody may not react with

these species. Please contact us for more information.

Properties

Form Liquid

Storage instructions Shipped at 4°C. Store at -20°C.

Storage buffer

Preservative: 0.01% Sodium azide

Constituents: 40% Glycerol (glycerin, glycerine), 0.05% BSA, 9% PBS

Purity Protein A purified

Clonality Monoclonal Clone number EPSISR6-42

Isotype IgG

Applications

The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab157452 in the following tested applications.

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

Application	Abreviews	Notes
ICC/IF		1/100 - 1/250.
WB		1/1000 - 1/5000. Predicted molecular weight: 122 kDa.

Application notes

Is unsuitable for IHC-P or IP.

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Function Ceruloplasmin is a blue, copper-binding (6-7 atoms per molecule) glycoprotein. It has ferroxidase

activity oxidizing Fe(2+) to Fe(3+) without releasing radical oxygen species. It is involved in iron

transport across the cell membrane.

Tissue specificity Expressed by the liver and secreted in plasma.

Involvement in disease Defects in CP are the cause of aceruloplasminemia (ACERULOP) [MIM:604290]. It is an

autosomal recessive disorder of iron metabolism characterized by iron accumulation in the brain as well as visceral organs. Clinical features consist of the triad of retinal degeneration, diabetes

mellitus and neurological disturbances.

Note=Ceruloplasmin levels are decreased in Wilson disease, in which copper cannot be

incorporated into ceruloplasmin in liver because of defects in the copper-transporting ATPase 2.

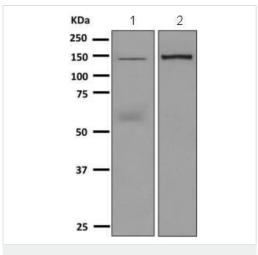
Sequence similaritiesBelongs to the multicopper oxidase family.

Contains 3 F5/8 type A domains.

Contains 6 plastocyanin-like domains.

Cellular localization Secreted.

Images



Western blot - Anti-Ceruloplasmin antibody [EPSISR6-42] (ab157452)

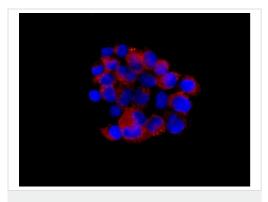
All lanes : Anti-Ceruloplasmin antibody [EPSISR6-42] (ab157452) at 1/1000 dilution

Lane 1: Human plasma cell lysate

Lane 2: Human serum cell lysate

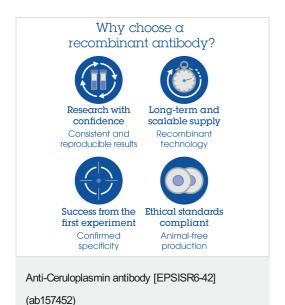
Lysates/proteins at 10 µg per lane.

Predicted band size: 122 kDa



Immunocytochemistry/ Immunofluorescence - Anti-Ceruloplasmin antibody [EPSISR6-42] (ab157452)

Immunofluorescent analysis of HepG2 cells labeling Ceruloplasmin with ab157452 at 1/100 dilution.



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

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