abcam

Product datasheet

HRP Anti-Fatty Acid Synthase antibody [EPR7466] ab196854



Recombinant

RabMAb

3 References 3 Images

Overview

Product name HRP Anti-Fatty Acid Synthase antibody [EPR7466]

Description HRP Rabbit monoclonal [EPR7466] to Fatty Acid Synthase

Host species Rabbit Conjugation HRP

Tested applications Suitable for: WB **Species reactivity** Reacts with: Human

Predicted to work with: Mouse, Rat

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

Positive control WB: HeLa, HEK293 and A549 whole cell lysates.

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**.

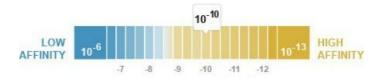
Properties

Form Liquid

Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C. Storage instructions

Avoid freeze / thaw cycle. Store In the Dark.

 $K_D = 1.34 \times 10^{-10} M$ Dissociation constant (K_D)



Learn more about K_D

Storage buffer pH: 7.40

Preservative: 0.1% Proclin 300 Solution

Constituents: 30% Glycerol (glycerin, glycerine), 1% BSA, PBS

Purity Protein A purified

Clonality Monoclonal
Clone number EPR7466

Isotype IgG

Applications

The Abpromise guarantee Our Abpromise guarantee covers the use of ab196854 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
WB		1/5000. Detects a band of approximately 273 kDa (predicted molecular weight: 273 kDa).

Function Fatty acid synthetase catalyzes the formation of long-chain fatty acids from acetyl-CoA, malonyl-

CoA and NADPH. This multifunctional protein has 7 catalytic activities and an acyl carrier protein.

Tissue specificity Ubiquitous. Prominent expression in brain, lung, and liver.

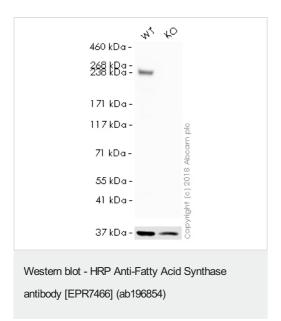
Sequence similarities Contains 1 acyl carrier domain.

Cellular localization Cytoplasm. Melanosome. Identified by mass spectrometry in melanosome fractions from stage I

to stage IV.

Images

Target



All lanes : HRP Anti-Fatty Acid Synthase antibody [EPR7466] (ab196854) at 1/5000 dilution

Lane 1: Wild-type HAP1 whole cell lysate

Lane 2: FASN (Fatty Acid Synthase) knockout HAP1 whole cell

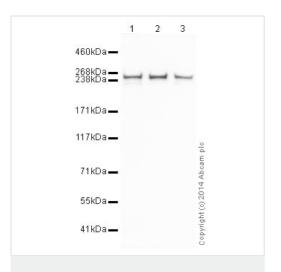
lysate

Lysates/proteins at 20 µg per lane.

Predicted band size: 273 kDa Observed band size: 260 kDa

Exposure time: 1 minute

ab196854 was shown to specifically react with Fatty Acid Synthase in wild-type HAP1 cells as signal was lost in FASN (Fatty Acid Synthase) knockout cells. Wild-type and FASN (Fatty Acid Synthase) knockout samples were subjected to SDS-PAGE. Ab196854 and ab184095 (Mouse monoclonal [mAbcam 9484] to GAPDH - Loading Control (Alexa Fluor 680) loading control) were incubated overnight at 4°C at 1/5000 dilution and 1/1000 dilution respectively. The loading control was imaged using the Licor Odyssey CLx prior to blots being developed with ECL technique.



Western blot - HRP Anti-Fatty Acid Synthase antibody [EPR7466] (ab196854)

All lanes : HRP Anti-Fatty Acid Synthase antibody [EPR7466] (ab196854) at 1/5000 dilution

Lane 1: HeLa whole cell lysate (ab150035)

Lane 2 : HEK293 (Human embryonic kidney cell line) Whole Cell Lysate

Lane 3: A549 (Human lung adenocarcinoma epithelial cell line) Whole Cell Lysate

Lysates/proteins at 10 µg per lane.

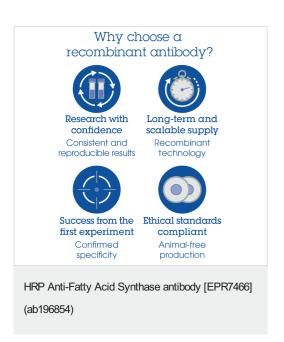
Developed using the ECL technique.

Performed under reducing conditions.

Predicted band size: 273 kDa **Observed band size:** 273 kDa

Exposure time: 8 minutes

This blot was produced using a 3-8% Tris Acetate gel under the TA buffer system. The gel was run at 150V for 60 minutes before being transferred onto a Nitrocellulose membrane at 30V for 70 minutes. The membrane was then blocked for an hour using 2% Bovine Serum Albumin before being incubated with ab196854 overnight at 4°C. Antibody binding was visualised using ECL development solution ab133406.



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