

## Product datasheet

# Anti-NSDHL antibody [EPR14489(2)] ab199730

**KO VALIDATED** Recombinant RabMAB

6 Images

### Overview

<b>Product name</b>	Anti-NSDHL antibody [EPR14489(2)]
<b>Description</b>	Rabbit monoclonal [EPR14489(2)] to NSDHL
<b>Host species</b>	Rabbit
<b>Tested applications</b>	<b>Suitable for:</b> WB, IP
<b>Species reactivity</b>	<b>Reacts with:</b> Mouse, Rat, Human
<b>Immunogen</b>	Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.
<b>Positive control</b>	WB: HEK293T, A431, HeLa, HepG2, NIH/3T3, PC-12 and C6 whole cell lysates; Human fetal brain lysate. IP: Human fetal brain whole cell lysate.
<b>General notes</b>	<p>This product is a recombinant monoclonal antibody, which offers several advantages including:</p> <ul style="list-style-type: none"> <li>- High batch-to-batch consistency and reproducibility</li> <li>- Improved sensitivity and specificity</li> <li>- Long-term security of supply</li> <li>- Animal-free production</li> </ul> <p>For more information <a href="#">see here</a>.</p> <p>Our RabMAB<sup>®</sup> technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to <a href="#">RabMAB<sup>®</sup> patents</a>.</p>

### Properties

<b>Form</b>	Liquid
<b>Storage instructions</b>	Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C long term. Avoid freeze / thaw cycle.
<b>Storage buffer</b>	<p>pH: 7.2</p> <p>Preservative: 0.01% Sodium azide</p> <p>Constituents: 59% PBS, 40% Glycerol (glycerin, glycerine), 0.05% BSA</p>
<b>Purity</b>	Protein A purified
<b>Clonality</b>	Monoclonal
<b>Clone number</b>	EPR14489(2)
<b>Isotype</b>	IgG

## Applications

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**The Abpromise guarantee** Our [Abpromise guarantee](#) covers the use of ab199730 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		1/1000. Detects a band of approximately 42 kDa (predicted molecular weight: 42 kDa).
IP		1/50.

## Target

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**Tissue specificity** Brain, heart, liver, lung, kidney, skin and placenta.

**Pathway** Steroid biosynthesis; zymosterol biosynthesis; zymosterol from lanosterol: step 4/6.

**Involvement in disease** Defects in NSDHL are the cause of congenital hemidysplasia with ichthyosiform erythroderma and limb defects (CHILD) [MIM:308050]. CHILD is an X-linked dominant disorder of lipid metabolism with disturbed cholesterol biosynthesis, which typically results in male lethality. Clinically, it is characterized by congenital, unilateral, ichthyosiform erythroderma with striking lateralization, sharp midline demarcation, and ipsilateral limb defects and hypoplasia of the body. Limbs defects range from hypoplasia of digits or ribs to complete amelia, often including scoliosis.

Defects in NSDHL are the cause of CK syndrome (CKS) [MIM:300831]. CKS is a disorder characterized by mild to severe cognitive impairment, seizures, microcephaly, cerebral cortical malformations, dysmorphic facial features, and thin body habitus.

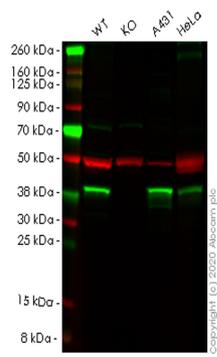
**Sequence similarities** Belongs to the 3-beta-HSD family.

**Cellular localization** Membrane.

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## Images

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Western blot - Anti-NSDHL antibody [EPR14489(2)] (ab199730)

**All lanes** : Anti-NSDHL antibody [EPR14489(2)] (ab199730) at 1/1000 dilution

**Lane 1** : Wild-type HEK293T cell lysate

**Lane 2** : NSDHL knockout HEK293T cell lysate

**Lane 3** : A431 cell lysate

**Lane 4** : HeLa cell lysate

Lysates/proteins at 20 µg per lane.

### Secondary

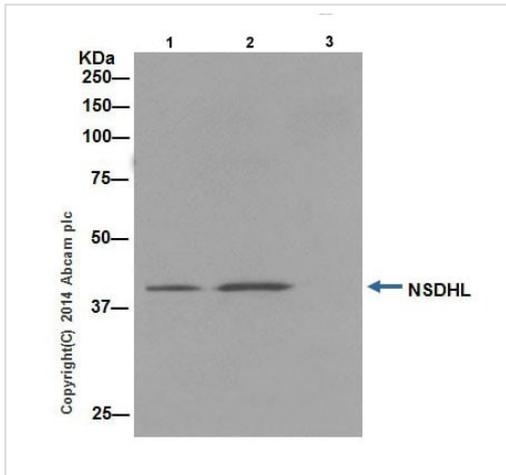
**All lanes** : Goat anti-Rabbit IgG H&L (IRDye® 800CW) preadsorbed (ab216773) at 1/10000 dilution

**Predicted band size:** 42 kDa

**Observed band size:** 38 kDa

**Lanes 1-4:** Merged signal (red and green). Green - ab199730 observed at 38 kDa. Red - loading control ab7291 observed at 50 kDa.

ab199730 Anti-NSDHL antibody [EPR14489(2)] was shown to specifically react with NSDHL in wild-type HEK293T cells. Loss of signal was observed when knockout cell line ab266682 (knockout cell lysate ab258082) was used. Wild-type and NSDHL knockout samples were subjected to SDS-PAGE. ab199730 and Anti-alpha Tubulin antibody [DM1A] - Loading Control (ab7291) were incubated overnight at 4°C at 1 in 1000 dilution and 1 in 20000 dilution respectively. Blots were developed with Goat anti-Rabbit IgG H&L (IRDye® 800CW) preadsorbed (ab216773) and Goat anti-Mouse IgG H&L (IRDye® 680RD) preadsorbed (ab216776) secondary antibodies at 1 in 20000 dilution for 1 hour at room temperature before imaging.



Immunoprecipitation - Anti-NSDHL antibody [EPR14489(2)] (ab199730)

NSDHL was immunoprecipitated from 1 mg of Human fetal brain whole cell lysate with ab199730 at 1/50 dilution.

Lane 1: Human fetal brain whole cell lysate 10ug (Input).

Lane 2: ab199730 IP in Human fetal brain whole cell lysate.

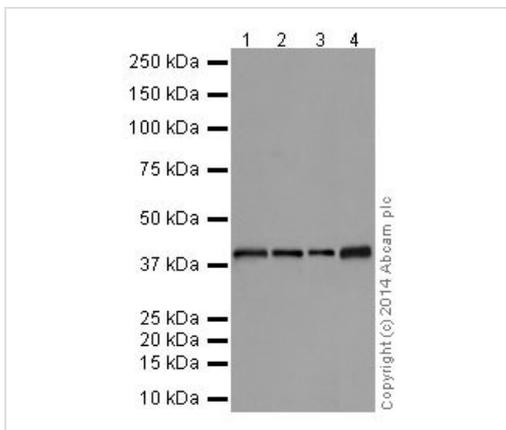
Lane 3: Rabbit monoclonal IgG (ab172730) instead of ab199730 in Human fetal brain whole cell lysate.

Western blot was performed from the immunoprecipitate using ab199730 at 1/1000 dilution.

Anti-Rabbit IgG (HRP), specific to the non-reduced form of IgG, was used as secondary antibody at 1/1500.

Blocking and dilution buffer and concentration: 5% NFDm/TBST.

Exposure time: 5 seconds.



Western blot - Anti-NSDHL antibody [EPR14489(2)] (ab199730)

**All lanes :** Anti-NSDHL antibody [EPR14489(2)] (ab199730) at 1/1000 dilution

**Lane 1 :** A431 (Human epidermoid carcinoma) whole cell lysate

**Lane 2 :** HeLa (Human epithelial cells from cervix adenocarcinoma) whole cell lysate

**Lane 3 :** HepG2 (Human liver hepatocellular carcinoma) whole cell lysate

**Lane 4 :** Human fetal brain lysate

Lysates/proteins at 10 µg per lane.

### Secondary

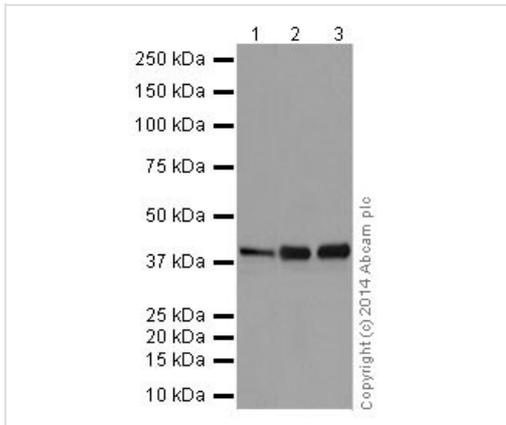
**All lanes :** Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/1000 dilution

**Predicted band size:** 42 kDa

**Observed band size:** 42 kDa

**Exposure time:** 5 seconds

Blocking/Dilution buffer: 5% NFDm/TBST.



Western blot - Anti-NSDHL antibody [EPR14489(2)] (ab199730)

**All lanes :** Anti-NSDHL antibody [EPR14489(2)] (ab199730) at 1/1000 dilution

**Lane 1 :** NIH/3T3 (Mouse embryo fibroblast cells) whole cell lysate

**Lane 2 :** PC-12 (Rat adrenal gland pheochromocytoma) whole cell lysate

**Lane 3 :** C6 (Rat glial tumor cells) whole cell lysate

Lysates/proteins at 10 µg per lane.

### Secondary

**All lanes :** Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/1000 dilution

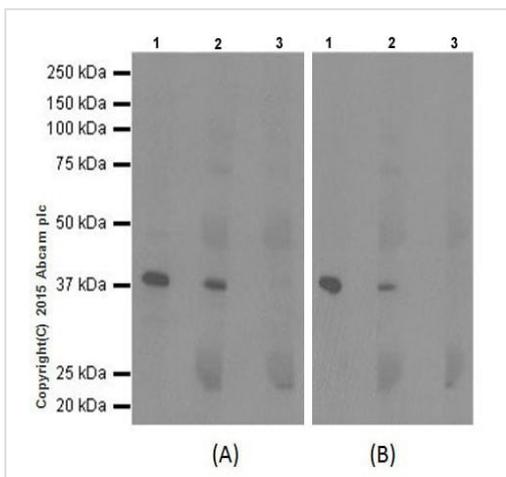
**Predicted band size:** 42 kDa

**Observed band size:** 42 kDa

**Exposure time:** 10 seconds

Blocking buffer: 5% NFDm/TBST.

Dilution buffer: 5% NFDm /TBST or 1%BSA /TBST.



Immunoprecipitation - Anti-NSDHL antibody [EPR14489(2)] (ab199730)

NSDHL was immunoprecipitated from 1 mg of HeLa (Human epithelial cells from cervix adenocarcinoma) whole cell lysate with ab199730 at 1/50 dilution.

Lane 1: HeLa whole cell lysate 10ug (Input).

Lane 2: ab199730 IP in HeLa whole cell lysate.

Lane 3: Rabbit monoclonal IgG (ab172730) instead of ab199730 in HeLa whole cell lysate.

Western blot was performed from the immunoprecipitate using ab199730 at 1/1000 dilution (Panel A) or ab190353 at 1/1000 dilution (Panel B).

Anti-Rabbit IgG (HRP), specific to the non-reduced form of IgG, was used as secondary antibody at 1/1500.

Blocking and dilution buffer and concentration: 5% NFDm/TBST.

Exposure time: 30 seconds (Panel A and B).

ab199730 was used to immunoprecipitate NSDHL. These precipitates were resolved with [ab190353](#).

Why choose a recombinant antibody?



- Research with confidence**  
Consistent and reproducible results
- Long-term and scalable supply**  
Recombinant technology
- Success from the first experiment**  
Confirmed specificity
- Ethical standards compliant**  
Animal-free production

Anti-NSDHL antibody [EPR14489(2)] (ab199730)

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

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