abcam

Product datasheet

Anti-L1CAM antibody [EPR23241-224] - BSA and Azide free ab272321





RabMAb

1 References 6 Images

Overview

Product name Anti-L1CAM antibody [EPR23241-224] - BSA and Azide free

Description Rabbit monoclonal [EPR23241-224] to L1CAM - BSA and Azide free

Host species Rabbit

Tested applications Suitable for: Flow Cyt, IP, ICC/IF, WB, IHC-P

Species reactivity Reacts with: Human

Immunogen Recombinant fragment. This information is proprietary to Abcam and/or its suppliers.

Positive control WB: Human brain tissue lysate; A375 and MCF7 whole cell lysates. IHC-P: Human kidney, spleen

and cerebellum tissue. Flow Cyt: A375 cells. IP: A375 whole cell lysate.

General notes ab272321 is the carrier-free version of <u>ab270455</u>.

Our <u>carrier-free</u> antibodies are typically supplied in a PBS-only formulation, purified and free of BSA, sodium azide and glycerol. The carrier-free buffer and high concentration allow for increased conjugation efficiency.

This conjugation-ready format is designed for use with fluorochromes, metal isotopes, oligonucleotides, and enzymes, which makes them ideal for antibody labelling, functional and cell-based assays, flow-based assays (e.g. mass cytometry) and Multiplex Imaging applications.

Use our **conjugation kits** for antibody conjugates that are ready-to-use in as little as 20 minutes with <1 minute hands-on-time and 100% antibody recovery: available for fluorescent dyes, HRP, biotin and gold.

This product is compatible with the Maxpar[®] Antibody Labeling Kit from Fluidigm, without the need for antibody preparation. Maxpar[®] is a trademark of Fluidigm Canada Inc.

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**.

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Properties

Form Liquid

Storage instructions Shipped at 4°C. Store at +4°C. Do Not Freeze.

Storage buffer pH: 7.2

Constituent: PBS

Carrier free Yes

Purity Protein A purified

Clonality Monoclonal

Clone number EPR23241-224

Isotype IgG

Applications

The Abpromise guarantee Our Abpromise guarantee covers the use of ab272321 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
Flow Cyt		Use at an assay dependent concentration.
IP		Use at an assay dependent concentration.
ICC/IF		Use at an assay dependent concentration.
WB		Use at an assay dependent concentration. Detects a band of approximately 250, 80 kDa (predicted molecular weight: 140 kDa).
IHC-P		Use at an assay dependent concentration. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.

Target

Function Cell adhesion molecule with an important role in the development of the nervous system. Involved

in neuron-neuron adhesion, neurite fasciculation, outgrowth of neurites, etc. Binds to axonin on

neurons.

Involvement in diseaseDefects in L1CAM are the cause of hydrocephalus due to stenosis of the aqueduct of Sylvius

(HSAS) [MIM:307000]. Hydrocephalus is a condition in which abnormal accumulation of cerebrospinal fluid in the brain causes increased intracranial pressure inside the skull. This is usually due to blockage of cerebrospinal fluid outflow in the brain ventricles or in the subarachnoid space at the base of the brain. In children is typically characterized by enlargement of the head, prominence of the forehead, brain atrophy, mental deterioration, and convulsions. In adults the syndrome includes incontinence, imbalance, and dementia. HSAS is characterized by mental retardation and enlarged brain ventricles.

Defects in L1CAM are the cause of mental retardation-aphasia-shuffling gait-adducted thumbs syndrome (MASA) [MIM:303350]; also known as corpus callosum hypoplasia, psychomotor retardation, adducted thumbs, spastic paraparesis, and hydrocephalus or CRASH syndrome. MASA is an X-linked recessive syndrome with a highly variable clinical spectrum. Main clinical features include spasticity and hyperreflexia of lower limbs, shuffling gait, mental retardation, aphasia and adducted thumbs. The features of spasticity have been referred to as complicated spastic paraplegia type 1 (SPG1). Some patients manifest corpus callosum hypoplasia and hydrocephalus. Inter- and intrafamilial variability is very wide, such that patients with hydrocephalus, MASA, SPG1, and agenesis of corpus callosum can be present within the same family.

Defects in L1CAM are the cause of spastic paraplegia X-linked type 1 (SPG1) [MIM:303350]. Spastic paraplegia is a degenerative spinal cord disorder characterized by a slow, gradual, progressive weakness and spasticity of the lower limbs.

Note=Defects in L1CAM may contribute to Hirschsprung disease by modifying the effects of Hirschsprung disease-associated genes to cause intestinal aganglionosis.

Defects in L1CAM are a cause of partial agenesis of the corpus callosum (ACCPX) [MIM:304100]. A syndrome characterized by partial corpus callosum agenesis, hypoplasia of inferior vermis and cerebellum, mental retardation, seizures and spasticity. Other features include microcephaly, unusual facies, and Hirschsprung disease in some patients.

Sequence similarities

Belongs to the immunoglobulin superfamily. L1/neurofascin/NgCAM family. Contains 5 fibronectin type-III domains.

Contains 6 lg-like C2-type (immunoglobulin-like) domains.

Cellular localization

Cell membrane.

Images



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-L1CAM antibody

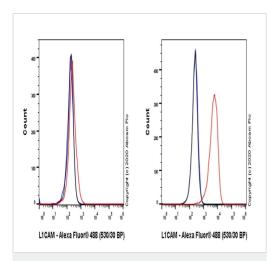
[EPR23241-224] - BSA and Azide free (ab272321)

Immunohistochemical analysis of paraffin-embedded Human cerebellum tissue labeling L1CAM with <u>ab270455</u> at 1/100 dilution followed by a ready to use Rabbit specific IHC polymer detection kit HRP/DAB (<u>ab209101</u>). Positive staining on the molecular layer of the human cerebellum (PMID: 12514225). The section was incubated with <u>ab270455</u> for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND[®] RX instrument. Counterstained with Hematoxylin.

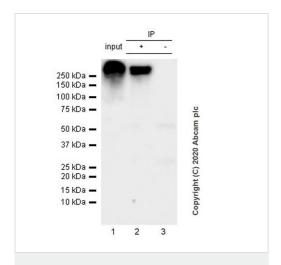
Secondary antibody only control: Secondary antibody is a ready to use Rabbit specific IHC polymer detection kit HRP/DAB (ab209101).

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution2) for 20 mins

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (**ab270455**).



Flow Cytometry - Anti-L1CAM antibody [EPR23241-224] - BSA and Azide free (ab272321)



Immunoprecipitation - Anti-L1CAM antibody
[EPR23241-224] - BSA and Azide free (ab272321)

Flow cytometric analysis of A549 (Human lung carcinoma epithelial cell, Left) / A375 (Human malignant melanoma epithelial cell, Right) cells labelling L1CAM with <u>ab270455</u> at 1/500 dilution (Red) compared with a Rabbit monoclonal IgG (<u>ab172730</u>) (Black) isotype control and an unlabelled control (cells without incubation with primary antibody and secondary antibody) (Blue). Goat anti rabbit IgG (Alexa Fluor[®] 488, <u>ab150077</u>) at 1/2000 dilution was used as the secondary antibody.

A549 is negative or very low expression cell line for L1CAM (PMID: 23511563).

Gated on viable cells.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab270455).

L1CAM was immunoprecipitated from 0.35 mg A375 (human malignant melanoma epithelial cell), whole cell lysate using ab270455 at 1/30 dilution (2ug in 0.35mg lysates). Western blot was performed on the immunoprecipitate using ab270455 at 1/500 dilution. VeriBlot for IP Detection Reagent (HRP)(ab270455 at 1/5000 dilution.

Lane 1: A375 (human malignant melanoma epithelial cell), whole cell lysate 10ug

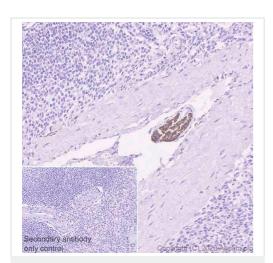
Lane 2: ab270455 IP in A375 whole cell lysate

Lane 3: Rabbit monoclonal IgG ($\underline{ab172730}$) instead of $\underline{ab270455}$ in A375 whole cell lysate

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

Exposure time: 5.5 seconds.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab270455).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-L1CAM antibody

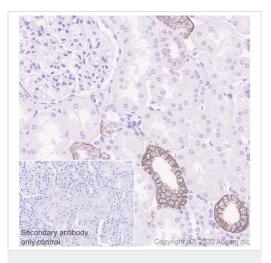
[EPR23241-224] - BSA and Azide free (ab272321)

Immunohistochemical analysis of paraffin-embedded Human spleen tissue labeling L1CAM with <u>ab270455</u> at 1/100 dilution followed by a ready to use Rabbit specific IHC polymer detection kit HRP/DAB (<u>ab209101</u>). Positive staining on the nerve in human spleen (PMID: 26743472). The section was incubated with <u>ab270455</u> for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Counterstained with Hematoxylin.

Secondary antibody only control: Secondary antibody is a ready to use Rabbit specific IHC polymer detection kit HRP/DAB (ab209101).

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution2) for 20 mins.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab270455</u>).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-L1CAM antibody

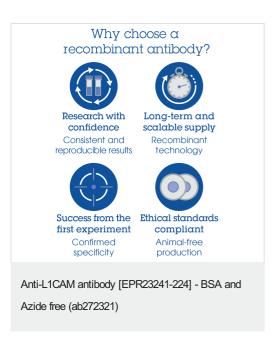
[EPR23241-224] - BSA and Azide free (ab272321)

Immunohistochemical analysis of paraffin-embedded Human kidney tissue labeling L1CAM with <u>ab270455</u> at 1/100 dilution followed by a ready to use Rabbit specific IHC polymer detection kit HRP/DAB (<u>ab209101</u>). Membranous staining on distal renal tubules in human kidney. The section was incubated with <u>ab270455</u> for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND[®] RX instrument. Counterstained with Hematoxylin.

Secondary antibody only control: Secondary antibody is a ready to use Rabbit specific IHC polymer detection kit HRP/DAB (ab209101).

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution2) for 20 mins.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab270455).



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