

**MFSD2A Rabbit pAb**  
**Rabbit Polyclonal Antibody**  
**Catalog # AP93287****Specification**

---

**MFSD2A Rabbit pAb - Product Information**

Application	<b>IHC, WB, FC</b>
Primary Accession	<a href="#">O8NA29</a>
Reactivity	<b>Human, Mouse, Rat</b>
Predicted	<b>Rabbit, Pig, Chicken</b>
Host	<b>Rabbit</b>
Clonality	<b>Polyclonal</b>
Calculated MW	<b>60 kDa KDa</b>

**MFSD2A Rabbit pAb - Additional Information****Gene ID** 84879**Other Names**

Major facilitator superfamily domain containing 2; Major facilitator superfamily domain containing 2A; Major facilitator superfamily domain-containing protein 2A; MFS2A\_HUMAN; MFSD2; MFSD2A.

**Target/Specificity**

0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol.

**Dilution**

&lt;span class = "dilution\_IHC"&gt;IHC~~1:100-500&lt;/span&gt;&lt;br \&gt;&lt;span class = "dilution\_WB"&gt;WB~~1:500-2000&lt;/span&gt;&lt;br \&gt;&lt;span class = "dilution\_FC"&gt;FC~~0.2ug/test&lt;/span&gt;

**Format**

0.01M TBS(pH7.4), 0.09% (W/V) sodium azide and 50% Glyce

**Storage**

Store at -20 °C for one year. Avoid repeated freeze/thaw cycles. When reconstituted in sterile pH 7.4 0.01M PBS or diluent of antibody the antibody is stable for at least two weeks at 2-4 °C.

**MFSD2A Rabbit pAb - Protein Information****Name** MFSD2A {ECO:0000303|PubMed:18694395, ECO:0000312|HGNC:HGNC:25897}**Function**

Sodium-dependent lysophosphatidylcholine (LPC) symporter, which plays an essential role for blood-brain barrier formation and function (PubMed:&lt;a href="http://www.uniprot.org/citations/24828040" target="\_blank"&gt;24828040&lt;/a&gt;, PubMed:&lt;a href="http://www.uniprot.org/citations/32572202" target="\_blank"&gt;32572202&lt;/a&gt;, PubMed:&lt;a href="http://www.uniprot.org/citations/34135507" target="\_blank"&gt;34135507&lt;/a&gt;). Specifically expressed in endothelium of the blood-brain barrier of micro-vessels and transports LPC into the

brain (By similarity). Transport of LPC is essential because it constitutes the major mechanism by which docosahexaenoic acid (DHA), an omega-3 fatty acid that is essential for normal brain growth and cognitive function, enters the brain (PubMed:<a href="http://www.uniprot.org/citations/26005868" target="\_blank">26005868</a>, PubMed:<a href="http://www.uniprot.org/citations/34135507" target="\_blank">34135507</a>). Transports LPC carrying long-chain fatty acids such LPC oleate and LPC palmitate with a minimum acyl chain length of 14 carbons (By similarity). Does not transport docosahexaenoic acid in unesterified fatty acid (By similarity). Specifically required for blood-brain barrier formation and function, probably by mediating lipid transport (By similarity). Not required for central nervous system vascular morphogenesis (By similarity). Acts as a transporter for tunicamycin, an inhibitor of asparagine-linked glycosylation (PubMed:<a href="http://www.uniprot.org/citations/21677192" target="\_blank">21677192</a>). In placenta, acts as a receptor for ERVFRD-1/syncytin-2 and is required for trophoblast fusion (PubMed:<a href="http://www.uniprot.org/citations/18988732" target="\_blank">18988732</a>, PubMed:<a href="http://www.uniprot.org/citations/23177091" target="\_blank">23177091</a>).

### Cellular Location

Cell membrane; Multi-pass membrane protein. Endoplasmic reticulum membrane {ECO:0000250|UniProtKB:Q9DA75}; Multi-pass membrane protein. Note=Cytoplasmic punctae that may represent vesicles shuttling between the endoplasmic reticulum and the plasma membrane (PubMed:21677192).

### Tissue Location

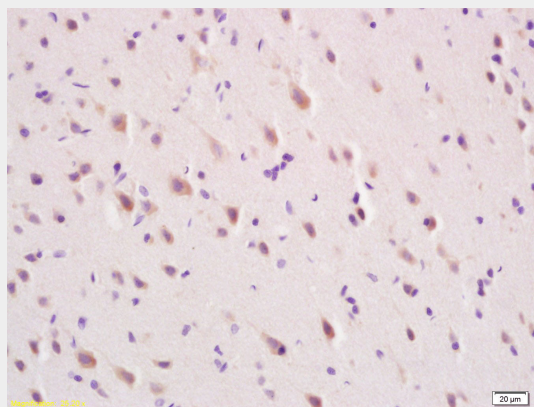
In placenta, associated with trophoblast cells.

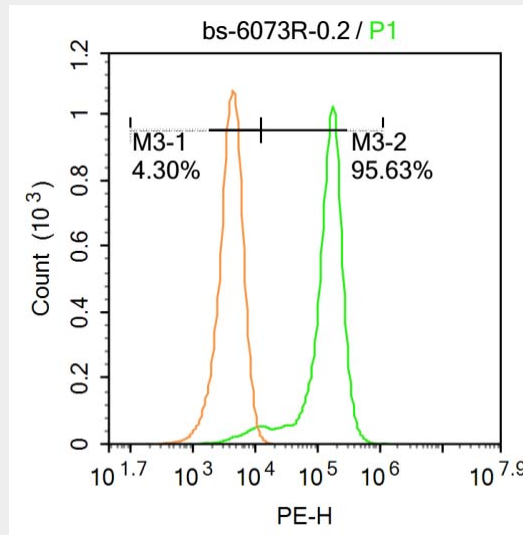
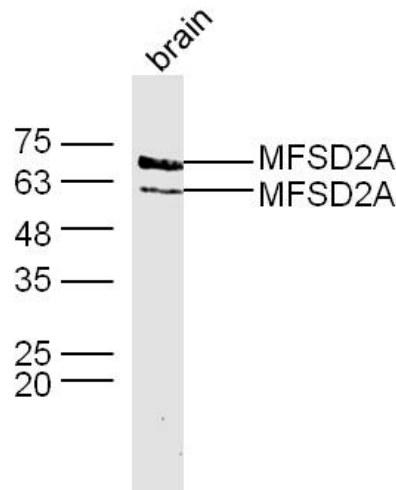
## MFSD2A Rabbit pAb - Protocols

Provided below are standard protocols that you may find useful for product applications.

- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

## MFSD2A Rabbit pAb - Images





### MFSD2A Rabbit pAb - Background

MFSD2 is a 543 amino acid multi-pass membrane protein of the endoplasmic reticulum that is involved in beta-adrenergic signaling during thermogenesis. Existing as three alternatively spliced isoforms, MFSD2 plays a role in G1 regulation and is encoded by a gene that maps to human chromosome 1p34.2. Human chromosome 1 spans 260 million base pairs, contains over 3,000 genes, comprises nearly 8% of the human genome and houses a large number of disease-associated genes, including those that are involved in familial adenomatous polyposis, Stickler syndrome, Parkinson's disease, Gaucher disease, schizophrenia and Usher syndrome.