

Anti-Human ARSA DyLight® 550 conjugated Antibody(monoclonal, 4C10)

Catalog # ABO14809

Specification

Anti-Human ARSA DyLight® 550 conjugated Antibody(monoclonal, 4C10) - Product Information

Application FC
Primary Accession P15289
Host Mouse

Isotype Mouse IgG2a
Reactivity Human
Clonality Monoclonal
Format Liquid

Description

Anti-Human ARSA DyLight® 550 conjugated Antibody (monoclonal, 4C10). Tested in Flow Cytometry applications. This antibody reacts with Human.

Anti-Human ARSA DyLight® 550 conjugated Antibody(monoclonal, 4C10) - Additional Information

Gene ID 410

Other Names

Arylsulfatase A, ASA, 3.1.6.8, Cerebroside-sulfatase, Arylsulfatase A component B, Arylsulfatase A component C, ARSA

Application Details

Flow Cytometry, 1-3 μg/1x10⁶ cells

Subcellular Localization

Lysosome.

Contents

Each vial contains 50% glycerol, 0.9% NaCl, 0.2% Na2HPO4, 0.02% NaN3.

Immunogen

A synthetic peptide corresponding to a sequence at the C-terminus of human ARSA, different from the related mouse sequence by six amino acids.

Cross Reactivity

No cross-reactivity with other proteins.

Storage At -20°C for one year from date of receipt.

Avoid repeated freezing and thawing.

Protect from light.

Anti-Human ARSA DyLight® 550 conjugated Antibody(monoclonal, 4C10) - Protein Information



Name ARSA

Function Hydrolyzes cerebroside sulfate.

Cellular Location Endoplasmic reticulum. Lysosome

Anti-Human ARSA DyLight® 550 conjugated Antibody(monoclonal, 4C10) - Protocols

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

- Western Blot
- Blocking Peptides
- Dot Blot
- Immunohistochemistry
- Immunofluorescence
- <u>Immunoprecipitation</u>
- Flow Cytomety
- Cell Culture

Anti-Human ARSA DyLight® 550 conjugated Antibody(monoclonal, 4C10) - Images

Anti-Human ARSA DyLight® 550 conjugated Antibody(monoclonal, 4C10) - Background

Arylsulfatase A (ARSA) is an enzyme that breaks down sulfatides, namely cerebroside 3-sulfate intocerebroside and sulfate. In humans, arylsulfatase A is encoded by the ARSA gene. ARSA is mapped to 22q13.33. The protein encoded by this gene hydrolyzes cerebroside sulfate to cerebroside and sulfate. Defects in this gene lead to metachromatic leucodystrophy (MLD), a progressive demyelination disease which results in a variety of neurological symptoms and ultimately death. Alternatively spliced transcript variants have been described for this gene.