

# ARSB Antibody (C-term)

Purified Rabbit Polyclonal Antibody (Pab) Catalog # AP7460b

#### **Product Information**

**Application** WB, IHC-P, E **Primary Accession** P15848 Reactivity Human Host Rabbit Clonality Polyclonal Isotype Rabbit IgG **Clone Names** RB18353 **Calculated MW** 59687 **Antigen Region** 464-493

## **Additional Information**

Gene ID 411

Other Names Arylsulfatase B, ASB, N-acetylgalactosamine-4-sulfatase, G4S, ARSB

**Target/Specificity** This ARSB antibody is generated from rabbits immunized with a KLH

conjugated synthetic peptide between 464-493 amino acids from the

C-terminal region of human ARSB.

**Dilution** WB~~1:1000 IHC-P~~1:100~500 E~~Use at an assay dependent concentration.

**Format** Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide.

This antibody is prepared by Saturated Ammonium Sulfate (SAS) precipitation

followed by dialysis against PBS.

**Storage** Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store

at -20°C in small aliquots to prevent freeze-thaw cycles.

**Precautions** ARSB Antibody (C-term) is for research use only and not for use in diagnostic

or therapeutic procedures.

## **Protein Information**

Name ARSB

**Function** Removes sulfate groups from chondroitin-4-sulfate (C4S) and regulates its

degradation (PubMed:<u>19306108</u>). Involved in the regulation of cell adhesion, cell migration and invasion in colonic epithelium (PubMed:<u>19306108</u>). In the central nervous system, is a regulator of neurite outgrowth and neuronal plasticity, acting through the control of sulfate glycosaminoglycans and

neurocan levels (By similarity).

**Cellular Location** 

Lysosome {ECO:0000250 | UniProtKB:P50429}. Cell surface {ECO:0000250 | UniProtKB:P50429}

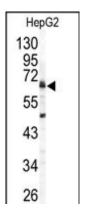
# **Background**

ARSB belongs to the sulfatase family. The arylsulfatase B homodimer hydrolyzes sulfate groups of N-Acetyl-D-galactosamine, chondriotin sulfate, and dermatan sulfate. The protein is targetted to the lysozyme. Mucopolysaccharidosis type VI is an autosomal recessive lysosomal storage disorder resulting from a deficiency of arylsulfatase B.

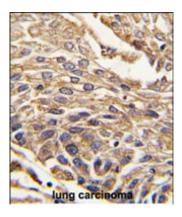
## References

Peters C., Schmidt B.J. Biol. Chem. 265:3374-3381(1990) Modaressi S., Rupp K.Biol. Chem. Hoppe-Seyler 374:327-335(1993) Kobayashi T., Honke K.Biochim. Biophys. Acta 1159:243-247(1992)

# **Images**



Western blot analysis of ARSB antibody (C-term) (Cat.#AP7460b) in HepG2 cell line lysates (35ug/lane). ARSB (arrow) was detected using the purified Pab.



Formalin-fixed and paraffin-embedded human lung carcinoma tissue reacted with ARSB antibody (C-term) (Cat. #AP7460b), which was peroxidase-conjugated to the secondary antibody, followed by DAB staining. This data demonstrates the use of this antibody for immunohistochemistry; clinical relevance has not been evaluated.

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.