

Product datasheet

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ARG59505 anti-SFTPC / SP-C antibody

Package: 100 μl Store at: -20°C

Summary

Product Description Rabbit Polyclonal antibody recognizes SFTPC / SP-C

Tested Reactivity Hu, Ms, Rat

Tested Application ICC/IF, IHC-P, WB

Host Rabbit

Clonality Polyclonal

Isotype IgG

Target Name SFTPC / SP-C

Species Human

Immunogen Recombinant fusion protein corresponding to aa. 1-197 of Human SFTPC / SP-C (NP_001165881.1).

Conjugation Un-conjugated

Alternate Names BRICD6; SP-C; Val; SFTP2; SP5; PSP-C; Pulmonary surfactant-associated protein C; Pulmonary surfactant-

associated proteolipid SPL; SMDP2

Application Instructions

redict Reactivit	y Note	Rat

Application table

Application	Dilution
ICC/IF	1:50 - 1:200
IHC-P	1:50 - 1:200
WB	1:500 - 1:2000

Application Note * The dilutions indicate recommended starting dilutions and the optimal dilutions or concentrations

should be determined by the scientist.

Positive Control Mouse pancreas and K562

Observed Size 21 kDa

Properties

Form Liquid

Purification Affinity purified.

Buffer PBS (pH 7.3), 0.02% Sodium azide and 50% Glycerol.

Preservative 0.02% Sodium azide

Stabilizer 50% Glycerol

Storage instruction For continuous use, store undiluted antibody at 2-8°C for up to a week. For long-term storage, aliquot

and store at -20°C. Storage in frost free freezers is not recommended. Avoid repeated freeze/thaw

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For laboratory research only, not for drug, diagnostic or other use.

Bioinformation

Gene Symbol

Gene Full Name surfactant protein C

SFTPC

Background This gene encodes the pulmonary-associated surfactant protein C (SPC), an extremely hydrophobic

surfactant protein essential for lung function and homeostasis after birth. Pulmonary surfactant is a surface-active lipoprotein complex composed of 90% lipids and 10% proteins which include plasma proteins and apolipoproteins SPA, SPB, SPC and SPD. The surfactant is secreted by the alveolar cells of the lung and maintains the stability of pulmonary tissue by reducing the surface tension of fluids that coat the lung. Multiple mutations in this gene have been identified, which cause pulmonary surfactant metabolism dysfunction type 2, also called pulmonary alveolar proteinosis due to surfactant protein C deficiency, and are associated with interstitial lung disease in older infants, children, and adults.

Alternatively spliced transcript variants encoding different protein isoforms have been

identified.[provided by RefSeq, Feb 2010]

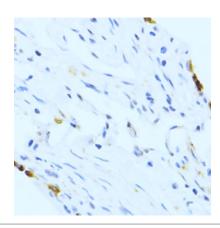
Function Pulmonary surfactant associated proteins promote alveolar stability by lowering the surface tension at

the air-liquid interface in the peripheral air spaces. [UniProt]

Calculated Mw 21 kDa

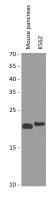
Cellular Localization Secreted, extracellular space, surface film. [UniProt]

Images



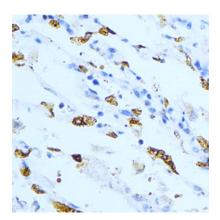
ARG59505 anti-SFTPC / SP-C antibody IHC-P image

Immunohistochemistry: Paraffin-embedded Human lung stained with ARG59505 anti-SFTPC / SP-C antibody at 1:100 dilution.



ARG59505 anti-SFTPC / SP-C antibody WB image

Western blot: 25 μg of Mouse pancreas and K562 cell lysate stained with ARG59505 anti-SFTPC / SP-C antibody at 1:1000 dilution.



ARG59505 anti-SFTPC / SP-C antibody IHC-P image

Immunohistochemistry: Paraffin-embedded Human lung cancer stained with ARG59505 anti-SFTPC / SP-C antibody at 1:100 dilution.