

Rabbit Anti-ASBT/SLC10A2 antibody

SL23146R

ASBT/SLC10A2
顶膜钠依赖性胆盐转运体蛋白抗体
SLC10A2; Apical sodium dependent bile acid transporter; Apical sodium- dependent bile acid transporter; IBAT; ileal apical sodium-dependent bile acid transporter; Ileal sodium dependent bile acid transporter; Ileal sodium-dependent bile acid transporter; Ileal sodium/bile acid cotransporter; ISBT; Na+ bile acid cotransporter; Na+ dependent ileal bile acid transporter; NTCP2; Sodium/taurocholate cotransporting polypeptide, ileal; solute carrier family 10 (sodium/bile acid cotransporter family); Solute carrier family 10 member 2.
Rabbit
Polyclonal
Human,Mouse,Rat,Dog,Cow,Horse,Rabbit,
WB=1:500-2000 not yet tested in other applications. optimal dilutions/concentrations should be determined by the end user.
38kDa
The cell membrane
Lyophilized or Liquid
1mg/ml
KLH conjugated synthetic peptide derived from human ASBT/SLC10A2:131- 230/348 <extracellular></extracellular>
IgG
affinity purified by Protein A
0.01M TBS(pH7.4) with 1% BSA, 0.03% Proclin300 and 50% Glycerol.
Store at -20 °C for one year. Avoid repeated freeze/thaw cycles. The lyophilized antibody is stable at room temperature for at least one month and for greater than a year when kept at -20°C. 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.
PubMed
SLC10A2 plays a critical role in reabsorption of bile acids from the the small intestine

lumen. Passive flow of sodium ions down their concentration gradient is coupled to bile acid movement, resulting in an increase in the concentration of bile acids in the interior of the cell. This action conserves the body's pool of re-circulating bile acid. SLC10A2 also plays a key role in cholesterol metabolism as cholesterol is the precursor molecule in bile acid synthesis mediated by CYP7A and FXR.

Function:

Plays a critical role in the sodium-dependent reabsorption of bile acids from the lumen of the small intestine. Plays a key role in cholesterol metabolism.

Subunit: Monomer and homodimer.

Subcellular Location: Membrane; Multi-pass membrane protein.

DISEASE:

Primary bile acid malabsorption (PBAM) [MIM:613291]: An intestinal disorder associated with chronic watery diarrhea, excess fecal bile acids, steatorrhea and interruption of the enterohepatic circulation of bile acids. {ECO:0000269|PubMed:9109432}. Note=The disease is caused by mutations affecting the gene represented in this entry.

Similarity:

Belongs to the bile acid:sodium symporter (BASS) (TC 2.A.28) family.

SWISS: 012908

Gene ID: 6555

Database links:

Entrez Gene: 6555Human

Entrez Gene: 20494Mouse

Entrez Gene: 29500Rat

Omim: 601295Human

SwissProt: Q12908Human

SwissProt: P70172Mouse

