

Basic Information

Product Name	Anti-SLC25A51/52 Antibody	
Gene Name	SLC25A51/SLC25A52	
Source	Rabbit	
Clonality	Polyclonal	
Isotype	IgG	
Species Reactivity	human, monkey, mouse	
Tested Application	WB, IHC, FCM, ICC/IF, ELISA	
Contents	500 ug/ml antibody with PBS, 0.02% NaN ₃ , 1 mg/ml BSA and 50% glycerol.	
Immunogen	E. coli-derived human SLC25A51/52 recombinant protein (Position: M1-E291). Human SLC25A51/52 shares 87% and 86.3% amino acid (aa) sequence identity with mouse and rat SLC25A51/52, respectively.	
Concentration	500 ug/ml	
Purification	Immunogen affinity purified.	
Observed MW	45 kDa	
Dilution Ratios	Western blot (WB):	1:500-2000
	Immunohistochemistry (IHC):	1:50-400
	Immunocytochemistry/Immunofluorescence (ICC/IF):	1:50-400
	Flow Cytometry (Fixed):	1:50-200
	Enzyme linked immunosorbent assay (ELISA):	1:100-1000
	(Boiling the paraffin sections in 10mM citrate buffer, pH6.0, or PH8.0 EDTA repair liquid for 20 mins is required for the staining of formalin/paraffin sections.) Optimal working dilutions must be determined by end user.	

Storage

12 months from date of receipt, -20°C as supplied. 6 months 2 to 8°C after reconstitution. Avoid repeated freezing and thawing.

Background Information

SLC25A51 gene ontology annotations related to this gene include transmembrane transporter activity.

Selected Validation Data

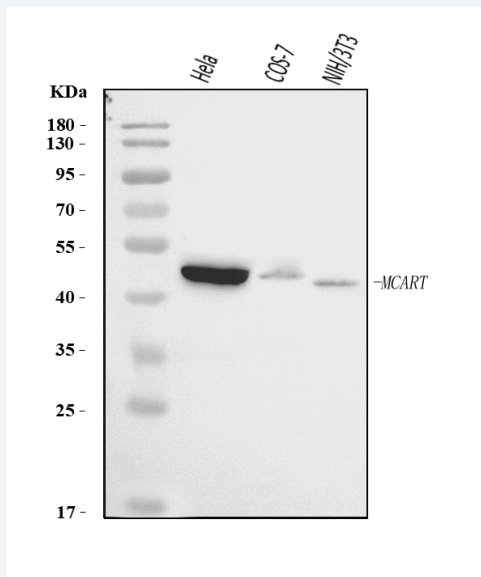


Figure 1. Western blot analysis of SLC25A51/52 using anti-SLC25A51/52 antibody (A17353). The sample well of each lane was loaded with 30 ug of sample under reducing conditions.

Lane 1: HeLa whole cell lysates,

Lane 2: COS-7 whole cell lysates,

Lane 3: NIH/3T3 whole cell lysates.

After electrophoresis, proteins were transferred to a membrane. Then the membrane was incubated with rabbit anti-SLC25A51/52 antigen affinity purified polyclonal antibody (A17353) at a dilution of 1:1000 and probed with a goat anti-rabbit IgG-HRP secondary antibody (Catalog # BA1054). The signal is developed using ECL Plus Western Blotting Substrate (Catalog # AR1197). A specific band was detected for SLC25A51/52 at approximately 45 kDa. The expected band size for SLC25A51/52 is at 34 kDa.

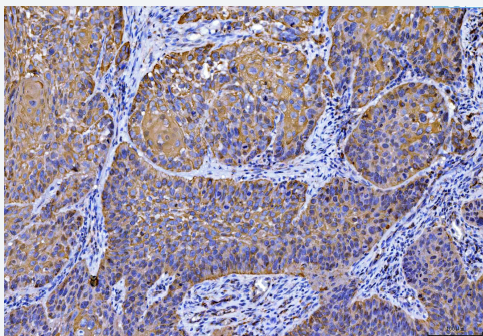


Figure 2. IHC analysis of SLC25A51/52 using anti-SLC25A51/52 antibody (A17353).

SLC25A51/52 was detected in a paraffin-embedded section of human Laryngeal squamous cell carcinomas tissue. The tissue section was incubated with rabbit anti-SLC25A51/52 Antibody (A17353) at a dilution of 1:200 and developed using HRP Conjugated Rabbit IgG Super Vision Assay Kit (Catalog # SV0002) with DAB (Catalog # AR1022) as the chromogen.

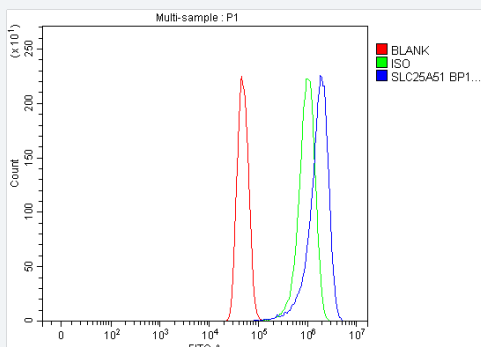


Figure 4. Flow Cytometry analysis of U251 cells using anti-SLC25A51/52 antibody (A17353).

Overlay histogram showing U251 cells stained with A17353 (Blue line). To facilitate intracellular staining, cells were fixed with 4% paraformaldehyde and permeabilized with permeabilization buffer. The cells were blocked with 10% normal goat serum. And then incubated with rabbit anti-SLC25A51/52 Antibody (A17353) at 1:100 dilution for 30 min at 20°C. DyLight®488 conjugated goat anti-rabbit IgG (BA1127) was used as secondary antibody at 1:100 dilution for 30 minutes at 20°C. Isotype control antibody (Green line) was rabbit IgG at 1:100 dilution used under the same conditions. Unlabelled sample without incubation with primary antibody

and secondary antibody (Red line) was used as a blank control.

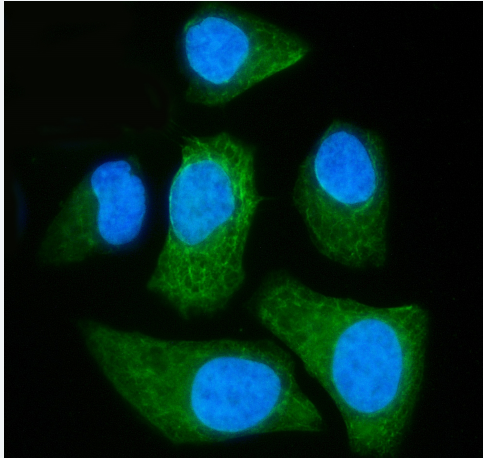


Figure 5. IF analysis of SLC25A51/52 using anti-SLC25A51/52 antibody (A17353).

SLC25A51/52 was detected in an immunocytochemical section of HeLa cells. The section was incubated with rabbit anti-SLC25A51/52 Antibody (A17353) at a dilution of 1:100. DyLight® 488 Conjugated Goat Anti-Rabbit IgG (Green) (Catalog # BA1127) was used as secondary antibody. The section was counterstained with DAPI (Catalog # AR1176) (Blue).