

## Basic Information

<b>Product Name</b>	Anti-SQRDL/SQOR Antibody	
<b>Gene Name</b>	SQOR	
<b>Source</b>	Rabbit	
<b>Clonality</b>	Polyclonal	
<b>Isotype</b>	IgG	
<b>Species Reactivity</b>	human	
<b>Tested Application</b>	WB, IHC, ICC/IF, FCM, ELISA	
<b>Contents</b>	500 ug/ml antibody with PBS, 0.02% NaN <sub>3</sub> , 1 mg/ml BSA and 50% glycerol.	
<b>Immunogen</b>	E.coli-derived human SQRDL/SQOR recombinant protein (Position: R12-E321). Human SQOR shares 86.1% amino acid (aa) sequence identity with mouse SQOR.	
<b>Concentration</b>	500 ug/ml	
<b>Purification</b>	Immunogen affinity purified.	
<b>Observed MW</b>	50 kDa	
<b>Dilution Ratios</b>	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.

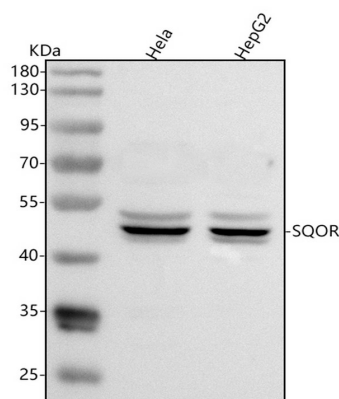
## Background Information

The protein encoded by this gene may function in mitochondria to catalyze the conversion of sulfide to persulfides, thereby decreasing toxic concentrations of sulfide. Alternative splicing results in multiple transcript variants that encode the same protein.

## Reference

Anti-SQRDL/SQOR Antibody被引用在1文献中。

## Selected Validation Data

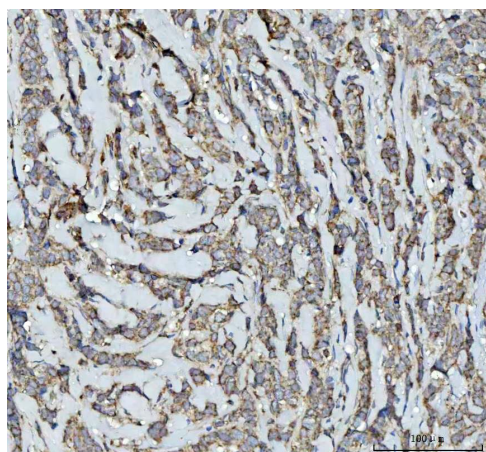


Western blot analysis of SQRDL/SQOR using anti-SQRDL/SQOR antibody (A31817). The sample well of each lane was loaded with 30 ug of sample under reducing conditions.

Lane 1: human HeLa whole cell lysates,

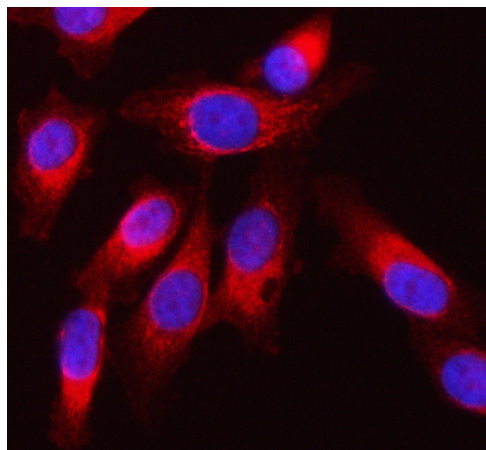
Lane 2: human HepG2 whole cell lysates.

After electrophoresis, proteins were transferred to a membrane. Then the membrane was incubated with rabbit anti-SQRDL/SQOR antigen affinity purified polyclonal antibody (A31817) 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 SQRDL/SQOR at approximately 50 kDa. The expected band size for SQRDL/SQOR is at 50 kDa.



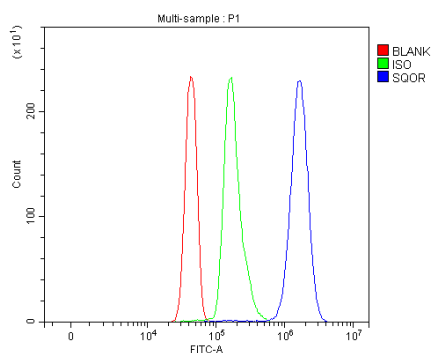
IHC analysis of SQRDL/SQOR using anti-SQRDL/SQOR antibody (A31817) .

SQRDL/SQOR was detected in a paraffin-embedded section of human breast cancer tissue. The tissue section was incubated with rabbit anti-SQRDL/SQOR Antibody (A31817) at a dilution of 1:200 and developed using HRP Conjugated Rabbit IgG Super Vision Assay Kit (Catalog # SV0002) with DAB (Catalog # AR1027) as the chromogen.



ICC/IF analysis of SQRDL/SQOR using anti-SQRDL/SQOR antibody (A31817).

SQRDL/SQOR was detected in an immunocytochemical section of PC-3 cells. The section was incubated with rabbit anti-SQRDL/SQOR Antibody (A31817) at a dilution of 1:100. Fluoro550-conjugated Anti-rabbit IgG Secondary Antibody (red)(Catalog#BA1135) was used as secondary antibody. The section was counterstained with DAPI (Catalog # AR1176) (Blue).



Flow Cytometry analysis of Jurkat cells using anti-SQRDL/SQOR antibody (A31817).

Overlay histogram showing Jurkat cells stained with A31817 (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-SQRDL/SQOR Antibody (A31817) at 1:100 dilution for 30 min at 20°C. Fluoro488 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 (Red line) was also used as a control.