

SCYL1 polyclonal antibody

Catalog: BS8395

Host: Rabbit

Reactivity: Human, Mouse, Rat

BackGround:

With approximately 135 million base pairs and 1,400 genes, chromosome 11 makes up around 4% of human genomic DNA and is considered a gene and disease association dense chromosome. The chromosome 11 encoded Atm gene is important for regulation of cell cycle arrest and apoptosis following double strand DNA breaks. Atm mutation leads to the disorder known as ataxia-telangiectasia. The blood disorders Sickle cell anemia and β thalassemia are caused by HBB gene mutations. Wilms' tumors, WAGR syndrome and Denys-Drash syndrome are associated with mutations of the WT1 gene. Jervell and Lange-Nielsen syndrome, Jacobsen syndrome, Niemann-Pick disease, hereditary angioedema and Smith-Lemli-Opitz syndrome are also associated with defects in chromosome 11.

Product:

Rabbit IgG, 1mg/ml in PBS with 0.02% sodium azide, 50% glycerol, pH7.2

Molecular Weight:

~ 110 kDa

Swiss-Prot:

Q96KG9

Purification&Purity:

The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen and the purity is > 95% (by SDS-PAGE).

Applications:

WB: 1:500~1:2000

IHC/IF: 1:50~1:200

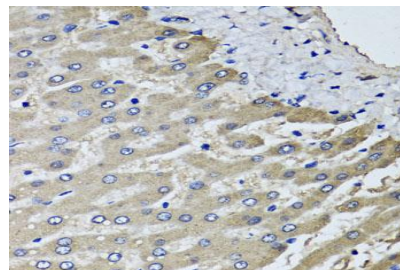
Storage&Stability:

Store at 4 °C short term. Aliquot and store at -20 °C long term. Avoid freeze-thaw cycles.

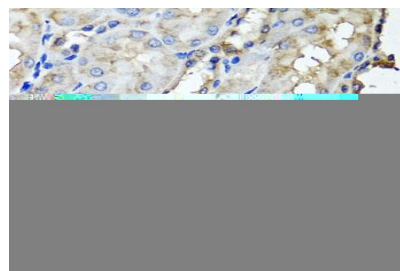
Specificity:

SCYL1 polyclonal antibody detects endogenous levels of SCYL1 protein.

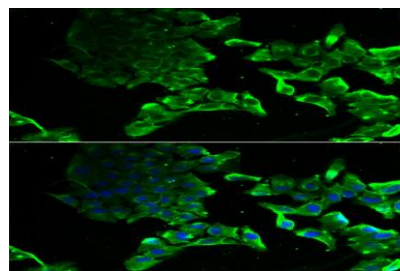
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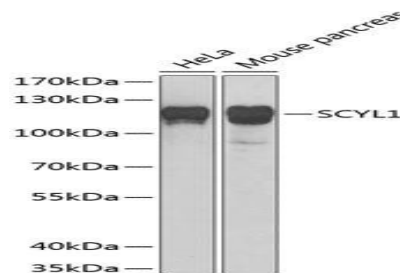
Immunohistochemistry of paraffin-embedded human liver injury using SCYL1 antibody at dilution of 1:100 (x40 lens).



Immunohistochemistry of paraffin-embedded rat kidney using SCYL1 antibody at dilution of 1:100 (x40 lens).



Immunofluorescence analysis of MCF-7 cell using SCYL1 antibody. Blue: DAPI for nuclear staining.



Western blot analysis of extracts of various cell lines, using SCYL1 An-

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PRODUCT DATA SHEET

Bioworld Technology, Inc.

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For research use only, not for use in diagnostic procedure.

Note:

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