

## Rabbit Anti-NOL4 antibody

SL19310R

Product Name:	NOL4
Chinese Name:	核仁蛋白4抗体
Alias:	HRIHFB2255; NOL 4; NOL4; NOL4 HUMAN; NOLP; Nucleolar localized protein;
	Nucleolar protein 4; Nucleolar-localized protein.
Organism Species:	Rabbit
Clonality:	Polyclonal
React Species:	Human,Mouse,Rat,Dog,Cow,Rabbit,Sheep,
Applications:	ELISA=1:500-1000IHC-P=1:400-800IHC-F=1:400-800ICC=1:100-500IF=1:100-
	500 (Paraffin sections need antigen repair)
	not yet tested in other applications.
	optimal dilutions/concentrations should be determined by the end user.
Molecular weight:	72kDa
Cellular localization:	The nucleus 🥏
Form:	Lyophilized or Liquid
<b>Concentration:</b>	1mg/ml
immunogen:	KLH conjugated synthetic peptide derived from human NOL4:1-100/638
Lsotype:	IgG
Purification:	affinity purified by Protein A
Storage Buffer:	0.01M TBS(pH7.4) with 1% BSA, 0.03% Proclin300 and 50% Glycerol.
Storage:	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:	PubMed
Product Detail:	NOL4 is a 524 amino acid protein that is predominantly expressed in testis and brain.
	NOL4 contains at least two domains that direct it to subnuclear locations. The gene
	encoding NOL4 is located on human chromosome 18, which houses over 300 protein-
	coding genes and contains nearly 76 million bases. There are a variety of diseases
	associated with defects in chromosome 18-localized genes, some of which include
	Trisomy 18 (also known as Edwards syndrome), Niemann-Pick disease, hereditary

hemorrhagic telangiectasia, erythropoietic protoporphyria and follicular lymphomas. There are two isoforms of NOL4 that are produced as a result of alternative splicing events.
Subcellular Location: Nucleus > nucleolus.
<b>Tissue Specificity:</b> Expressed predominantly in fetal brain, adult brain and testis.
SWISS: O94818
Gene ID: 8715 Database links: Entrez Gene: 8715 Human Omim: 603577 Human SwissProt: O94818 Human
Database links:
Entrez Gene: 8715 Human
<u>Omim: 603577</u> Human
Unigene: 514795 Human
Important Note:
This product as supplied is intended for research use only, not for use in human,
therapeutic or diagnostic applications.
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