

Product datasheet for AP53382PU-N

POLR1B (N-term) Rabbit Polyclonal Antibody

Product data:

OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

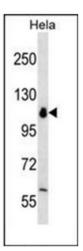
Product Type:	Primary Antibodies
Applications:	FC, IHC, WB
Recommended Dilution:	ELISA: 1/1000. Western Blot: 1/100-1/500. Flow Cytometry: 1/10-1/50. Immunohistochemistry on Paraffin Sections: 1/10-1/50.
Reactivity:	Human
Host:	Rabbit
lsotype:	lg
Clonality:	Polyclonal
Immunogen:	conjugated synthetic peptide between 88-117 amino acids from the N-terminal region of human POLR1B
Specificity:	This antibody recognizes Human POLR1B (N-term).
Formulation:	PBS containing 0.09% (W/V) Sodium Azide as preservative State: Aff - Purified State: Liquid purified Ig fraction
Concentration:	lot specific
Purification:	Affinitty Chromatography on Protein A
Conjugation:	Unconjugated
Storage:	Store undiluted at 2-8°C for one month or (in aliquots) at -20°C for longer. Avoid repeated freezing and thawing.
Stability:	Shelf life: one year from despatch.
Gene Name:	polymerase (RNA) l subunit B
Database Link:	<u>Entrez Gene 84172 Human</u> <u>Q9H9Y6</u>



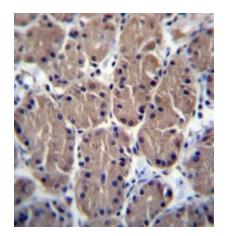
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	POLR1B (N-term) Rabbit Polyclonal Antibody – AP53382PU-N
Background:	Eukaryotic RNA polymerase I (pol I) is responsible for the transcription of ribosomal RNA (rRNA) genes and production of rRNA, the primary component of ribosomes. Pol I is a multisubunit enzyme composed of 6 to 14 polypeptides, depending on the species. Most of the mass of the pol I complex derives from the 2 largest subunits, Rpa1 and Rpa2 in yeast. POLR1B is homologous to Rpa2 (Seither and Grummt, 1996 [PubMed 8921381]).
Synonyms:	RNA polymerase I subunit 2, RPA135
Note:	Molecular Weight: 128229 Da
Protein Families	: Transcription Factors
Protein Pathwa	ys: Metabolic pathways, Purine metabolism, Pyrimidine metabolism, RNA polymerase

Product images:

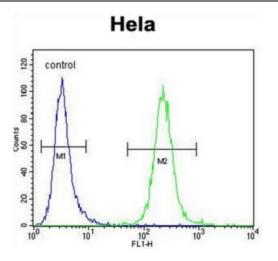


Western blot analysis of POLR1B Antibody (Nterm) in Hela cell line lysates (35ug/lane). This demonstrates the POLR1B antibody detected the POLR1B protein (arrow).



Formalin fixed and paraffin embedded human stomach tissue reacted with POLR1B Antibody (N-term) followed by peroxidase conjugation of the secondary antibody and DAB staining.

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Flow cytometric analysis of Hela cells using POLR1B Antibody (N-term) (right histogram) compared to a negative control cell (left histogram). FITC-conjugated goat-anti-rabbit secondary antibodies were used for the analysis.

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