



RECQ4 Polyclonal Antibody

Catalog No	YP-Ab-05361
Isotype	IgG
Reactivity	Human;Rat;Mouse;
Applications	WB;ELISA
Gene Name	RECQL4 RECQ4
Immunogen	Synthesized peptide derived from human protein . at AA range: 1030-1110
Specificity	RECQ4 Polyclonal Antibody detects endogenous levels of protein.
Formulation	Liquid in PBS containing 50% glycerol, and 0.02% sodium azide.
Source	Polyclonal, Rabbit,IgG
Purification	The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.
Dilution	WB 1:500-2000 ELISA 1:5000-20000
Concentration	1 mg/ml
Purity	≥90%
Storage Stability	-20°C/1 year
Synonyms	ATP dependent DNA helicase Q4;ATP-dependent DNA helicase Q4;DNA 3'-5' helicase RecQ4;DNA helicase;RecQ-like type 4;EC:5.6.2.4;RECQ4;RECQL4 RECQ4;ATP-dependent DNA helicase Q4 (EC 3.6.4.12) (DNA helicase;RecQ-like type 4) (RecQ4) (RTS) (RecQ protein-like 4)
Observed Band	145-150 kDa
Calculated Molecular Weight	1208 aa, 133 kDa
Cell Pathway	Cytoplasm . Nucleus .
Tissue Specificity	Ubiquitously expressed, with highest levels in thymus and testis.
Function	disease:Defects in RECQL4 are a cause of Baller-Gerold syndrome (BGS) [MIM:218600]; also known as craniosynostosis with radial defects. BGS is an autosomal recessive syndrome characterized by short stature, craniosynostosis, absent or hypoplastic radii, short and curved ulna, fused carpal bones and absent carpals, metacarpals and phalanges. Some patients manifest poikiloderma. Cases reported as Baller-Gerold syndrome have phenotypic overlap with several other disorders, including Saethre-Chotzen syndrome. BGS is part of the clinical spectrum of Rothmund-Thomson and RAPADILINO syndromes.,disease:Defects in RECQL4 are a cause of RAPADILINO syndrome [MIM:266280]. A disease characterized by radial and patellar aplasia or



hypoplasia.,disease:Defects in RECQL4 are a cause of Rothmund-Thomson syndrome (RTS) [MIM:268400]. A disease characterized by dermatological features such as atrophy, pigmen

Background

The protein encoded by this gene is a DNA helicase that belongs to the RecQ helicase family. DNA helicases unwind double-stranded DNA into single-stranded DNAs and may modulate chromosome segregation. This gene is predominantly expressed in thymus and testis. Mutations in this gene are associated with Rothmund-Thomson, RAPADILINO and Baller-Gerold syndromes. [provided by RefSeq, Jan 2010],

matters needing attention

Avoid repeated freezing and thawing!

Usage suggestions

This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.

Products Images