

Product datasheet for TA388155M

SUPV3L1 Rabbit Polyclonal Antibody

Product data:

Product Type: Primary Antibodies

IHC, WB **Applications:**

Recommended Dilution: Recommended dilution: WB:1:1000-1:5000, IHC:1:200-1:500

Reactivity: Rabbit Host: Isotype: lgG

Clonality: Polyclonal

Immunogen: Recombinant Human ATP-dependent RNA helicase SUPV3L1, mitochondrial protein (499-

786AA)

Formulation: Preservative: 0.03% Proclin 300

Constituents: 50% Glycerol, 0.01M PBS, pH 7.4

Concentration: lot specific

Purification: >95%, Protein G purified

Conjugation: Unconjugated

Upon receipt, store at -20°C or -80°C. Avoid repeated freeze. Storage:

1 year from dispatch. Stability:

Database Link: Q8IYB8

Background: Major helicase player in mitochondrial RNA metabolism. Component of the mitochondrial

> degradosome (mtEXO) complex, that degrades 3' overhang double-stranded RNA with a 3'-to-5' directionality in an ATP-dependent manner. ATPase and ATP-dependent multisubstrate helicase, able to unwind double-stranded (ds) DNA and RNA, and RNA/DNA heteroduplexes in the 5'-to-3' direction. Plays a role in the RNA surveillance system in mitochondria; regulates the stability of mature mRNAs, the removal of aberrantly formed mRNAs and the rapid degradation of non coding processing intermediates. Also implicated in recombination and

chromatin maintenance pathways. May protect cells from apoptosis. Associates with

mitochondrial DNA.



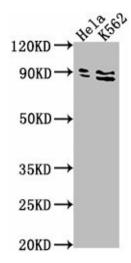
OriGene Technologies, Inc. 9620 Medical Center Drive, Ste 200

CN: techsupport@origene.cn

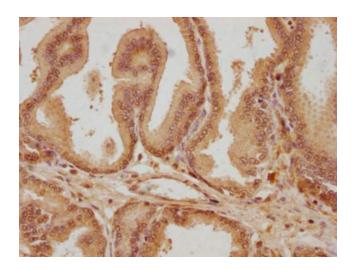
Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com



Product images:



Western Blot
Positive WB detected in: Hela whole cell lysate,
K562 whole cell lysate
All lanes: SUPV3L1 antibody at 1:2000
Secondary
Goat polyclonal to rabbit IgG at 1/50000 dilution
Predicted band size: 88 kDa
Observed band size: 88 kDa



IHC image of [TA388155] diluted at 1:300 and staining in paraffin-embedded human prostate tissue performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.