

# **Product datasheet for AP14367PU-N**

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# Integrin Linked Kinase (ILK) Rabbit Polyclonal Antibody

**Product data:** 

**Product Type:** Primary Antibodies

**Applications:** IHC, WB

Recommended Dilution: ELISA: 1/1,000.

Western blotting: 1/50 - 1/100. Immunohistochemistry: 1/10 - 1/50.

Reactivity: Human
Host: Rabbit
Isotype: Ig

Clonality: Polyclonal

Immunogen: KLH conjugated synthetic peptide corresponding to amino acid residues surrounding T173 of

human ILK1.

**Specificity:** This antibody reacts to ILK.

**Formulation:** PBS with 0.09% (W/V) sodium azide

State: Aff - Purified State: Liquid purified Ig

**Concentration:** lot specific

**Purification:** Affinity chromatography on Protein A

Conjugation: Unconjugated

Storage: Store the antibody 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:** integrin linked kinase

Database Link: Entrez Gene 3611 Human

Q13418



## Integrin Linked Kinase (ILK) Rabbit Polyclonal Antibody - AP14367PU-N

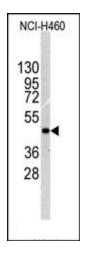
#### Background:

Transduction of extracellular matrix signals through integrins influences intracellular and extracellular functions, and appears to require interaction of integrin cytoplasmic domains with cellular proteins. Integrin-linked kinase (ILK), interacts with the cytoplasmic domain of beta-1 integrin. ILK is a serine/threonine protein kinase with 4 ankyrin-like repeats, which associates with the cytoplasmic domain of beta integrins and acts as a proximal receptor kinase regulating integrin-mediated signal transduction.

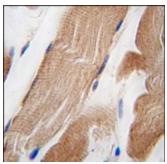
Synonyms:

ILK-1, ILK-2, p59ILK, ILK, ILK1, ILK2

# **Product images:**



Western blot analysis of anti-ILK Antibody (T173) in NCI-H460 cell line lysates (35ug/lane). EN1 (arrow) was detected using the purified Pab (1:60 dilution).



Formalin-fixed and paraffin-embedded human skeletal muscle tissue reacted with Phospho ILK Antibody (T173) antibody, which was peroxidase-conjugated to the secondary antibody, followed by DAB staining.