

UK Office

Everest Biotech Ltd

Cherwell Innovation Centre
77 Heyford Park
Upper Heyford
Oxfordshire
OX25 5HD
UK

Enquiries:

info@everestbiotech.com

Sales:

sales@everestbiotech.com

Tech support:

support@everestbiotech.com

Tel: +44 (0)1869 238326

www.everestbiotech.com

**Research Use Only. Not for
diagnostic or therapeutic use.**

EB06619-T - Goat Anti-IRAK4 (N Terminus) Antibody - Trial

Size: 20µg specific antibody in 40µl



Target Protein

Principal Names: IRAK4, REN64, NY-REN-64, interleukin-1 receptor-associated kinase 4, IRAK-4 mutated form 1, interleukin-1 receptor associated kinase 4, interleukin-1 receptor associated kinase 4 mutant form 1, IPD1

Official Symbol: IRAK4

Accession Number(s): NP_057207.2; NP_001107654.1

Human GeneID(s): [51135](#)

Non-Human GeneID(s): 266632 (mouse)

Immunogen

Peptide with sequence NKPITPSTYVRC, from the N Terminus of the protein sequence according to NP_057207.2; NP_001107654.1.

Please note the [peptide](#) is available for sale.

Purification and Storage

Purified from goat serum by ammonium sulphate precipitation followed by antigen affinity chromatography using the immunizing peptide.

Supplied at 0.5 mg/ml in Tris saline, 0.02% sodium azide, pH7.3 with 0.5% bovine serum albumin.

Aliquot and store at -20°C. Minimize freezing and thawing.

Applications Tested

Peptide ELISA: antibody detection limit dilution 1:32000.

Western blot: Approx 50kDa band observed in lysates of cell line HeLa and approx 55kDa in lysates of cell line Jurkat (calculated MW of 51.5kDa according to NP_057207.1). This molecular weight is routinely observed by other sources.

Recommended concentration: 0.5-2µg/ml. Primary incubation 1 hour at room temperature.

IHC: Paraffin embedded Human Heart and Human Liver. Recommended concentration: 3.75µg/ml.

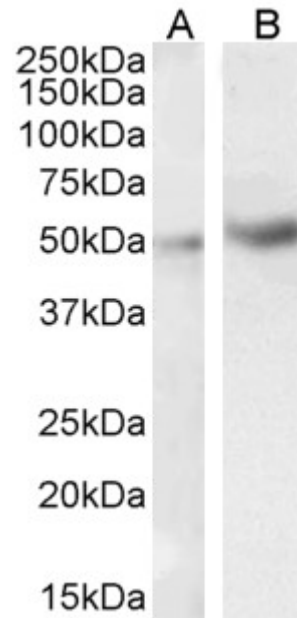
Immunofluorescence: Strong expression of the protein seen in the cytoplasm and microtubules of HeLa cells. Recommended concentration: 10µg/ml.

Flow Cytometry: Flow cytometric analysis of HeLa cells. Recommended concentration: 10ug/ml.

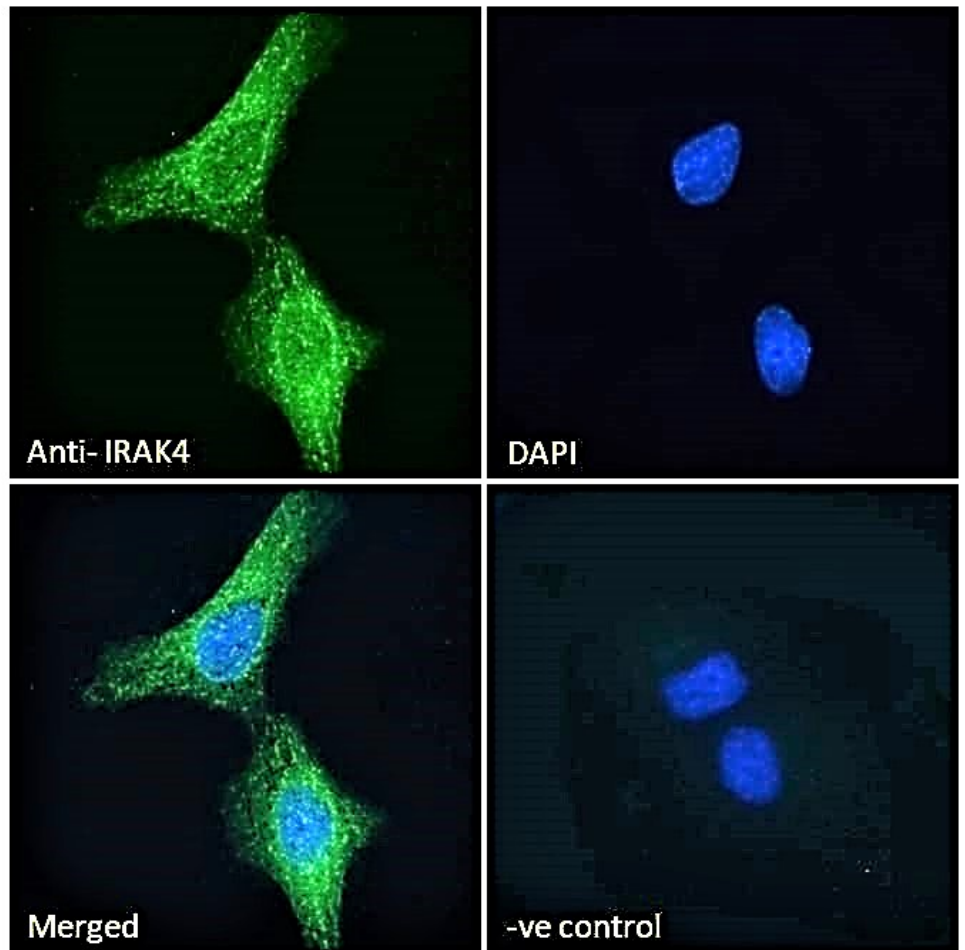
Species Reactivity

Tested: Human

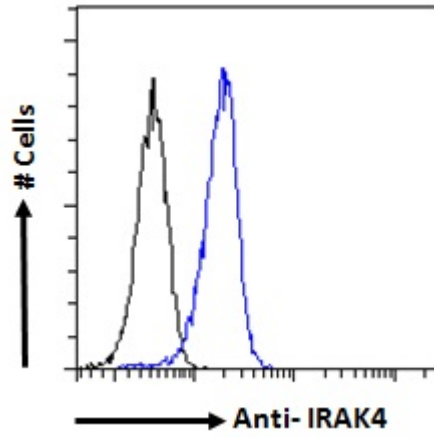
Expected from sequence similarity: Human, Mouse



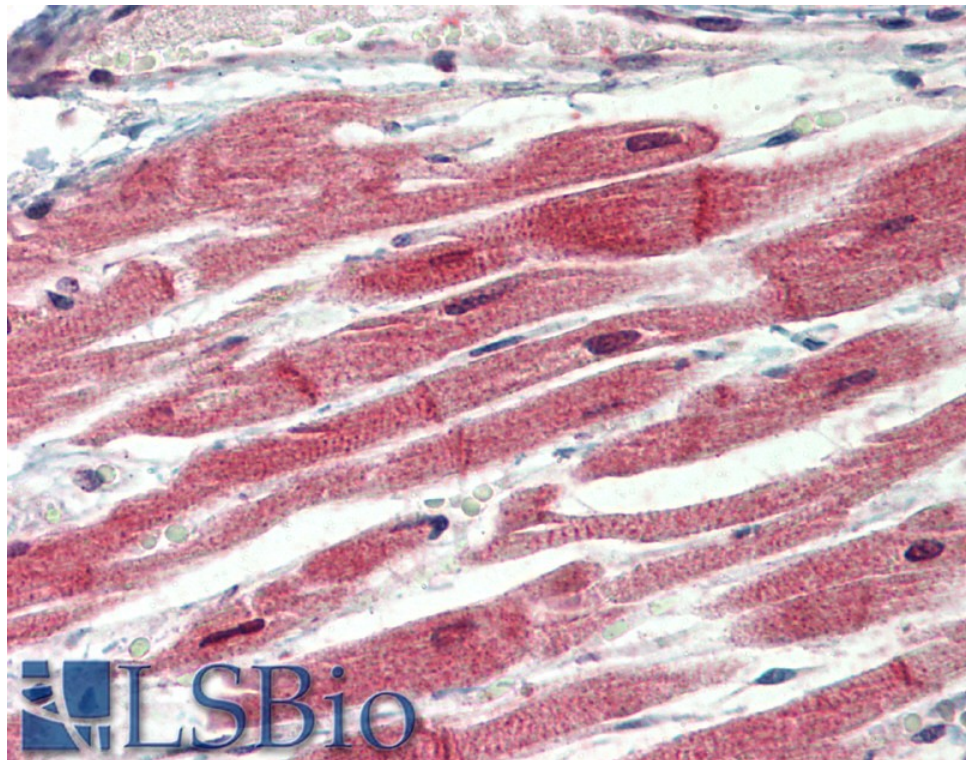
EB06619 (1 μ g/ml) staining of HeLa (A) and Jurkat (B) cell lysate (35 μ g protein in RIPA buffer). Detected by chemiluminescence.



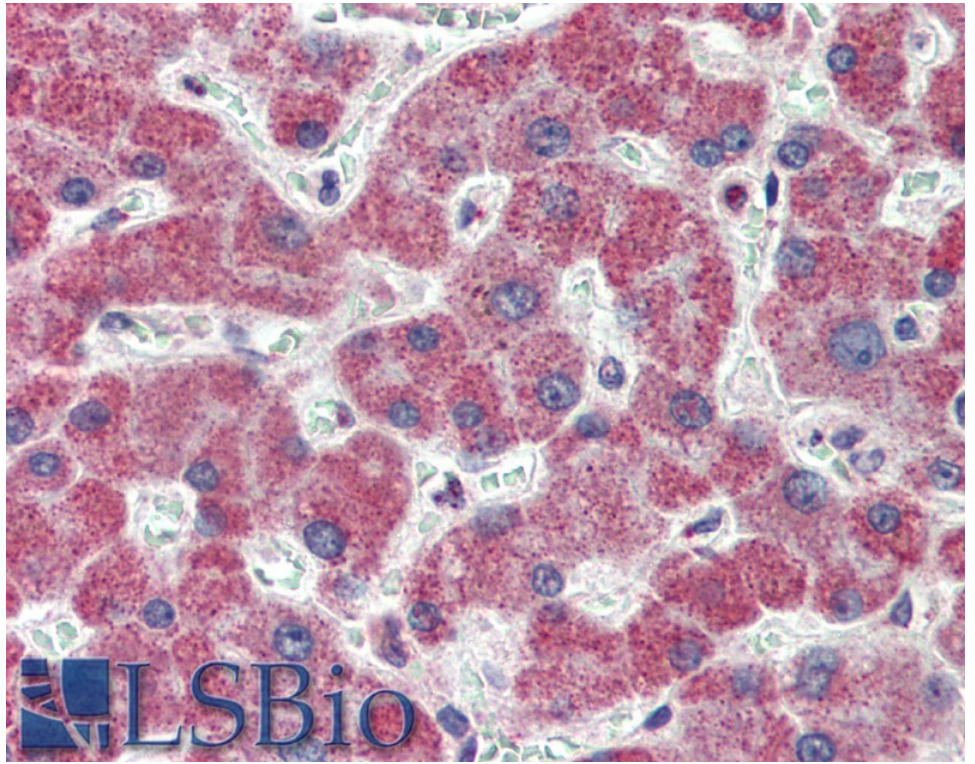
EB06619 Immunofluorescence analysis of paraformaldehyde fixed HeLa cells, permeabilized with 0.15% Triton. Primary incubation 1hr (10 μ g/ml) followed by Alexa Fluor 488 secondary antibody (2 μ g/ml), showing cytoplasmic and microtubule staining. The nuclear stain is DAPI (blue). Negative control: Unimmunized goat IgG (10 μ g/ml) followed by Alexa Fluor 488 secondary antibody (2 μ g/ml).



EB06619 Flow cytometric analysis of paraformaldehyde fixed HeLa cells (blue line), permeabilized with 0.5% Triton. Primary incubation 1hr (10ug/ml) followed by Alexa Fluor 488 secondary antibody (1ug/ml). IgG control: Unimmunized goat IgG (black line) followed by Alexa Fluor 488 secondary antibody.



EB06619 (3.75 μ g/ml) staining of paraffin embedded Human Heart. Steamed antigen retrieval with citrate buffer pH 6, AP-staining.



EB06619 (3.75µg/ml) staining of paraffin embedded Human Liver. Steamed antigen retrieval with citrate buffer pH 6, AP-staining.