



## UK Office

### Everest Biotech Ltd

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

Enquiries:

[info@everestbiotech.com](mailto:info@everestbiotech.com)

Sales:

[sales@everestbiotech.com](mailto:sales@everestbiotech.com)

Tech support:

[support@everestbiotech.com](mailto:support@everestbiotech.com)

Tel: +44 (0)1869 238326

[www.everestbiotech.com](http://www.everestbiotech.com)

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

## EB08772 - Goat Anti-APOLD1 / VERGE Antibody

Size: 100µg specific antibody in 200µl



### Target Protein

**Principal Names:** APOLD1, apolipoprotein L domain containing 1, DKFZP434F0318, FLJ25138, VERGE

**Official Symbol:** APOLD1

**Accession Number(s):** NP\_110444.3; NP\_001123887.1

**Human GeneID(s):** [81575](#)

### Immunogen

Peptide with sequence C-KSSRGHDLKISADQ, from the C Terminus of the protein sequence according to NP\_110444.3; NP\_001123887.1.

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 1:2000.

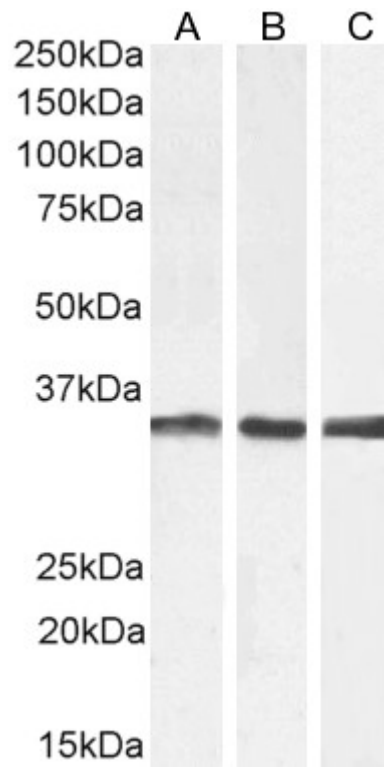
**Western blot:** Approx. 35kDa band observed in lysates of cell lines A431, Jurkat and U251 (calculated MW of 30.5kDa according to Human NP\_001123887.1). This molecular weight is routinely observed by other sources. Recommended concentration: 0.5-2µg/ml. Primary incubation 1 hour at room temperature.

**Flow Cytometry:** Flow cytometric analysis of A431 cells. Recommended concentration: 10µg/ml.

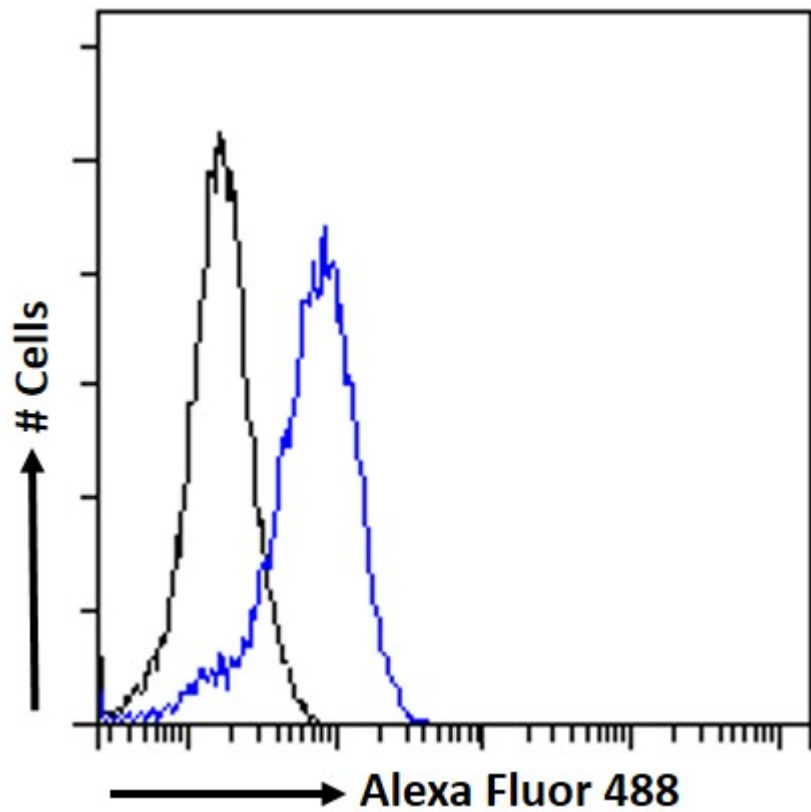
### Species Reactivity

**Tested:** Human

**Expected from sequence similarity:** Human



EB09741 (2 $\mu$ g/ml) staining of A431 (A), Jurkat (B), and U251 (C) cell lysate (35 $\mu$ g protein in RIPA buffer).  
Detected by chemiluminescence.



EB08772 Flow cytometric analysis of paraformaldehyde fixed A431 cells (blue line), permeabilized with 0.5% Triton. Primary incubation 1hr (10 $\mu$ g/ml) followed by Alexa Fluor 488 secondary antibody (1 $\mu$ g/ml). IgG control: Unimmunized goat IgG (black line) followed by Alexa Fluor 488 secondary antibody.