

## **Product Datasheet**

# Anti-Exonuclease 1/EXO1 Antibody (orb1290014)

**Description** Anti-Exonuclease 1/EXO1 Antibody. Tested in ELISA, IF, ICC, WB applications.

This antibody reacts with Human.

Species/Host Rabbit

**Reactivity** Human

**Conjugation** Unconjugated

**Tested Applications** ELISA, FC, ICC, IF, IP, WB

Immunogen E.coli-derived human Exonuclease 1/EXO1 recombinant protein (Position: Q25-

Q846).

Form/Appearance Lyophilized

**Concentration** Adding 0.2 ml of distilled water will yield a concentration of 500 μg/ml.

**Storage** Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -

20°C in small aliquots to prevent freeze-thaw cycles.

**Note** For research use only

**Application notes** Western blot, 0.25-0.5 μg/ml, Human

Immunocytochemistry/Immunofluorescence, 5 μg/ml, Human

Immunoprecipitation, 0.5-2  $\mu$ g/ml, Human Flow Cytometry(Fixed), 1-3  $\mu$ g/1x106 cells, Human ELISA, 0.1-0.5  $\mu$ g/ml, -. Adding 0.2 ml of distilled water will yield a

concentration of 500 µg/ml

**Isotype** Rabbit IgG

**Clonality** Polyclonal

**Antibody Type** Primary Antibody

**MW** 120 kDa



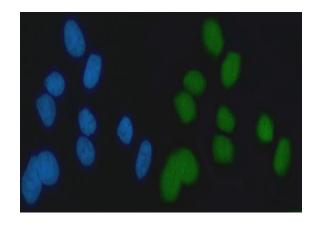


#### **Uniprot ID**

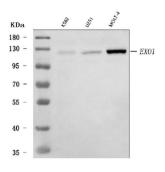
#### **Q9UQ84**

### **Expiration Date**

12 months from date of receipt.



IF analysis of Exonuclease 1/EXO1 using anti-Exonuclease 1/EXO1 antibody. Exonuclease 1/EXO1 was detected in an immunocytochemical section of MCF-7 cells. Enzyme antigen retrieval was performed using IHC enzyme antigen retrieval reagent for 15 mins. The cells were blocked with 10% goat serum. And then incubated with 5 µg/mL rabbit anti-Exonuclease 1/EXO1 Antibody overnight at 4°C. DyLight®488 Conjugated Goat Anti-Rabbit IgG was used as secondary antibody at 1:100 dilution and incubated for 30 minutes at 37°C. The section was counterstained with DAPI. Visualize using a fluorescence microscope and filter sets appropriate for the label used.



Western blot analysis of Exonuclease 1/EXO1 using anti-Exonuclease 1/EXO1 antibody. Electrophoresis was performed on a 5-20% SDS-PAGE gel at 70V (Stacking gel) / 90V (Resolving gel) for 2-3 hours. The sample well of each lane was loaded with 30 ug of sample under reducing conditions. Lane 1: human K562 whole cell lysates, Lane 2: human U251 whole cell lysates, Lane 3: human MOLT-4 whole cell lysates. After electrophoresis, proteins were transferred to a nitrocellulose membrane at 150 mA for 50-90 minutes. Blocked the membrane with 5% non-fat milk/TBS for 1.5 hour at RT. The membrane was incubated with rabbit anti-Exonuclease 1/EXO1 antigen affinity purified polyclonal antibody at 0.5 μg/mL overnight at 4°C, then washed with TBS-0.1% Tween 3 times with 5 minutes each and probed with a goat anti-rabbit IgG-HRP secondary antibody at a dilution of 1:5000 for 1.5 hour at RT. The signal is developed using an Enhanced Chemiluminescent detection (ECL) kit with Tanon 5200 system. A specific band was detected for Exonuclease 1/EXO1 at approximately 120 kDa. The expected band size for Exonuclease 1/EXO1 is at 94 kDa.