

## **Product Datasheet**

## Anti-Neutrophil Elastase/ELANE Antibody (orb381060)

**Description** Anti-Neutrophil Elastase/ELANE Antibody

**Species/Host** Rabbit

**Reactivity** Mouse, Rat

**Conjugation** Unconjugated

**Tested Applications** ELISA, IHC, WB

Immunogen E. coli-derived mouse Elastase/ELANE/ELA2 recombinant protein (Position: S27-

N265). Mouse Elastase/ELANE/ELA2 shares 92% amino acid (aa) sequence

identity with rat Elastase/ELANE/ELA2.

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, 0.1-0.5μg/ml, Mouse Immunohistochemistry (Paraffin-embedded

Section), 2-5µg/ml, Mouse, Rat ELISA, 0.1-0.5µg/ml,-. Add 0.2ml of distilled water

will yield a concentration of 500ug/ml

**Isotype** Rabbit IgG

**Clonality** Polyclonal

**MW** 29 kDa

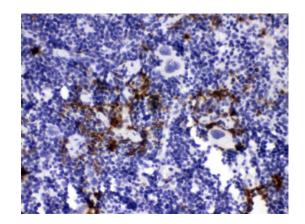
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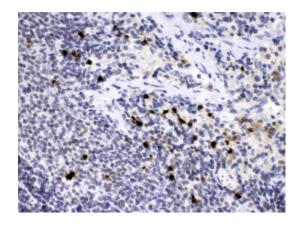


## **Expiration Date**

12 months from date of receipt.



IHC analysis of Elastase/ELANE/ELA2 using anti-Elastase/ELANE/ELA2 antibody. Elastase/ELANE/ELA2 was detected in a paraffin-embedded section of mouse spleen tissue. Heat mediated antigen retrieval was performed in EDTA buffer (pH8.0, epitope retrieval solution). The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 1  $\mu$ g/ml rabbit anti-Elastase/ELANE/ELA2 Antibody overnight at 4°C. Biotinylated goat anti-rabbit IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using Strepavidin-Biotin-Complex (SABC) with DAB as the chromogen.



IHC analysis of Elastase/ELANE/ELA2 using anti-Elastase/ELANE/ELA2 antibody. Elastase/ELANE/ELA2 was detected in a paraffin-embedded section of rat spleen tissue. Heat mediated antigen retrieval was performed in EDTA buffer (pH8.0, epitope retrieval solution). The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 1  $\mu$ g/ml rabbit anti-Elastase/ELANE/ELA2 Antibody overnight at 4°C. Biotinylated goat anti-rabbit IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using Strepavidin-Biotin-Complex (SABC) with DAB as the chromogen.

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Western blot analysis of Elastase/ELANE/ELA2 using anti-Elastase/ELANE/ELA2 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: mouse bone tissue lysates, Lane 2: mouse spleen tissue 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-Elastase/ELANE/ELA2 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 Elastase/ELANE/ELA2 at approximately 29 kDa. The expected band size for Elastase/ELANE/ELA2 is at 29 kDa.