

## Basic Information

|                    |   |  |
|--------------------|---|--|
| Product Name       | Anti-CD90/THY1 Antibody   |  |
| Gene Name          | THY1  |  |
| Source             | Rabbit  |  |
| Isotype            | IgG   |  |
| Species Reactivity | human, mouse, rat   |  |
| Tested Application | WB, IHC, IF, FCM  |  |
| Contents           | 500 ug/ml antibody with PBS , 0.02% NaN <sub>3</sub> , 1 mg BSA and 50% glycerol.   |  |
| Immunogen          | E.coli-derived mouse CD90/Thy1 recombinant protein (Position: Q20-C131). Mouse CD90/Thy1 shares 63.4% and 81.3% amino acid (aa) sequence identity with human and rat CD90/Thy1, respectively.   |  |
| concentration      | 500 ug/ml   |  |
| Purification       | Immunogen affinity purified.  |  |
| Observed MW        | 22-30KD   |  |
| Dilution Ratios    | Western blot(WB): 1:500-2000<br>Immunohistochemistry in paraffin section (IHC): 1:50-400<br>Immunofluorescence (IF): 1:50-400<br>Flow cytometry (FCM): 1-3 µg/1x10 <sup>6</sup> cells<br>(Boiling the paraffin sections in 10mM citrate buffer,pH6.0,or PH8.0 EDTA repair liquid for 20 mins is required for the staining of formalin/paraffin sections.) Optimal working dilutions must be determined by end user. |  |

## Storage

12 months from date of receipt, -20°C as supplied. 6 months 2 to 8°C after reconstitution. Avoid repeated freezing and thawing.

## Background Information

CD90 (Cluster of Differentiation 90) or Thy-1 is a 25-37 kDa heavily N-glycosylated, glycoposphatidylinositol (GPI) anchored conserved cell surface protein with a single V-like immunoglobulin domain, originally discovered as a thymocyte antigen. The CD90 gene is mapped to 11q23.3. Thy-1 can be used as a marker for a variety of stem cells and for the axonal processes of mature neurons. Structural study of Thy-1 lead to the foundation of the Immunoglobulin superfamily, of which it is the smallest member, and led to the first biochemical description and characterization of a vertebrate GPI anchor.

## Reference

Anti-CD90/THY1 Antibody被引用在1文献中。

## Selected Validation Data

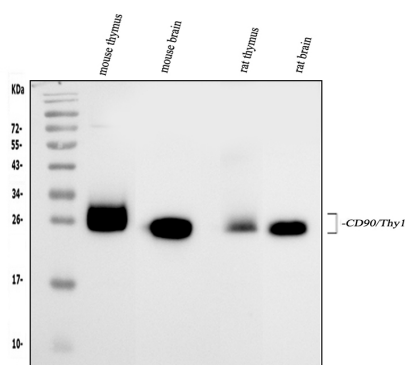


Figure 1. Western blot analysis of anti- CD90/THY1 antibody (A01818). The sample well of each lane was loaded with 50ug of sample under reducing conditions.

Lane 1: mouse thymus tissue lysates,

Lane 2: mouse brain tissue lysates,

Lane 3: rat thymus tissue lysates,

Lane 4: rat brain tissue lysates.

Use rabbit anti- CD90/THY1 1:1000, probed with a goat anti-rabbit IgG-HRP secondary antibody. The signal is developed using an Enhanced Chemiluminescent detection (ECL) kit (Catalog # EK1002). A specific band was detected for CD90/THY1 at approximately 24-30 kDa. The expected band size for CD90/THY1 is at 18 kDa.

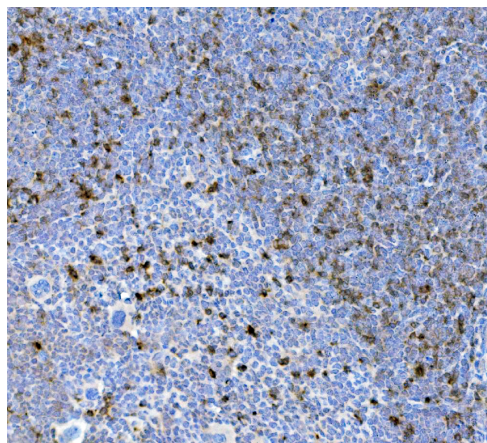


Figure 2. IHC analysis of CD90/Thy1 using anti- CD90/Thy1 antibody (A01818). anti-CD90/Thy1 Antibody (A01818) was detected in paraffin-embedded section of mouse spleen tissues. Biotinylated goat anti-rabbit IgG was used as secondary antibody. The tissue section was developed using Streptavidin-Biotin-Complex (SABC) (Catalog # SA1022) with DAB as the chromogen.

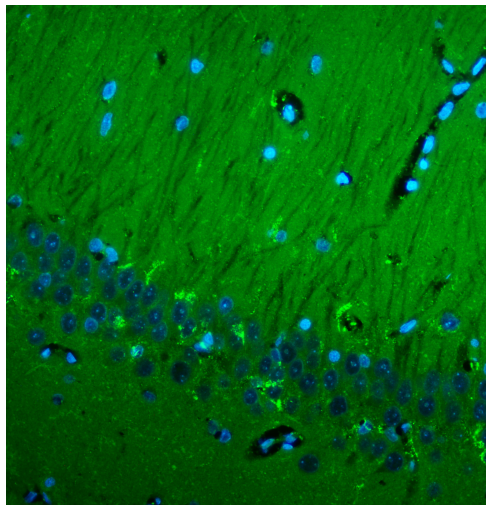


Figure 4. IF analysis using anti- CD90/THY1 antibody (A01818). detected in paraffin-embedded section of rat brain tissue. The tissue section were stained using the Dylight488-conjugated Anti-rabbit IgG Secondary Antibody (green) (Catalog # BA1127) and counterstained with DAPI (blue).

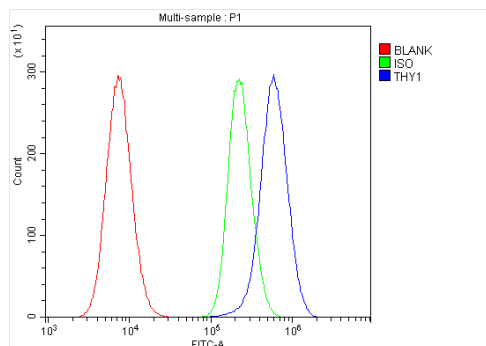


Figure 5. Flow cytometry analysis of HEPA1-6 cell (1x106) DyLight 488 conjugated goat anti- rabbit IgG(blue) was used as secondary antibody. Isotype control antibody (Green line) was rabbit IgG DyLight 488. Unlabelled sample (Red line).