### SONY

# **Product Data Sheet**

#### FITC anti-human CD45

**Catalog # / Size:** 2120025 / 25 tests

2120030 / 100 tests

2120190 / 500 tests

2120270 / 100 µg

Clone: HI30

**Isotype:** Mouse IgG1, κ

Reactivity: Human

**Preparation:** The antibody was purified by affinity

chromatography, and conjugated with FITC under optimal conditions. The solution is free of unconjugated FITC.

**Formulation:** microg size: Phosphate-buffered

solution, pH 7.2, containing 0.09%

sodium azide.

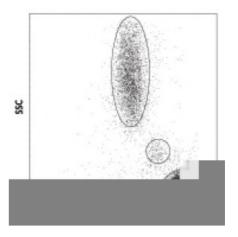
test sizes: Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide and 0.2% (w/v) BSA (origin USA).

Workshop IV N816

Number:

Concentration: microg sizes: 0.5 mg/ml

test sizes: lot-specific



Human peripheral blood lymphocytes, monocytes and granulocytes stained with HI30 FITC

## **Applications:**

**Applications:** Flow Cytometry

Recommended

**Usage:** 

Each lot of this antibody is quality control tested by immunofluorescent staining with flow cytometric analysis. For flow cytometric staining using the microg size, the suggested use of this reagent is  $\leq 1.0$  microg per million cells in 100 microL volume. **Test size products are transitioning from 20 microL to 5 microL per test**. Please check your vial or your CoA to find the suggested use of this reagent per million cells in 100 microL staining volume or per 100 microL of whole blood. It is recommended that the reagent be titrated for optimal performance for each application.

Application Notes:

Additional reported applications (for the relevant formats) include: immunohistochemical staining of acetone-fixed frozen tissue sections and formalin-fixed paraffin-embedded tissue sections<sup>9</sup>, inhibition of CD45 functions4, immunofluorescence<sup>11</sup>, and Western blotting3.

It was found that the HI30 clone and the 2D1 clone can cross block each other's binding.

Application References:

- 1. Knapp W, et al. 1989. Leucocyte Typing IV. Oxford University Press. New York.
- 2. Kishihara K, et al. 1993. Cell 74:143.
- 3. Esser M, et al. 2001. J. Virol. 75:6173. (WB)
- 4. Yamada T, et al. 2002. J. Biol. Chem. 277:28830.
- 5. Nagano M, et al. 2007. Blood 110:151.
- 6. Jiang Q, et al. 2008. Blood 112:2858. PubMed
- 7. Morozov A, et al. 2010. Clin Cancer Res. 16:5630. PubMed
- 8. Yoshino N, et al. 2000. Exp. Anim. (Tokyo) 49:97. (FC)

- 9. Friedman T, et al. 1999. J. Immunol. 162:5256. (IHC)
- 10. Oeztuerk-Winder F, et al. 2012. EMBO J. 31:3431. (FC) PubMed
- 11. Rees LE, et al. 2003. Clin. Exp. Immunol. 134:497. (IF)
- 12. Lee J, et al. 2015. J Exp Med. 212:385. PubMed
- 13. Breton G, et al. 2015. J Exp Med. 212:401. PubMed
- 14. Marquardt N, et al. 2015. J Immunol. 6:2467. PubMed

#### **Description:**

CD45 is a 180-240 kD single chain type I membrane glycoprotein also known as leukocyte common antigen (LCA) and T200. It is a tyrosine phosphatase expressed on the plasma membrane of all hematopoietic cells, except erythrocytes and platelets. CD45 is a signaling molecule that regulates a variety of cellular processes including cell growth, differentiation, cell cycle, and oncogenic transformation. CD45 plays a critical role in T and B cell antigen receptor-mediated activation by dephosphorylating substrates including p56Lck, p59Fyn, and other Src family kinases. CD45 non-covalently associates with lymphocyte phosphatase-associated phosphoprotein (LPAP) on T and B lymphocytes. CD45 has been reported to bind galectin-1 and to be associated with several other cell surface antigens including CD1, CD2, CD3, and CD4.

# Antigen References:

- 1. Thomas M. 1989. Annu. Rev. Immunol. 7:339.
- **s:** 2. Trowbridge I, *et al.* 1994. *Annu. Rev. Immunol.*12:85.