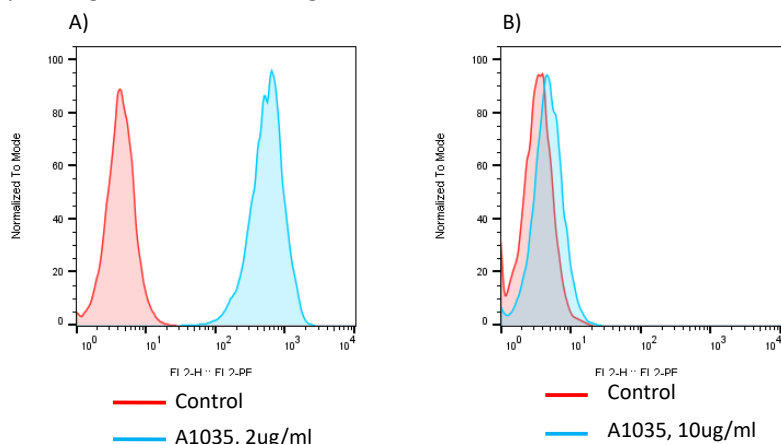


SPECIFICATIONS

Catalog Number	A1035
Product Name	Anti-human B7-H3 antibody
Source	Mouse hybridoma
Clone	6B1
Species Reactivity	Human and cynomolgus B7-H3, does not bind to mouse and hamster B7-H3
Isotype	mouse IgG1
Formulation	1x PBS, pH7.0. Sterile
Stability & Storage	1 month at 4°C; 12 months at <-20°C; Avoid repeated freeze-thaw
Purity	>95%
Protein Aggregation	Not obvious on SDS-PAGE
Application	Flow cytometry, ELISA, cell-based assay

DATA

Detection of human B7-H3 expression on A) a human B7-H3-CHO-K1 cell line (Cat.#C3092) by flow cytometry. Anti-human B7-H3 mouse monoclonal antibody 6B1 (Cat.#A1035) was incubated with A) human B7-H3-CHO-K1 cells (Cat.#C3092) and B) vector control CHO-K1 cells (Cat.#C3022), followed by staining with PE-anti-mouse IgG.



BACKGROUND

B7-H3, also known as CD276, is a cell surface protein that belongs to the B7 family of immune regulatory molecules. B7-H3 has two isoforms determined by its extracellular domain. In mice, the extracellular domain consists of a single pair of immunoglobulin variable (IgV)-like and immunoglobulin constant (IgC)-like domains, whereas in humans it consists of one pair (2Ig-B7-H3) or two identical pairs (4Ig-B7-H3) due to exon duplication. B7-H3 mRNA is expressed in most normal tissues. Flow cytometric analysis demonstrated inducible expression of B7H3 on monocytes, dendritic cells, and T cells after stimulation with selected cytokines and mitogens. B7-H3 protein is expressed at high frequency on many different cancer types (60% of all cancers). B7-H3 has both costimulatory and coinhibitory properties that can affect the proliferation of CD4⁺ and CD8⁺ T cells, production of cytokines, and activity of T cells and NK cells depending on the microenvironment. B7-H3 also exhibits nonimmunological pro-tumorigenic functions such as migration and invasion, apoptosis, cell viability and chemoresistance.

References

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- Picarda E, Ohaegbulam KC, Zang X. Molecular Pathways: Targeting B7-H3 (CD276) for Human Cancer Immunotherapy. *Clin Cancer Res*.22:3425-3431. 2016.

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