

# Product datasheet for AM05187AC-N

## CD28 Mouse Monoclonal Antibody [Clone ID: B-23]

### **Product data:**

#### **Product Type: Primary Antibodies Clone Name:** B-23 FC **Applications: Recommended Dilution:** Flow Cytometry. Monitoring of activated T cells in peripheral blood. Study of T lymphocyte cytokine function. Study of B cell activation. Study of Cell-adhesion molecules relating T and B lymphocytes. **Reactivity:** Human Host: Mouse lgG1 Isotype: **Clonality:** Monoclonal Immunogen: CD28=Derived from the hybridization of mouse Sp2/O-Ag14 myeloma cells with spleen cells of BALB/c mice immunized with the HPB-ALL T-cell line. This antibody recats with CD28 receptor. Specificity: Formulation: PBS containing 0.2% protein carrier and 0.08% Sodium Azide as preservative. Label: APC State: Liquid purified IgG fraction. **Concentration:** lot specific APC **Conjugation:** Store the antibody undiluted at 2-8°C. Storage: **Do Not Freeze!** Stability: Shelf life: One year from despatch. Gene Name: CD28 molecule Database Link: Entrez Gene 940 Human P10747



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	CD28 Mouse Monoclonal Antibody [Clone ID: B-23] – AM05187AC-N
Background:	Anti-human CD28 binds the 44kDa MW cell surface protein on the surface of most T cells. CD28 acts as the ligand for the B7/BB-1 molecule on the surface of activated B cells. B7/BB-1 costimulates T cells through CD28, along with CD2 and CD3.CD28 antigen is a disulfide-linked homodimeric glycoprotein. The CD28 antigen is present on approximately 60%-80% T lymphocytes (95% of CD4 and 50% of CD8 lymphocytes). CD28 regulates the expression of cytokines by T cells, not only IL-2, but also IL-1 alpha and CSF-1, usually synthesized by accessory cells. CD28 functions as a cell adhesion molecule (CAM) for certain T cell subsets.
Synonyms:	TP44
Note:	Protocol: <b>PBMC:</b> Add 10 ul of MAB/10e6 PBMC in 100 ul PBS. Mix gently and incubate for 15 minutes at 2-8°C. Wash twice with PBS and analyze or fix with 0.5% v/v of paraformaldehyde in PBS and analyze.
	<b>WHOLE BLOOD:</b> Add 10 ul of MAB/100 ul of whole blood. Mix gently and incubate for 15 minutes at room temperature (20°C). Lyse the whole blood. Wash once with PBS and analyze or fix with 0.5% v/v of paraformaldehyde in PBS and

analyze. See instrument manufacturers instructions for Lysed Whole Blood and Immunofluorescence analysis with a flow cytometer or microscope.

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