

## DESCRIPTION

<b>Species Reactivity</b>	Human
<b>Specificity</b>	Detects human GATA-3 in direct ELISAs.
<b>Source</b>	Monoclonal Mouse IgG <sub>2B</sub> Clone # 634919
<b>Purification</b>	Protein A or G purified from hybridoma culture supernatant
<b>Immunogen</b>	<i>E. coli</i> -derived recombinant human GATA-3 Pro135-Ser258 Accession # P23771
<b>Formulation</b>	Lyophilized from a 0.2 µm filtered solution in PBS with Trehalose. See Certificate of Analysis for details. *Small pack size (-SP) is supplied either lyophilized or as a 0.2 µm filtered solution in PBS.

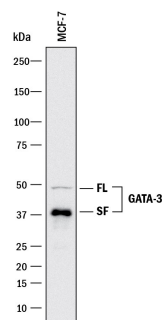
## APPLICATIONS

**Please Note:** Optimal dilutions should be determined by each laboratory for each application. [General Protocols](#) are available in the Technical Information section on our website.

	<b>Recommended Concentration</b>	<b>Sample</b>
<b>Western Blot</b>	2 µg/mL	See Below
<b>Intracellular Staining by Flow Cytometry</b>	0.25 µg/10 <sup>6</sup> cells	See Below

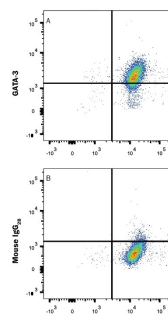
## DATA

### Western Blot



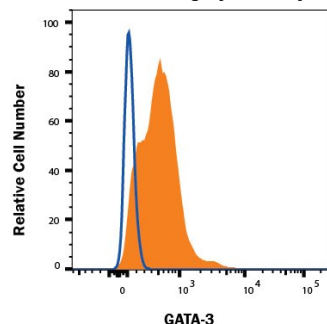
**Detection of Human GATA-3 by Western Blot.** Western blot shows lysates of MCF-7 human breast cancer cell line. PVDF membrane was probed with 2 µg/mL of Mouse Anti-Human GATA-3 Monoclonal Antibody (Catalog # MAB26052) followed by HRP-conjugated Anti-Mouse IgG Secondary Antibody (Catalog # Catalog # [HAF018](#)). Specific bands were detected for GATA-3 full length (FL) at approximately 50 kDa and the splice form (SF) at approximately 37 kDa (as indicated). This experiment was conducted under reducing conditions and using [Immunoblot Buffer Group 3](#).

### Intracellular Staining by Flow Cytometry



**Detection of GATA-3 in Human PBMCs by Flow Cytometry.** Human peripheral blood mononuclear cells (PBMCs) stimulated to induce Th2 cells were stained with Mouse Anti-Human CD4 PE-conjugated Monoclonal Antibody (Catalog # [FAB3791P](#)) and either (A) Mouse Anti-Human GATA-3 Monoclonal Antibody (Catalog # MAB26052) or (B) Mouse IgG<sub>2B</sub> Flow Cytometry Isotype Control (Catalog # [MAB0041](#)) followed by Allophycocyanin-conjugated Anti-Mouse IgG Secondary Antibody (Catalog # [F0101B](#)). To facilitate intracellular staining, cells were fixed and permeabilized with FlowX FoxP3 Fixation & Permeabilization Buffer Kit (Catalog # [FC012](#)). View our protocol for [Staining Intracellular Molecules](#).

### Intracellular Staining by Flow Cytometry



**Detection of GATA-3 in MCF-7 cells by Flow Cytometry.** MCF-7 cells were stained with Mouse Anti-Human GATA-3 Monoclonal Antibody (Catalog # MAB26052, filled histogram) or isotype control antibody (Catalog # [MAB0041](#), open histogram), followed by Allophycocyanin-conjugated Anti-Mouse IgG Secondary Antibody (Catalog # [F0101B](#)). To facilitate intracellular staining, cells were fixed with [FC012](#) and permeabilized with FoxP3 Perm. View our protocol for [Staining Intracellular Molecules](#).

**PREPARATION AND STORAGE**

**Reconstitution** Reconstitute at 0.5 mg/mL in sterile PBS.

**Shipping** The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below.  
\*Small pack size (-SP) is shipped with polar packs. Upon receipt, store it immediately at -20 to -70 °C

**Stability & Storage** Use a manual defrost freezer and avoid repeated freeze-thaw cycles.

- 12 months from date of receipt, -20 to -70 °C as supplied.
- 1 month, 2 to 8 °C under sterile conditions after reconstitution.
- 6 months, -20 to -70 °C under sterile conditions after reconstitution.

**BACKGROUND**

GATA-3 belongs to the GATA family of transcription factors, which bind to the consensus DNA sequence (A/T) GATA (A/G) to control diverse tissue-specific programs of gene expression and morphogenesis. It is widely expressed in mesodermal- and endodermal-derived tissues. GATA-3 has been shown to be an essential regulator for immune cell function, sympathetic neuron development and the maintenance of the differentiated state in epithelial cells.