

## DESCRIPTION

<b>Species Reactivity</b>	Human
<b>Specificity</b>	Detects human TRA-1-85 antigen in Western blots.
<b>Source</b>	Monoclonal Mouse IgG <sub>1</sub> Clone # TRA-1-85
<b>Purification</b>	Protein A or G purified from hybridoma culture supernatant
<b>Immunogen</b>	2120Ep human embryonal carcinoma cell line
<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.

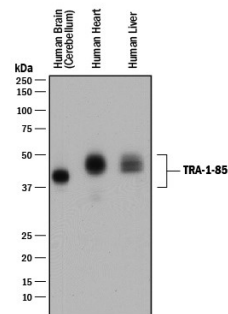
## APPLICATIONS

**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>	1 µg/mL	See Below
<b>Flow Cytometry</b>	0.25 µg/10 <sup>6</sup> cells	See Below
<b>Immunocytochemistry</b>	8-25 µg/mL	See Below
<b>CyTOF-ready</b>	Ready to be labeled using established conjugation methods. No BSA or other carrier proteins that could interfere with conjugation.	

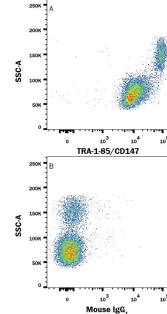
## DATA

### Western Blot



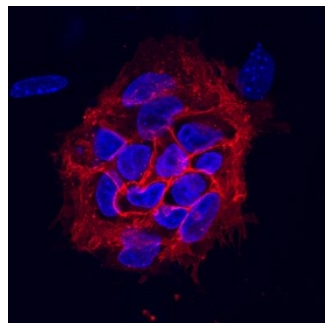
**Detection of Human TRA-1-85/CD147 by Western Blot.** Western blot shows lysates of human brain (cerebellum) tissue, human heart tissue, and human liver tissue. PVDF membrane was probed with 1 µg/mL of Mouse Anti-Human TRA-1-85/CD147 Monoclonal Antibody (Catalog # MAB3195) followed by HRP-conjugated Anti-Mouse IgG Secondary Antibody (Catalog # HAF018). A specific band was detected for TRA-1-85/CD147 at approximately 40-60 kDa (as indicated). This experiment was conducted under reducing conditions and using *Immunoblot Buffer Group 1*.

### Flow Cytometry



**Detection of TRA-1-85/CD147 in Human PBMCs by Flow Cytometry.** Human peripheral blood lymphocytes and monocytes were stained with either (A) Mouse Anti-Human TRA-1-85/CD147 Monoclonal Antibody (Catalog # MAB3195) or (B) Mouse IgG<sub>1</sub> Isotype Control (Catalog # MAB002) followed by Allophycocyanin-conjugated Anti-Mouse IgG Secondary Antibody (Catalog # F0101B).

### Immunocytochemistry



**TRA-1-85/CD147 in BG01V Human Embryonic Stem Cells.** TRA-1-85/CD147 was detected in immersion fixed BG01V human embryonic stem cells on irradiated mouse embryonic fibroblasts using Mouse Anti-Human TRA-1-85/CD147 Monoclonal Antibody (Catalog # MAB3195) at 10 µg/mL for 3 hours at room temperature. Cells were stained using the NorthernLights™ 557-conjugated Anti-Mouse IgG Secondary Antibody (red; Catalog # NL007) and counterstained with DAPI (blue). Specific staining was localized to cytoplasm. View our protocol for *Fluorescent ICC Staining of Cells on Coverslips*.

## PREPARATION AND STORAGE

<b>Reconstitution</b>	Reconstitute at 0.5 mg/mL in sterile PBS.
<b>Shipping</b>	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
<b>Stability &amp; Storage</b>	<b>Use a manual defrost freezer and avoid repeated freeze-thaw cycles.</b> <ul style="list-style-type: none"> <li>• 12 months from date of receipt, -20 to -70 °C as supplied.</li> <li>• 1 month, 2 to 8 °C under sterile conditions after reconstitution.</li> <li>• 6 months, -20 to -70 °C under sterile conditions after reconstitution.</li> </ul>

## BACKGROUND

The TRA-1-85 antigen, also known as OKa blood group antigen, is a specific epitope within the protein known as Basigin, EMMPRIN and CD147. It is a cell surface determinant expressed on almost all human cell types. This antibody has been used in somatic cell hybrid studies to identify tissues of partial human origin (1, 2).

## References:

1. Williams, B.P. *et al.* (1988) *Immunogenetics*. **27**:322.
2. Spring, F.A. *et al.* (1997) *Eur. J. Immunol.* **27**:891.