

DESCRIPTION

Species Reactivity	Human
Specificity	Detects human Integrin α L in Western blots.
Source	Monoclonal Mouse IgG ₁ Clone # 345908
Purification	Protein A or G purified from hybridoma culture supernatant
Immunogen	Chinese hamster ovary cell line CHO-derived recombinant human Integrin α L Tyr26-Met1089 (Tyr660Ile) Accession # CAA68747
Formulation	Lyophilized from a 0.2 μ m filtered solution in PBS with Trehalose.

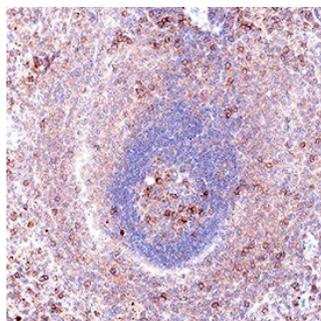
APPLICATIONS

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.

	Recommended Concentration	Sample
Western Blot	2 μ g/mL	Jurkat human acute T cell leukemia cell line
Immunohistochemistry	5-15 μ g/mL	Immersion fixed paraffin-embedded sections of Human Spleen
Simple Western	50 μ g/mL	Human PBMCs

DATA

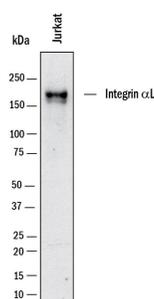
Immunohistochemistry



Detection of Integrin α L/CD11a in Human Spleen.

Integrin α L/CD11a was detected in immersion fixed paraffin-embedded sections of Human Spleen using Mouse Anti-Human Integrin α L/CD11a Monoclonal Antibody (Catalog # MAB35952) at 5 μ g/mL for 1 hour at room temperature followed by incubation with the Anti-Mouse IgG VisUCyte™ HRP Polymer Antibody (Catalog # VC001). Before incubation with the primary antibody, tissue was subjected to heat-induced epitope retrieval using VisUCyte Antigen Retrieval Reagent-Basic (Catalog # VCTS021). Tissue was stained using DAB (brown) and counterstained with hematoxylin (blue). Specific staining was localized to lymphocytes. View our protocol for IHC Staining with VisUCyte HRP Polymer Detection Reagents.

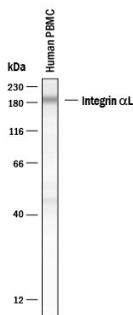
Western Blot



Detection of Human Integrin α L/CD11a by Western Blot.

Western blot shows lysates of Jurkat Human Acute T Cell Leukemia Cells. PVDF membrane was probed with 2 μ g/mL of Mouse Anti-Human Integrin α L/CD11a Monoclonal Antibody (Catalog # MAB35952) followed by HRP-conjugated Anti-Mouse IgG Secondary Antibody (Catalog # HAF018). A specific band was detected for Integrin α L/CD11a at approximately 160 kDa (as indicated). This experiment was conducted under reducing conditions and using Western Blot Buffer Group 1.

Simple Western



Detection of Human Integrin α L/CD11a by Simple Western™.

Simple Western lane view shows lysates of Human PBMCs, loaded at 0.2 mg/mL. A specific band was detected for Integrin α L/CD11a at approximately 190 kDa (as indicated) using 50 μ g/mL of Mouse Anti-Human Integrin α L/CD11a Monoclonal Antibody (Catalog # MAB35952). This experiment was conducted under reducing conditions and using the 12-230 kDa separation system.



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.
Stability & Storage	<p>Use a manual defrost freezer and avoid repeated freeze-thaw cycles.</p> <ul style="list-style-type: none"> • 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

Integrin subunit α L/CD11a is a 180 kDa type I TM glycoprotein that interacts only with Integrin β 2/CD18 to form LFA-1, a leukocyte adhesion protein which binds endothelial cell ICAM. Human Integrin α L contains a 1064 aa extracellular domain (ECD), a 20 aa TM sequence and a 58 aa cytoplasmic domain. The ECD contains seven repeats that form a beta-propeller structure and one inserted vWA domain (I domain) containing a metal ion-dependent adhesion site (MIDAS). Human and mouse Integrin α L ECD share 74% aa identity. A second isoform has a 53 aa insert in the ECD.