

DESCRIPTION

Species Reactivity	Human/Canine
Specificity	Detects human Integrin β 1/CD29 in direct ELISAs and Western blots. In direct ELISAs, approximately 25% cross-reactivity with recombinant mouse (rm) Integrin β 1 is observed and less than 1% cross-reactivity with recombinant human (rh) Integrin β 2, rhIntegrin β 3, rhIntegrin β 5, and rhIntegrin β 7 is observed.
Source	Polyclonal Goat IgG
Purification	Antigen Affinity-purified
Immunogen	Chinese hamster ovary cell line CHO-derived recombinant human Integrin β 1 isoform 1A Gln21-Asp728 Accession # P05556
Formulation	Lyophilized from a 0.2 μ m 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 μ m filtered solution in PBS.

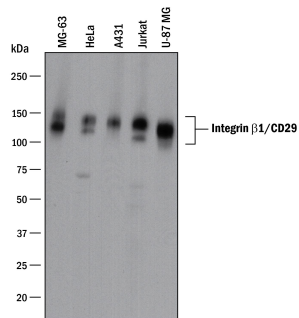
APPLICATIONS

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.

	Recommended Concentration	Sample
Western Blot	μ g/mL	See Below
Flow Cytometry	0.25 μ g/ 10^6 cells	See Below
Immunocytochemistry	5-15 μ g/mL	See Below
Simple Western	10 μ g/mL	See Below
CyTOF-ready	Ready to be labeled using established conjugation methods. No BSA or other carrier proteins that could interfere with conjugation.	
Knockout Validated	Integrin β 1/CD29 is specifically detected in HeLa human cervical epithelial carcinoma parental cell line but is not detectable in Integrin β 1/CD29 knockout HeLa cell line.	

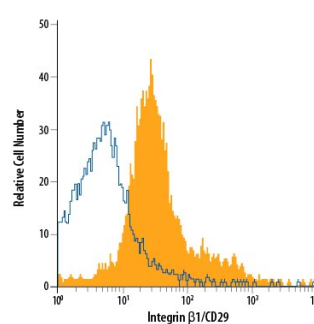
DATA

Western Blot



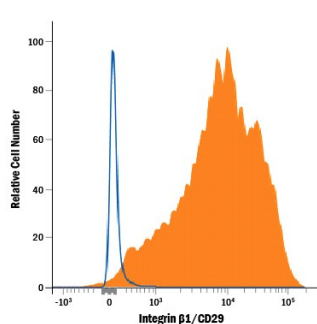
Detection of Human Integrin β 1/CD29 by Western Blot. Western blot shows lysates of MG-63 human osteosarcoma cell line, HeLa human cervical epithelial carcinoma cell line, A431 human epithelial carcinoma cell line, Jurkat human acute T cell leukemia cell line, and U-87 MG human glioblastoma/astrocytoma cell line. PVDF membrane was probed with 1 μ g/mL of Goat Anti-Human/Canine Integrin β 1/CD29 Antigen Affinity-purified Polyclonal Antibody (Catalog # AF1778) followed by HRP-conjugated Anti-Goat IgG Secondary Antibody (Catalog # HAF017). Specific bands were detected for Integrin β 1/CD29 at approximately 130-140 kDa (as indicated). This experiment was conducted under reducing conditions and using Immunoblot Buffer Group 1.

Flow Cytometry



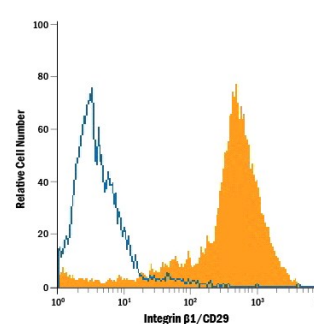
Detection of Integrin β 1/CD29 in Canine PBMCs by Flow Cytometry. Canine peripheral blood mononuclear cells (PBMCs) were stained with Goat Anti-Human/Canine Integrin β 1/CD29 Antigen Affinity-purified Polyclonal Antibody (Catalog # AF1778, filled histogram) or isotype control antibody (Catalog # AB-108-C, open histogram), followed by Phycoerythrin-conjugated Anti-Goat IgG Secondary Antibody (Catalog # F0107).

Flow Cytometry



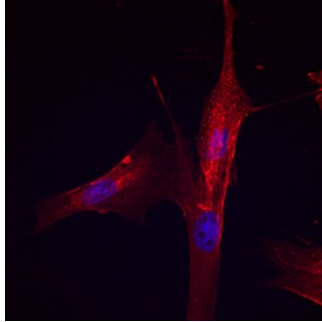
Detection of Integrin β 1/CD29 in Human PBMCs by Flow Cytometry. Human peripheral blood mononuclear cells (PBMCs) were stained with Goat Anti-Human/Canine Integrin β 1/CD29 Antigen Affinity-purified Polyclonal Antibody (Catalog # AF1778, filled histogram) or isotype control antibody (Catalog # AB-108-C, open histogram), followed by Phycoerythrin-conjugated Anti-Goat IgG Secondary Antibody (Catalog # F0107).

Flow Cytometry



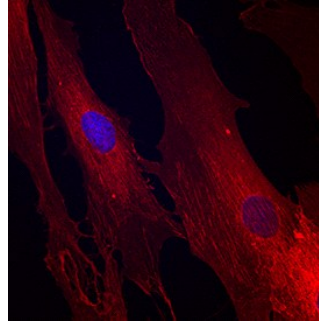
Detection of Integrin β 1/CD29 in Canine Mesenchymal Stem Cells by Flow Cytometry. Canine mesenchymal stem cells were stained with Goat Anti-Human/Canine Integrin β 1/CD29 Antigen Affinity-purified Polyclonal Antibody (Catalog # AF1778, filled histogram) or isotype control antibody (Catalog # AB-108-C, open histogram), followed by Phycoerythrin-conjugated Anti-Goat IgG Secondary Antibody (Catalog # F0107).

Immunocytochemistry



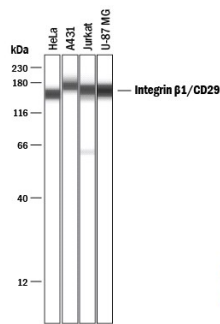
Integrin β 1/CD29 in Canine Mesenchymal Stem Cells. Integrin β 1/CD29 was detected in immersion fixed canine mesenchymal stem cells using Goat Anti-Human/Canine Integrin β 1/CD29 Antigen Affinity-purified Polyclonal Antibody (Catalog # AF1778) at 10 μ g/mL for 3 hours at room temperature. Cells were stained using the NorthernLights™ 557-conjugated Anti-Goat IgG Secondary Antibody (red; Catalog # NL001) and counterstained with DAPI (blue). Specific staining was localized to cell surfaces. View our protocol for [Fluorescent ICC Staining of Stem Cells on Coverslips](#).

Immunocytochemistry



Integrin β 1/CD29 in Human Mesenchymal Stem Cells. Integrin β 1/CD29 was detected in immersion fixed human mesenchymal stem cells using Goat Anti-Human/Canine Integrin β 1/CD29 Antigen Affinity-purified Polyclonal Antibody (Catalog # AF1778) at 10 μ g/mL for 3 hours at room temperature. Cells were stained using the NorthernLights™ 557-conjugated Anti-Goat IgG Secondary Antibody (red; Catalog # NL001) and counterstained with DAPI (blue). Specific staining was localized to cell surfaces. View our protocol for [Fluorescent ICC Staining of Stem Cells on Coverslips](#).

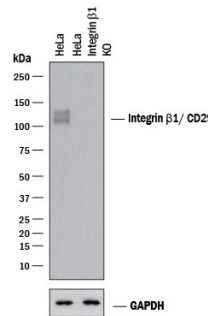
Simple Western



Detection of Human Integrin β 1/CD29 by Simple Western™. Simple Western lane view shows lysates of HeLa human cervical epithelial carcinoma cell line, A431 human epithelial carcinoma cell line, Jurkat human acute T cell leukemia cell line, and U-87 MG human glioblastoma/astrocytoma cell line, loaded at 0.2 mg/mL. Specific bands were detected for Integrin β 1/CD29 at approximately 157-176 kDa (as indicated) using 10 μ g/mL of Goat Anti-Human/Canine Integrin β 1/CD29 Antigen Affinity-purified Polyclonal Antibody (Catalog # AF1778) followed by 1:50 dilution of HRP-conjugated Anti-Goat IgG Secondary Antibody (Catalog # HAF109). This experiment was conducted under reducing conditions and using the 12-230 kDa separation system.



Knockout Validated



Western Blot Shows Human Integrin β 1/CD29 Specificity by Using Knockout Cell Line. Western blot shows lysates of HeLa human cervical epithelial carcinoma parental cell line and Integrin β 1/CD29 knockout HeLa cell line (KO). PVDF membrane was probed with 2 μ g/mL of Goat Anti-Human/Canine Integrin β 1/CD29 Antigen Affinity-purified Polyclonal Antibody (Catalog # AF1778) followed by HRP-conjugated Anti-Goat IgG Secondary Antibody (Catalog # HAF017). Specific bands were detected for Integrin β 1/CD29 at approximately 110-120 kDa (as indicated) in the parental HeLa cell line, but is not detectable in knockout HeLa cell line. GAPDH (Catalog # MAB5718) is shown as a loading control. This experiment was conducted under reducing conditions and using Immunoblot Buffer Group 1.

PREPARATION AND STORAGE

Reconstitution	Reconstitute at 0.2 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	<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 β 1, also called CD29 and VLA- β chain, associates with at least ten different integrin α subunits to form various VLA complexes. The β 1 subunit has a broad tissue distribution except erythrocytes. Over aa 21-720, human Integrin β 1 shares 95% aa identity with canine Integrin β 1.