

Purified anti-TIF1 β (KAP-1, TRIM28) Antibody

Catalog# / Size	941301 / 25 μ g 941302 / 100 μ g
Clone	W19027A
Regulatory Status	RUO
Other Names	Tripartite Motif Containing 28; RING Finger Protein 96; Transcription Intermediary Factor1- β , KRAB [Kruppel-Associated Box Domain]-Associated Protein 1; KRAB-Associated Protein 1
Isotype	Rat IgG2a, κ
Description	TIF β (transcription intermediary factor 1-beta) is an 89 kD member of the tripartite motif family. This protein contains three zinc binding domains, a RING domain, a B-box type 1 and type 2 domain, and a coiled-coil region. TIF β is found in the nucleus and associates with specific chromatin regions. This protein forms a complex with KRAB-domain transcription factors and recruits SETDB1 to histone 3 to increase KRAB-mediated transcriptional repression. TIF1 β has been reported to interact with SETDB1 and CBX3 proteins. Studies using knockout mice reveal the important function of TIF1 β in regulating genomic imprinting, T cell activation, and T cell tolerance.

Product Details

Verified Reactivity	Human, Mouse
Antibody Type	Monoclonal
Host Species	Rat
Immunogen	Full-length recombinant mouse TIF1 β protein
Formulation	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide
Preparation	The antibody was purified by affinity chromatography.
Concentration	0.5 mg/mL
Storage & Handling	The antibody solution should be stored undiluted between 2°C and 8°C.
Application	WB - Quality tested IP, ICC - Verified
Recommended Usage	Each lot of this antibody is quality control tested by western blotting. For western blotting, the suggested use of this reagent is 1.0 μ g/mL. For immunocytochemistry, a concentration range of 1.0 - 5.0 μ g/mL is recommended. For immunoprecipitation, the suggested use of this reagent is 2.5 μ g/test. It is recommended that the reagent be titrated for optimal performance for each application.
Application Notes	- BioLegend offers two monoclonal antibodies, clones 20A1 and W19027A that bind TIF1 β (KAP-1, TRIM28). <ul style="list-style-type: none">• 20A1 is a mouse IgG1, κ isotype; W19027A is a rat IgG2a, κ isotype• Both clones detect human and mouse TIF1β by western blot; 20A1 binds TIF1β with a higher affinity than W19027A for this application, but also shows a higher background• Both clones are validated for immunocytochemistry on human cells• W19027A is validated for immunoprecipitation, 20A1 has not been tested for this application <p>- Clone W19027A was tested for immunocytochemistry using 4% PFA-fixed HeLa cells permeabilized with either Triton X-100 or methanol. Both methods were compatible with TIF1β staining.</p>
RRID	AB_2888901 (BioLegend Cat. No. 941301) AB_2888901 (BioLegend Cat. No. 941302)

Antigen Details

Structure	TIF1 β is an 835 amino acid protein with a predicted molecular weight of 89 kD.
Distribution	Ubiquitously expressed/Nucleus
Function	Chromatin remodeling
Antigen References	<ol style="list-style-type: none">1. Ryan RF, <i>et al.</i> 1999. <i>Mol. Cell. Biol.</i> 19:4366.2. Schultz DC, <i>et al.</i> 2002. <i>Genes Dev.</i> 16:919.3. Moosmann PR, <i>et al.</i> 1996. <i>Nucleic Acids Res.</i> 24:4859.4. Friedman JR, <i>et al.</i> 1996. <i>Genes Dev.</i> 10:2067.5. Messerschmidt DM, <i>et al.</i> 2012. <i>Science</i> 335:1499.6. Chikuma S, <i>et al.</i> 2012. <i>Nat. Immunol.</i> 13:596.
Gene ID	10155

Related Protocols

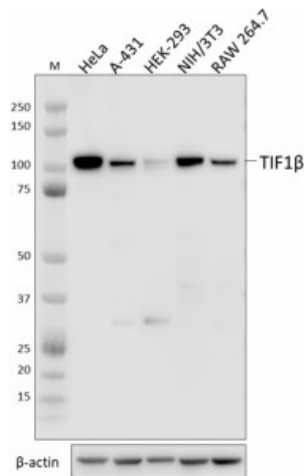
[Western Blotting Protocol](#)

[Immunoprecipitation Protocol](#)

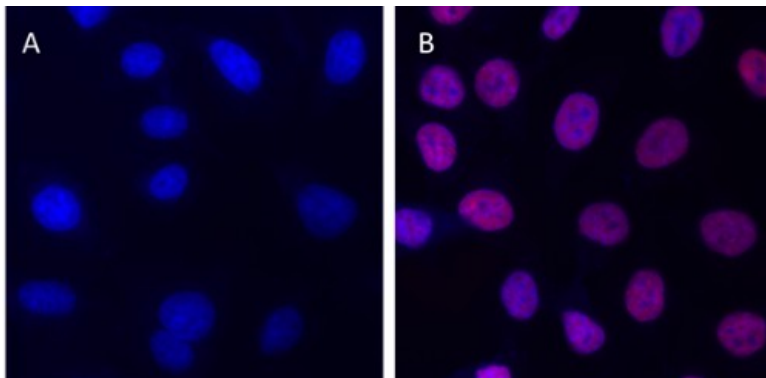
Other Formats

Purified anti-TIF1 β (KAP-1, TRIM28) Antibody

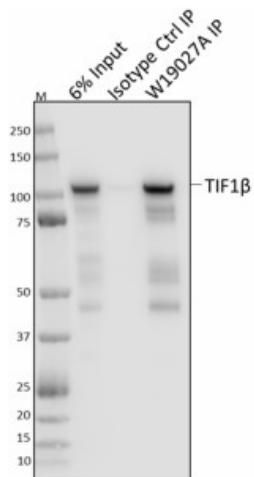
Product Data



Whole cell extracts (15 μ g total protein) from the indicated cell lines were resolved by 4-12% Bis-Tris gel electrophoresis, transferred to a PVDF membrane, and probed with 1.0 μ g/mL (1:500 dilution) purified anti-TIF1 β (KAP-1, TRIM28) antibody (clone W19027A) overnight at 4°C. Proteins were visualized by chemiluminescence detection using HRP goat anti-rat IgG antibody (Cat. No. 405405) at a 1:3000 dilution. Direct-Blot™ HRP anti- β -actin antibody (Cat. No. 664804) was used as a loading control at a 1:25000 dilution (lower). Lane M: Molecular weight marker.



HeLa cells were fixed with 4% paraformaldehyde for 10 minutes, permeabilized with methanol for 6 minutes, and blocked with 5% FBS for 60 minutes. Cells were then intracellularly stained with 5.0 μ g (1:100 dilution) purified rat IgG2a, κ isotype ctrl antibody (panel A) (Cat. No. 400501) or purified anti-TIF1 β (KAP-1, TRIM28) antibody (clone W19027A) (panel B) overnight at 4°C followed by incubation with Alexa Fluor® 594 goat anti-rat IgG antibody (Cat. No. 405422) at 2.0 μ g/mL. Nuclei were counterstained with DAPI and the image was captured with a 60X objective.



Whole cell extracts (250 µg total protein) prepared from HeLa cells were immunoprecipitated overnight with 2.5 µg purified rat IgG2a, κ isotype ctrl antibody (Cat. No. 400501) or purified anti-TIF1β (KAP-1, TRIM28) antibody (clone W19027A). The resulting IP fractions and whole cell extract input (6%) were resolved by 4-12% Bis-Tris gel electrophoresis, transferred to a PVDF membrane and probed with purified anti-TIF1β (KAP-1, TRIM28) antibody (clone 20A1) (Cat. No. 619301). Lane M: Molecular weight marker.

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