



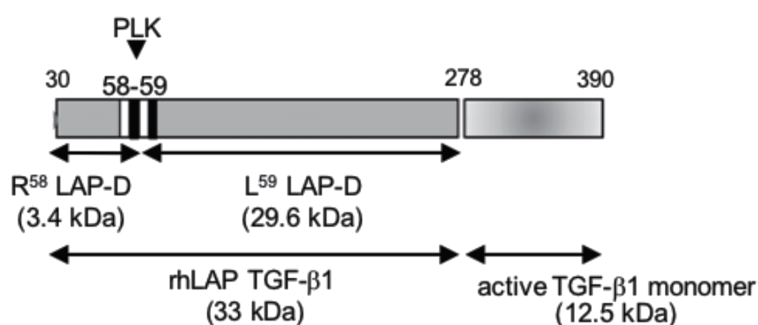
Anti TGF- β 1 LAP-D (L59)

(LAP Degrades N-Terminus side cut end L59)

Background :

TGF- β is produced as a latent form in which 25 kD active TGF- β is trapped by its pro-peptide called Latency Associated Protein (LAP). Upon receiving certain stimuli, a conformational change is induced in a latent complex to release the active TGF- β from the complex. The resultant TGF- β binds to cognate signaling receptors and exerts various physiological and pathological activities. This reaction is called TGF- β activation reaction, which is known to be induced by binding of the latent complex to cell adhesion proteins such as thrombospondin and integrins, and/or by being cleaved by the action of proteases such as serine proteases, cysteine proteases, and MMPs in an organ and context-depending manner.

Kojima and his colleagues in Cellular Molecular Pathology Research Unit (currently, Center for Integrative Medical Sciences, Liver Cancer Prevention Research Unit), RIKEN, Japan identified that a serine protease, plasma kallikrein induces release and activation of TGF- β by cleaving between 58Arg-59Leu within LAP and thereby participates in the pathogenesis of the liver diseases. The anti-TGF- β 1 LAP-degradates (LAP-D) antibodies are useful to investigate the molecular mechanism of TGF- β activation and its related diseases including liver fibrosis/cirrhosis and liver degeneration as tools to detect LAP-D.



Host Species:	Mouse
Form:	Liquid, PBS (pH 7.4), 0.05% NaN ₃
Volume:	100 μ g (1 mg/mL)
Specificity:	Recognizes N-terminus cut end of LAP degradates (LAP-D) L59 when latent TGF- β is digested with Plasma Kallikrein (PLK).
Antigen:	L59 peptide [LASPPSQGEVPGGC]
Clonality:	Monoclonal (clone # 6D6)
Isotype:	IgG1
Applications:	Western Blot: 2-10 μ g/mL ELISA (Enzyme-Linked ImmunoSorbent Assay): 20 μ g/mL * Optimal dilutions/concentrations should be determined by each researcher.
Purification method:	Purified from cell culture of serum-free medium by affinity column (Protein G)
Conjugation:	none
Storage condition:	Store below -20°C (below -70°C for prolonged storage) *Aliquot to avoid cycles of freeze/thaw

* Anti TGF- β 1 LAP-D (L59) was generated & licensed under RIKEN, Japan.

References:

1. LAP degradation product reflects plasma kallikrein-dependent TGF- β activation in patients with hepatic fibrosis, Hara M., Kirita A., Kondo W. et al. Springerplus. May 1; 3: 221. PMCID: PMC4033717 (2014)
2. L⁵⁹ TGF- β LAP degradation products serve as a promising blood biomarker for liver fibrogenesis in mice, Hara M., Inoue I., Yamazaki Y. et al. Fibrogenesis Tissue Repair. Sep 15; 8: 17. PMCID: PMC4570586 (2015)

Example Assay Data:

1. Western Blotting

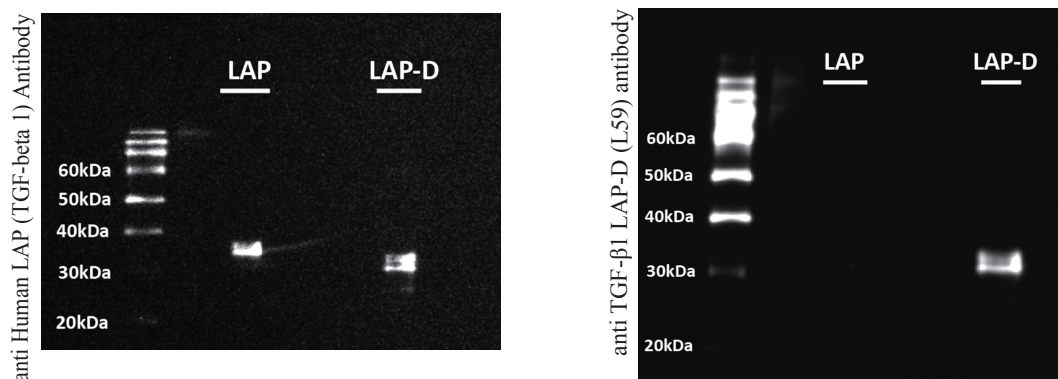


Figure 1. Western Blotting result with anti TGF- β 1 LAP-D (L59) antibody

[sample] uncut human recombinant LAP (R&D Systems, 246-LP-025) (LAP), PLK digested human recombinant LAP (LAP degradates, LAP-D) [10 ng/lane]

Antibody:

(left figure) anti Human LAP (TGF-beta 1) Antibody (R&D Systems, AF-246-NA, 0.1 μ g/mL), Peroxidase AffiniPure F(ab')₂ Fragment Rabbit Anti-Goat IgG (H+L) (Jackson ImmunoResearch Laboratories, 305-036-045, 0.08 μ g/mL)

(right figure) anti TGF- β 1 LAP-D (L59) antibody (10 μ g/mL), Peroxidase AffiniPure F(ab')₂ Fragment Goat Anti-Mouse IgG (H+L) (Jackson ImmunoResearch Laboratories, 115-036-062, 0.08 μ g/mL)

Detection: Pierce® ECL Plus Western Blotting Substrate (ThermoFischer Scientific, NCI32132JP)

* anti TGF- β 1 LAP-D (L59) antibody specificity has been confirmed by detection of LAP-D [31 kDa]

2. ELISA (Enzyme-Linked ImmunoSorbent Assay)

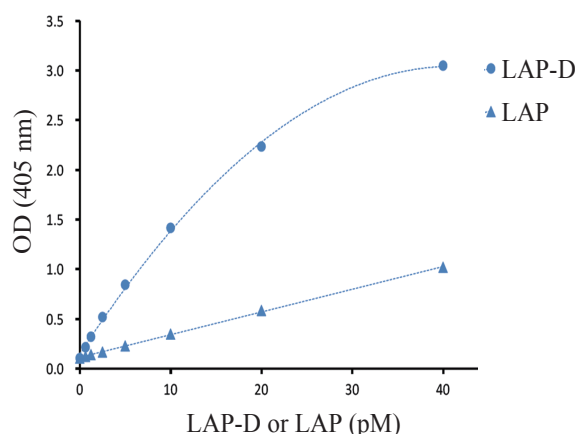


Figure 2. Sandwich ELISA with anti TGF- β 1 LAP-D (L59) antibody and anti Human LAP TGF-beta 1 Biotinylated antibody

[sample] PLK digested human recombinant LAP (L59 LAP degradates, LAP-D), uncut human recombinant LAP (LAP)

Coating Antibody: anti TGF- β 1 LAP-D (L59) antibody [20 μ g/mL]

Detection Antibody: anti Human LAP TGF-beta 1 Biotinylated Antibody (R&D Systems, BAM2462, 0.5 μ g/mL)

Detection: Streptavidin-AP (Jackson ImmunoResearch Laboratories, 016-050-084, 0.05 μ g/mL)

* This ELISA system detected uncut human recombinant LAP, which showed c.a. 1 OD at 40 pM, while LAP-D showed c.a. 3 OD at 40 pM, giving significant differences. This result proves the specificity of anti TGF- β 1 LAP-D (L59) antibody.

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