# Pan Lactic acid-Lysine Rabbit pAb

Catalog No.: A18831 1 Publications



### **Basic Information**

**Observed MW** 

**Calculated MW** 

Category

Primary antibody

**Applications** 

WB,ELISA

**Cross-Reactivity** 

Human, Mouse, Rat, Other (Wide Range)

# **Background**

Histone lysine lactation (Kla) is a newly discovered histone modification that regulates gene expression in macrophages. In M1 macrophages, lactic acid is derived from incompletely oxidized glucose and then produces lactyl-CoA, which is transferred via acetyltransferase p300 to the lysine tail of the histone. This modification is abundant in gene promoter regions that lack acetylation and are associated with gene expression activation.

# **Recommended Dilutions**

**WB** 1:500 - 1:1000

**ELISA** 

Recommended starting concentration is 1 µg/mL.
Please optimize the concentration based on your specific assay requirements.

# **Immunogen Information**

Gene ID Swiss Prot

#### **Immunogen**

A synthetic peptide corresponding to a sequence containing Pan Lactic acid-Lysine.

#### **Synonyms**

# **Contact**

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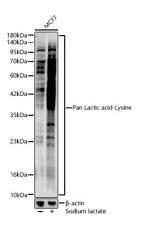
# **Product Information**

SourceIsotypePurificationRabbitIgGAffinity purification

# Storage

Store at -20°C. Avoid freeze / thaw cycles.

Buffer: PBS with 0.09% sodium azide,50% glycerol,pH7.3.



Western blot analysis of lysates from MCF7 cells, using Pan Lactic acid-Lysine Rabbit pAb (A18831) at 1:400 dilution. MCF7 cells were treated by Sodium lactate(100mM) for 24h.

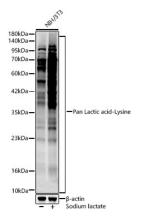
Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

Lysates/proteins: 25µg per lane.

Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Enhanced Kit (RM00021).

Exposure time: 60s.



Western blot analysis of lysates from NIH/3T3 cells, using Pan Lactic acid-Lysine Rabbit pAb (A18831) at 1:400 dilution. NIH/3T3 cells were treated by Sodium lactate(100 mM) for 24 h.

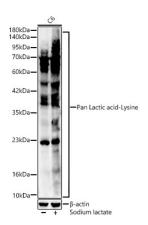
Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

Lysates/proteins: 25µg per lane.

Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Enhanced Kit (RM00021).

Exposure time: 60s.



Western blot analysis of lysates from C6 cells, using Pan Lactic acid-Lysine Rabbit pAb (A18831) at 1:400 dilution. C6 cells were treated by Sodium lactate(100mM) for 24h.

Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

Lysates/proteins: 25µg per lane.

Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Enhanced Kit (RM00021).

Exposure time: 60s.