



## AL (11A1): sc-59944

### BACKGROUND

AL is an altered amino acid, specifically a N-epsilon acetyl lysine. AL catalyzes the reversible transamination of hydrophobic D-amino acids with pyridoxal 5'-phosphate (PLP). Acetylation of specific lysine residues in the N-terminal domains of core histones is a biochemical marker of active genes, and thus antibodies to these lysines prove to be extremely useful as a research tool. AL can also be used as a lysine derivative used to treat individuals with a lysine deficiency which may lead to epithelial diamino acid transport in lysinuric protein intolerance (LPI), a defect that results from an abnormality of the basolateral cell membranes.

### REFERENCES

1. Rajantie, J., Simell, O. and Perheentupa, J. 1983. Oral administration of epsilon N-acetyllysine and homocitrulline in lysinuric protein intolerance. *J. Pediatr.* 102: 388-390.
2. Violand, B.N., Schlittler, M.R., Lawson, C.O., Kane, J.F., Siegel, N.R., Smith, C.E., Kolodziej, E.W. and Duffin, K.L. 1994. Isolation of *Escherichia coli* synthesized recontain  $\epsilon$ -N-acetyllysine. *Protein Sci.* 3: 1089-1097.
3. Crane-Robinson, C., Hebbes, T.R., Clayton, A.L. and Thorne, A.W. 1997. Chromosomal mapping of core histone acetylation by immunoselection. *Methods* 12: 48-56.
4. Hudson, B.P., Martinez-Yamout, M.A., Dyson, H.J. and Wright, P.E. 2000. Solution structure and acetyl-lysine binding activity of the GCN5 bromodomain. *J. Mol. Biol.* 304: 355-370.
5. Schultz, B.E., Misialek, S., Wu, J., Tang, J., Conn, M.T., Tahilramani, R. and Wong, L. 2004. Kinetics and comparative reactivity of human class I and class IIb histone deacetylases. *Biochemistry* 43: 11083-11091.
6. Fatkins, D.G., Monnot, A.D. and Zheng, W. 2006. Nepsilon-thioacetyl-lysine: a multi-facet functional probe for enzymatic protein lysine Nepsilon-deacetylation. *Bioorg. Med. Chem. Lett.* 16: 3651-3656.
7. Golinelli-Pimpaneau, B., Lüthi, C. and Christen, P. 2006. Structural basis for D-amino acid transamination by the pyridoxal 5'-phosphate-dependent catalytic antibody 15A9. *J. Biol. Chem.* 281: 23969-23977.

### SOURCE

AL (11A1) is a mouse monoclonal antibody.

### PRODUCT

Each vial contains 100  $\mu$ l ascites containing IgG<sub>1</sub> with < 0.1% sodium azide.

### STORAGE

For immediate and continuous use, store at 4° C for up to one month. For sporadic use, freeze in working aliquots in order to avoid repeated freeze/thaw cycles. If turbidity is evident upon prolonged storage, clarify solution by centrifugation.

### PROTOCOLS

See our web site at [www.scbt.com](http://www.scbt.com) or our catalog for detailed protocols and support products.

### APPLICATIONS

AL (11A1) is recommended for detection of the N $\alpha$ -acetyl lysine (AL), N-acetyl putrescine and acetylated histones by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunofluorescence and immunohistochemistry (including paraffin-embedded sections) (starting dilution 1:50, dilution range 1:50-1:500); non cross-reactive with either propionyl lysine or butyryl lysine.

### RECOMMENDED SECONDARY REAGENTS

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use goat anti-mouse IgG-HRP: sc-2005 (dilution range: 1:2000-1:32,000) or Cruz Marker™ compatible goat anti-mouse IgG-HRP: sc-2031 (dilution range: 1:2000-1:5000), Cruz Marker™ Molecular Weight Standards: sc-2035, TBS Blotto A Blocking Reagent: sc-2333 and Western Blotting Luminol Reagent: sc-2048. 2) Immunofluorescence: use goat anti-mouse IgG-FITC: sc-2010 (dilution range: 1:100-1:400) or goat anti-mouse IgG-TR: sc-2781 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941. 3) Immunohistochemistry: use ImmunoCruz™: sc-2050 or ABC: sc-2017 mouse IgG Staining Systems.

### RESEARCH USE

For research use only, not for use in diagnostic procedures.