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## 제 ATP-Citrate Lyase (D1X6P) Rabbit mAb



| Orders:  | 877-616-CELL (2355)<br>orders@cellsignal.com |
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3 Trask Lane | Danvers | Massachusetts | 01923 | USA

For Research Use Only. Not for Use in Diagnostic Procedures.

| Applications:<br>W           | <b>Reactivity:</b><br>H M R | <b>Sensitivity:</b><br>Endogenous   | <b>MW (kDa):</b><br>125  | <b>Source/Isotype:</b><br>Rabbit IgG   | UniProt ID:<br>#P53396   | Entrez-Gene Id:<br>47  |  |  |
|------------------------------|-----------------------------|---|--|--|--|--|--|--|
| Product Usage<br>Information | 9                           | Application<br>Western Blotting   |  |  | Dilution<br>1:1000   |  |  |  |
| Storage                      |                             | Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.  |  |  |  |  |  |  |
|                              |                             | For a carrier free (BSA and azide free) version of this product see product #51077.   |  |  |  |  |  |  |
| Specificity/Ser              | nsitivity                   | ATP-Citrate Lyase (D1X6P) Rabbit mAb recognizes endogenous levels of total ATP-citrate lyase p  |  |  |  |  |  |  |
| Source / Purifi              | cation                      | Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Val34 of human ATP citrate lyase protein.  |  |  |  |  |  |  |
| Background                   |                             | ATP-citrate lyase (ACL) is a homotetramer that catalyzes the formation of acetyl-CoA and oxaloacetate<br>(OAA) in the cytosol, which is the key step for the biosynthesis of fatty acids, cholesterol, and<br>acetylcholine, as well as for gluconeogenesis (1). Nutrients and hormones regulate the expression level<br>and phosphorylation of ATP-citrate lyase (1,2). It is phosphorylated by GSK-3 on Thr446 and Ser450 (3).<br>Ser455 of ATP-citrate lyase has been reported to be phosphorylated by PKA and Akt (4,5).<br>Phosphorylation on Ser455 abolishes the homotropic allosteric regulation by citrate and enhances the<br>catalytic activity of the enzyme (2). |  |  |  |  |  |  |
| Background R                 | eferences                   | 1. Towle, H.C. et al. (19<br>2. Potapova, I.A. et al.<br>3. Hughes, K. et al. (19<br>4. Pierce, M.W. et al. (1<br>5. Berwick, D.C. et al. (1  | (2000) <i>Biochemisti</i><br>92) <i>Biochem J</i> 288<br>982) <i>J Biol Chem</i> 2           | y 39, 1169-79.<br>( Pt 1), 309-14.<br>57, 10681-6.   |  |  |  |  |
| Species Reacti               | vity                        | Species reactivity is determined by testing in at least one approved application (e.g., western blot).  |  |  |  |  |  |  |
| Western Blot B               | Buffer                      | IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v nonfat<br>dry milk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.   |  |  |  |  |  |  |
| Applications K               | ey                          | W: Western Blotting   |  |  |  |  |  |  |
| Cross-Reactivi               | ty Key                      | H: Human M: Mouse R: Rat  |  |  |  |  |  |  |
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