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#5277

Phospho-LCP1 (Tyr28) Antibody

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

| | | | | | | |
|---------------------------|-------------------------|-----------------------------------|------------------------|----------------------------------|-------------------------------|--------------------------------|
| Applications: W | Reactivity: H | Sensitivity: Endogenous | MW (kDa): 70 | Source/Isotype: Rabbit | UniProt ID: #P13796 | Entrez-Gene Id: 3936 |
|---------------------------|-------------------------|-----------------------------------|------------------------|----------------------------------|-------------------------------|--------------------------------|

Product Usage Information

Application

Western Blotting

Dilution

1:1000

Storage

Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA and 50% glycerol. Store at -20°C. Do not aliquot the antibody.

Specificity/Sensitivity

Phospho-LCP1 (Tyr28) Antibody detects endogenous levels of LCP1 protein only when phosphorylated on Tyr28.

Source / Purification

Polyclonal antibodies are produced by immunizing animals with a synthetic phosphopeptide corresponding to residues surrounding Tyr28 of human LCP1 protein. Antibodies are purified by protein A and peptide affinity chromatography.

Background

Highly conserved and widely expressed platin proteins comprise a subset of actin-binding proteins that include proteins that promote actin bundling. Three platins exhibiting differential expression are found in mammals and include L-platin, T-platin, and I-platin. T-platin (platin-3) is found in cells of most solid tissues, while I-platin (platin-1) is expressed specifically in the kidney, colon, and small intestine (1-3). Research studies have shown that L-platin (platin-2) or lymphocyte cytosolic protein 1 (LCP1) is mainly expressed in hematopoietic cells and nonhematopoietic tumors, and increased expression correlates with metastatic progression in colon cancer cell lines (4). Investigators have found that overexpression of LCP1 in premetastatic cancer cell lines induces invasion and loss of E-cadherin expression, which is characteristic of metastatic cancer cell lines (5). LCP1 becomes phosphorylated at Ser5 upon stimulation through the T cell receptor/CD3 complex in association with the CD2 cell adhesion molecule or the CD28 receptor (6). Phosphorylation at Ser5 enhances the ability of LCP1 to bind to F-actin and increases cell motility (7,8). Phosphorylation of LCP1 on Tyr28 was identified at Cell Signaling Technology (CST) using PhosphoScan[®], CST's LC-MS/MS platform for phosphorylation site discovery as well as other publications using MS technology (9). Phosphorylation of LCP1 at Tyr28 is seen in many leukemic cell lines (9-12).

Background References

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- Otsuka, M. et al. (2001) *Biochem Biophys Res Commun* 289, 876-81.
- Foran, E. et al. (2006) *Int J Cancer* 118, 2098-104.
- Wabnitz, G.H. et al. (2007) *Eur J Immunol* 37, 649-62.
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- Klemke, M. et al. (2007) *Int J Cancer* 120, 2590-9.
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- Rikova, K. et al. (2007) *Cell* 131, 1190-203.
- Gu, T.L. et al. (2007) *Blood* 110, 323-33.
- Walters, D.K. et al. (2006) *Leuk Res* 30, 1097-104.

Species Reactivity

Species reactivity is determined by testing in at least one approved application (e.g., western blot).

Western Blot Buffer

IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

Applications Key

W: Western Blotting

Cross-Reactivity Key

H: Human

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