#3341 Store at -20C

Phospho-ALK (Tyr1604) Antibody



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

Applications: WB, W-S, IP	Reactivity: H	Sensitivity: Endogenous	MW (kDa): 80 (NPM-ALK) 220 (ALK)	Source: Rabbit	UniProt ID: #Q9UM73	Entrez-Gene Id 238	
Product Usage Information	Application			Dilution			
	We	Western Blotting			1:1000		
	Sin	Simple Western™			1:10 - 1:50		
	lmi	Immunoprecipitation			1:50		
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μ g/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.					
Specificity / Sensitiv	,	Phospho-ALK (Tyr1604) Antibody detects ALK only when phosphorylated at Tyr16 of NPM-ALK). This antibody may cross-react with other activated protein tyrosine					
Source / Purification	to re	Polyclonal antibodies are produced by immunizing animals with a synthetic phosphopeptide corresponding to residues surrounding Tyr1604 of human ALK. Antibodies are purified by protein A and peptide affinity chromatography					
Background	invo both	Anaplastic lymphoma kinase (ALK) is a tyrosine kinase receptor for pleiotrophin (PTN), a growth factor involved in embryonic brain development (1-3). In ALK-expressing cells, PTN induces phosphorylation of both ALK and the downstream effectors IRS-1, Shc, PLCy, and PI3 kinase (1). ALK was originally discovered as a nucleophosmin (NPM)-ALK fusion protein produced by a translocation (4). Investigators					

involved in embryonic brain development (1-3). In ALK-expressing cells, PTN induces phosphorylation of both ALK and the downstream effectors IRS-1, Shc, PLCy, and PI3 kinase (1). ALK was originally discovered as a nucleophosmin (NPM)-ALK fusion protein produced by a translocation (4). Investigators have found that the NPM-ALK fusion protein is a constitutively active, oncogenic tyrosine kinase associated with anaplastic lymphoma (4). Research literature suggests that activation of PLCy by NPM-ALK may be a crucial step for its mitogenic activity and involved in the pathogenesis of anaplastic lymphomas (5).

A distinct ALK oncogenic fusion protein involving ALK and echinoderm microtubule-associated protein like 4 (EML4) has been described in the research literature from a non-small cell lung cancer (NSCLC) cell line, with corresponding fusion transcripts present in some cases of lung adenocarcinoma. The short, amino-terminal region of the microtubule-associated protein EML4 is fused to the kinase domain of ALK (6-8).

Phosphorylated Tyr664 of NPM-ALK (equivalent to Tyr1604 of full length ALK) is required for the interaction with PLCgamma (5). Site-directed mutagenesis of this tyrosine residue results in the loss of oncogenic activity of NPM-ALK (5).

Background References

- 1. Stoica, G.E. et al. (2001) J Biol Chem 276, 16772-9.
- 2. Iwahara, T. et al. (1997) Oncogene 14, 439-49.
- 3. Morris, S.W. et al. (1997) *Oncogene* 14, 2175-88.
- 4. Morris, S.W. et al. (1994) Science 263, 1281-4.
- 5. Bai, R.Y. et al. (1998) *Mol Cell Biol* 18, 6951-61.
- 6. Rikova, K. et al. (2007) *Cell* 131, 1190-203.
- 7. Takeuchi, K. et al. (2008) Clin Cancer Res 14, 6618-24.
- 8. Soda, M. et al. (2007) Nature 448, 561-6.

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

Cross-Reactivity Key

WB: Western Blotting W-S: Simple Western™ IP: Immunoprecipitation

Phospho-ALK (Tyr1604) Antibody (#3341) Datasheet Without Images Cell Signaling Technology

H: human M: mouse R: rat Hm: hamster Mk: monkey Vir: virus Mi: mink C: chicken Dm: D. melanogaster X: Xenopus Z: zebrafish B: bovine Dg: dog Pg: pig Sc: S. cerevisiae Ce: C. elegans Hr: horse GP: Guinea Pig Rab: rabbit All: all species expected

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