

## ALK (D5F3<sup>®</sup>) XP<sup>®</sup> Rabbit mAb (HRP Conjugate)



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

Applications:	Reactivity:	Sensitivity:	MW (kDa):	Source/Isotype:	UniProt ID:	Entrez-Gene Id:
W	Н	Endogenous	220 (ALK), 80 (NPM-	Rabbit IgG	#Q9UM73	238
			ALK), 117 (EML4-			
			ALK v1), 86 (EML4-			
			ALK v3)			

Product Usage<br/>InformationApplication<br/>Western BlottingDilution<br/>1:1000

Storage Supplied in 136 mM NaCl, 2.6 mM KCl, 12 mM sodium phosphate (pH 7.4) dibasic, 2 mg/ml BSA, and

50% glycerol. Store at -20°C. Do not aliquot the antibody.

Specificity/Sensitivity ALK (D5F3®) XP® Rabbit mAb (HRP Conjugate) detects endogenous levels of total ALK protein as well as

ALK fusion proteins, such as EML4-ALK variants and NPM-ALK, even at low levels. This antibody does

not cross-react with other family members.

**Source / Purification** Monoclonal antibody is produced by immunizing animals with recombinant protein corresponding to

residues in the carboxy terminus of human ALK.

**Description** This Cell Signaling Technology antibody is conjugated to the carbohydrate groups of horseradish

peroxidase (HRP) via its amine groups. The HRP conjugated antibody is expected to exhibit the same

species cross-reactivity as the unconjugated ALK (D5F3®) XP® Rabbit mAb #3633.

Background

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

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 PLCγ 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).

**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 nonfat

dry milk, 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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