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# Fes Antibody

Store at -20C  
#2736

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

<b>Applications:</b> W, IP	<b>Reactivity:</b> H	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 93	<b>Source/Isotype:</b> Rabbit	<b>UniProt ID:</b> #P07332	<b>Entrez-Gene Id:</b> 2242
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## Product Usage Information

### Application

Western Blotting  
Immunoprecipitation

### Dilution

1:1000  
1:50

## 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

Fes Antibody detects endogenous levels of Fes proteins. This antibody does not cross-react with other proteins.

## Source / Purification

Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Thr421 of human Fes. Antibodies are purified by protein A and peptide affinity chromatography.

## Background

Fes/Fps and Fer are the only two members of a unique family of cytoplasmic protein tyrosine kinases (1,2). Fes and Fer contain a central Src homology-2 (SH2) domain and a carboxy-terminal tyrosine kinase catalytic domain. They are structurally distinguished from other members of cytoplasmic protein tyrosine kinase subfamilies by the presence of amino-terminal Fer/CIP4 homology and coiled-coil domains (3). Fes/Fps was originally identified as an oncogene from avian (Fps) and feline (Fes) retroviruses. Human c-Fes has been implicated in myeloid, vascular endothelial and neuronal cell differentiation. Mutations may activate the Fps kinase and thereby contribute to cancer (4). However, recent data strongly suggests that the c-Fes protein-tyrosine kinase is a tumor suppressor rather than a dominant oncogene in colorectal cancer (5).

## Background References

1. Smithgall, T.E. et al. (1998) *Crit Rev Oncog* 9, 43-62.
2. Greer, P. (2002) *Nat Rev Mol Cell Biol* 3, 278-89.
3. Sangrar, W. et al. (2005) *Cancer Res* 65, 3518-22.
4. Ley, T.J. et al. (2003) *Proc Natl Acad Sci U S A* 100, 14275-80.
5. Delfino, F.J. et al. (2006) *J Biol Chem* 281, 8829-35.

## 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 **IP:** Immunoprecipitation

## Cross-Reactivity Key

**H:** Human

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