

Product datasheet for **SC205145**

ZAP70 (NM_207519) Human 3' UTR Clone

Product data:

Product Type:	3' UTR Clones
Product Name:	ZAP70 (NM_207519) Human 3' UTR Clone
Symbol:	ZAP70
Synonyms:	ADMIO2; IMD48; SRK; STCD; STD; TZK; ZAP-70
Mammalian Cell Selection:	Neomycin
Vector:	pMirTarget (PS100062)
ACCN:	NM_207519
Insert Size:	392 bp
Insert Sequence:	>SC205145 3'UTR clone of NM_207519 The sequence shown below is from the reference sequence of NM_207519. The complete sequence of this clone may contain minor differences, such as SNPs. Blue =Stop Codon Red =Cloning site GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAAGCCAAGAAGGGCGGAAAGATCGCCGTG TAACAATTGGCAGAGCTCAGAATTCAA GCGATCGCC ACACAGAAAGGCTGAGGCTGCCTGTGCC TGA GCTCCCCTGCCAGGGGAGCCCTCCACGCCGGCTCTTC CCCACCCTCAGCCCCACCCAGGTCCTGCAGTCTGGCTGAGCCCTGCTTGGTTGTCTCCACACACAGCT GGGCTGTGGTAGGGGTGTCTCAGGCCACACCGCCTTGCAATTGCCTGGCCCCCTGCCTCTCTG GCTGGGAGCAGGGAGGTCGGGAGGGTCCGGCTGTGCAGCCTGTCTGGGCTGGTGGCTCCCGGAGGG CCCTGAGCTGAGGCATTGCTTACACGGATGCCTTCCCCTGGGCCCTGACATTGGAGCCTGGGCATCCT CAGGTGGTCAGGCGTAGATCACCAGAATAAACCCAGCTTCCCTCTTG ACGCGT AAGCGGCCGCGCATCTAGATTCAAGAAAAATGACCGACCAAGCGACGCCAACCTGCCATCA CGAGATTCGATTCCACCGCCGCTTCTATGAAAGG
Restriction Sites:	Sgfl-MluI
OTI Disclaimer:	Our molecular clone sequence data has been matched to the sequence identifier above as a point of reference. Note that the complete sequence of this clone is largely the same as the reference sequence but may contain minor differences, e.g., single nucleotide polymorphisms (SNPs).
Components:	The cDNA clone is shipped in a 2-D bar-coded Matrix tube as 10 ug dried plasmid DNA. The package also includes 100 pmols of both the corresponding 5' and 3' vector primers in separate vials.



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RefSeq: [NM_207519.2](#)

Summary: This gene encodes an enzyme belonging to the protein tyrosine kinase family, and it plays a role in T-cell development and lymphocyte activation. This enzyme, which is phosphorylated on tyrosine residues upon T-cell antigen receptor (TCR) stimulation, functions in the initial step of TCR-mediated signal transduction in combination with the Src family kinases, Lck and Fyn. This enzyme is also essential for thymocyte development. Mutations in this gene cause selective T-cell defect, a severe combined immunodeficiency disease characterized by a selective absence of CD8-positive T-cells. Two transcript variants that encode different isoforms have been found for this gene. [provided by RefSeq, Jul 2008]

Locus ID: 7535

MW: 13.1