

ZAP-70 (1E7.2): sc-32760



The Power to Question

BACKGROUND

The activation of T lymphocytes by antigens is mediated by the T cell receptor (TCR) which is a multisubunit complex assembled from at least six different genes. The TCR subunits include the $Ti\alpha$ and β chains, the $CD3\gamma$, δ and ϵ chains and a ζ -containing homodimer or heterodimer. The disulfide-linked $Ti\alpha$ - β heterodimer is responsible for antigen recognition, but the short five amino acid cytoplasmic domains of $Ti\alpha$ and β are unlikely to be sufficient to couple to intracellular signaling pathways. In contrast, the structured features of the $CD3$ and ζ subunits suggest a role in signal transduction. Of these, the ζ chain, which is expressed as either a homodimer or heterodimer, has a short extracellular domain of only nine amino acids, but a larger 113 amino acid cytoplasmic domain. A tyrosine phosphoprotein, ZAP-70, has been identified that associates with ζ and undergoes tyrosine phosphorylation following TCR stimulation.

CHROMOSOMAL LOCATION

Genetic locus: ZAP70 (human) mapping to 2q11.2; Zap70 (mouse) mapping to 1B.

SOURCE

ZAP-70 (1E7.2) is a mouse monoclonal antibody raised against amino acids 282-307 of ZAP-70 of human origin.

PRODUCT

Each vial contains 200 μ g IgG₁ kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

ZAP-70 (1E7.2) is available conjugated to agarose (sc-32760 AC), 500 μ g/0.25 ml agarose in 1 ml, for IP; to HRP (sc-32760 HRP), 200 μ g/ml, for WB, IHC(P) and ELISA; to either phycoerythrin (sc-32760 PE), fluorescein (sc-32760 FITC), Alexa Fluor® 488 (sc-32760 AF488), Alexa Fluor® 546 (sc-32760 AF546), Alexa Fluor® 594 (sc-32760 AF594) or Alexa Fluor® 647 (sc-32760 AF647), 200 μ g/ml, for WB (RGB), IF, IHC(P) and FCM; and to either Alexa Fluor® 680 (sc-32760 AF680) or Alexa Fluor® 790 (sc-32760 AF790), 200 μ g/ml, for Near-Infrared (NIR) WB, IF and FCM.

Alexa Fluor® is a trademark of Molecular Probes, Inc., Oregon, USA

APPLICATIONS

ZAP-70 (1E7.2) is recommended for detection of ZAP-70 of mouse, rat and human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2 μ g per 100-500 μ g of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and flow cytometry (1 μ g per 1×10^6 cells).

Suitable for use as control antibody for ZAP-70 siRNA (h): sc-29526, ZAP-70 siRNA (m): sc-36867, ZAP-70 shRNA Plasmid (h): sc-29526-SH, ZAP-70 shRNA Plasmid (m): sc-36867-SH, ZAP-70 shRNA (h) Lentiviral Particles: sc-29526-V and ZAP-70 shRNA (m) Lentiviral Particles: sc-36867-V.

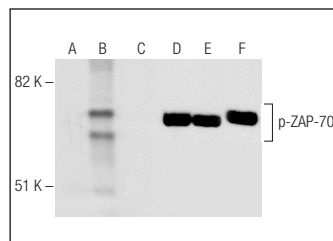
Molecular Weight of ZAP-70: 70 kDa.

Positive Controls: BYDP whole cell lysate: sc-364368, Jurkat whole cell lysate: sc-2204 or CCRF-CEM cell lysate: sc-2225.

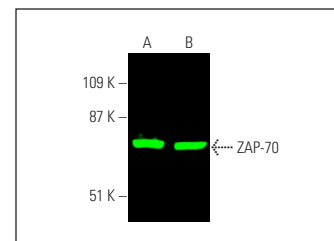
STORAGE

Store at 4° C, ****DO NOT FREEZE****. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

DATA



Western blot analysis of ZAP-70 phosphorylation in untreated (A,D), pervanadate treated (B,E) and pervanadate and lambda protein phosphatase (sc-200312A) treated (C,F) Jurkat whole cell lysates. Antibodies tested include p-ZAP-70 (Tyr 493): sc-101823 (A,B,C) and ZAP-70 (1E7.2): sc-32760 (D,E,F).



ZAP-70 (1E7.2): sc-32760. Near-infrared western blot analysis of ZAP-70 expression in CCRF-CEM (A) and BYDP (B) whole cell lysates. Blocked with UltraCruz® Blocking Reagent: sc-516214. Detection reagent used: m-IgGκ BP-CFL 680: sc-516180.

SELECT PRODUCT CITATIONS

1. Fallone, C.A. 2000. Epidemiology of the antibiotic resistance of *Helicobacter pylori* in Canada. *Can. J. Gastroenterol.* 14: 879-882.
2. Chen, C., et al. 2007. c-Abl is required for the signaling transduction induced by L-selectin ligation. *Eur. J. Immunol.* 37: 3246-3258.
3. Shim, J.H., et al. 2008. (-)-Epigallocatechin gallate regulates CD3-mediated T cell receptor signaling in leukemia through the inhibition of ZAP-70 kinase. *J. Biol. Chem.* 283: 28370-28379.
4. Leavenworth, J.W. and Pauza, M.E. 2009. Engagement of transgenic Ly49A inhibits mouse CD4 cell activation by disrupting T cell receptor, but not CD28, signaling. *Cell. Immunol.* 257: 88-96.
5. Tsagaratou, A., et al. 2010. Thymocyte-specific truncation of the deubiquitinating domain of CYLD impairs positive selection in a NFκB essential modulator-dependent manner. *J. Immunol.* 185: 2032-2043.
6. San Luis, B., et al. 2011. Sts-2 is a phosphatase that negatively regulates ζ -associated protein ZAP-70 and T cell receptor signaling pathways. *J. Biol. Chem.* 286: 15943-15954.
7. Markwart, R., et al. 2014. Immunosuppression after sepsis: systemic inflammation and sepsis induce a loss of naïve T-cells but no enduring cell-autonomous defects in T-cell function. *PLoS ONE* 9: e115094.
8. Zou, Q., et al. 2015. T cell development involves TRAF3IP3-mediated ERK signaling in the Golgi. *J. Exp. Med.* 212: 1323-1336.
9. Moogk, D., et al. 2016. Constitutive Ick activity drives sensitivity differences between CD8⁺ memory T-cell subsets. *J. Immunol.* 197: 644-654.
10. Geibler, K., et al. 2017. Functional characterization of T-cells from palatine tonsils in patients with chronic tonsillitis. *PLoS ONE* 12: e0183214.

RESEARCH USE

For research use only, not for use in diagnostic procedures.