Supporting Information

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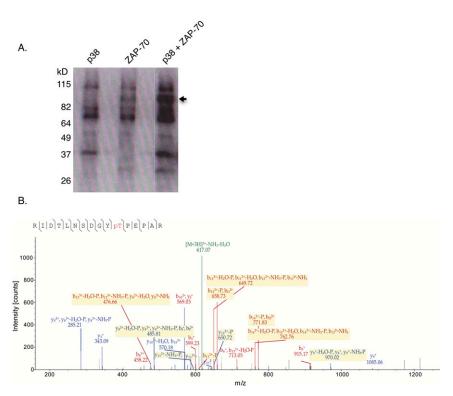


Fig. S1. (*A*) Human recombinant GST–ZAP-70 was incubated with or without murine His-tagged p38 α alone in kinase buffer containing [γ -³²P]-ATP. After electrophoresis the radioactive bands were visualized using a Phosphorimager. The arrow indicates the ~95-kDa band corresponding to GST–ZAP-70. (*B*) Tandem mass spectrum of the peptide RIDTLNSDGYTPEPAR from GST–ZAP-70, corresponding to residues 283–298. The identified b (red) and y (blue) ions are denoted in the spectrum; unfragmented precursor peptide is shown in green. The ions showing a loss of phosphate are highlighted in yellow. The peptide sequence and matched ions are provided above the spectrum.

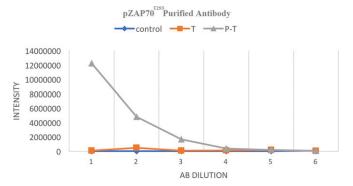


Fig. S2. Serial dilutions of the phospho-peptide-purified anti-pZAP-70^{T293} polyclonal rabbit antibody were tested against unphosphorylated or the phosphorylated peptide for reactivity by ELISA. The wells were coated with: control, sodium carbonate buffer; T, unphosphorylated peptide; P-T, immunizing phospho-peptide, NSDGY-pT-PEPAC.