

# PRODUCT DATA SHEET

Bioworld Technology, Inc.

# ZAP70 (phospho-Y315) polyclonal antibody

Catalog: BS4884 Host: Rabbit Reactivity: Human, Mouse, Rat

#### **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 a and b chains, the CD3 g, d and e chains and a z -containing homodimer or heterodimer . The protein tyrosine kinase ZAP-70 binds to the phosphorylated immunoreceptor tyrosine-base activation motifs (ITAMs) of the TCR z chain through two src-homology (SH2) domains. This binding results in the phosphorylation of ZAP-70 on multiple tyrosine residues, including Tyr292 and Tyr319. ZAP-70 is autophosphorylated on Tyr292, which is thought to negatively regulate ZAP-70 function in lymphocytes . Alternatively, ZAP-70 is positively regulated by phosphorylation on Tyr319, which mediates the SH2-dependent interaction between Lck and ZAP-70.

### **Product:**

Rabbit IgG, 1mg/ml in PBS with 0.02% sodium azide, 50% glycerol, pH7.2

#### **Molecular Weight:**

~ 70 kDa

#### **Swiss-Prot:**

P43403

## **Purification&Purity:**

The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen and the purity is > 95% (by SDS-PAGE).

# **Applications:**

WB: 1:500~1:1000 IHC: 1:50~1:200

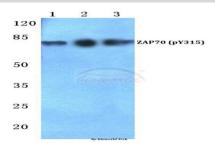
# Storage&Stability:

Store at  $4\,\mathrm{C}$  short term. Aliquot and store at  $-20\,\mathrm{C}$  long term. Avoid freeze-thaw cycles.

#### **Specificity:**

p-ZAP70 (Y315) polyclonal antibody detects endogenous levels of ZAP70 protein only when phosphorylated at Tyr315.

#### **DATA:**

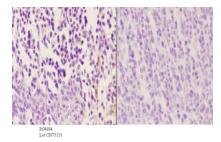


Western blot (WB) analysis of p-ZAP70 (Y315) polyclonal antibody at 1:500 dilution

Lane1:Hela cell lysate treated with H2O2(100 µM,15mins)

Lane2:Mouse kidney tissue lysate

Lane3:Rat kidney tissue lysate



Immunohistochemistry (IHC) analyzes of p-ZAP70 (Y315) pAb in paraffin-embedded human tonsil carcinoma tissue at 1:50.showing cytoplasmic and cell membrane staining. Negative control (the right)Using PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit IgG-biotin followed by avidin-peroxidase.

# Note:

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

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