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# Datasheet for ABIN5691024 STAT2 ELISA Kit

3 Images



#### Overview

Quantity:	96 tests
Target:	STAT2
Binding Specificity:	pTyr689
Reactivity:	Human
Method Type:	Sandwich ELISA
Application:	ELISA

### Product Details

Purpose:	Human Phospho-STAT2 (Tyr689) and Total STAT2 ELISA Kit. This assay semi-quantitatively measures STAT2 phosphorylated at Tyrosine-689 as well as total STAT2 in cell lysate samples.
Sample Type:	Cell Samples, Tissue Lysate
Analytical Method:	Semi-Quantitative
Detection Method:	Colorimetric
Specificity:	This ELISA kit recognizes Human STAT2 phosphorylated at site Tyrosine-689 as well as total STAT2.
Characteristics:	<ul> <li>Pre-Coated 96-well Strip Microplate</li> <li>Wash Buffer</li> <li>Anti-Phospho Antibody</li> <li>Anti-Pan Antibody</li> <li>HRP-Conjugated Secondary Antibody</li> <li>Streptavidin-Conjugated HRP</li> <li>Assay Diluent</li> </ul>

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	TMB One-Step Substrate
	Stop Solution
	Lysis Buffer
	Positive Control Sample
Components:	Pre-Coated 96-well Strip Microplate
	• Wash Buffer
	Anti-Phospho Antibody
	Anti-Pan Antibody
	HRP-Conjugated Secondary Antibody
	Streptavidin-Conjugated HRP
	Assay Diluent
	TMB One-Step Substrate
	Stop Solution
	Lysis Buffer
	Positive Control Sample
Material not included:	Distilled or deionized water
	<ul> <li>100 mL and 1 liter graduated cylinders</li> </ul>
	Tubes to prepare sample dilutions
	Protease and Phosphatase inhibitors
	- Precision pipettes to deliver 2 $\mu$ L to 1 mL volumes
	Adjustable 1-25 mL pipettes for reagent preparation
	Benchtop rocker or shaker
	Microplate reader capable of measuring absorbance at 450 nm

## Target Details

Target:	STAT2
Alternative Name:	STAT2 (STAT2 Products)
Gene ID:	6773
UniProt:	P52630
Pathways:	JAK-STAT Signaling, Hepatitis C, CXCR4-mediated Signaling Events

## Application Details

Plate:	Pre-coated
Protocol:	1. Prepare all reagents and samples as instructed in the manual. 2. Add 100 μL of sample or positive control to each well.

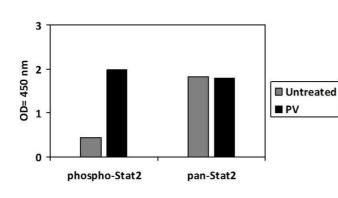
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	3. Incubate 2.5 h at RT or O/N at 4 °C.
	4. Add 100 µL of prepared primary antibody to each well.
	5. Incubate 1 h at RT.
	6. Add 100 µL of prepared 1X HRP-Streptavidin to each well.
	7. Incubate 1 h at RT.
	8. Add 100 $\mu$ L of TMB One-Step Substrate Reagent to each well.
	9. Incubate 30 min at RT.
	10. Add 50 μL of Stop Solution to each well.
	11. Read at 450 nm immediately.
Restrictions:	For Research Use only
Handling	
Storage:	4 °C
Storage Comment:	Upon receipt, the kit should be stored at -20 °C. Please use within 6 months from the date of

torage Comment:	Upon receipt, the kit should be stored at -20 °C. Please use within 6 months from the date of
	shipment. After initial use, Wash Buffer Concentrate (Item B), Assay Diluent (Item E), TMB One-
	Step Substrate Reagent (Item H), HRP-Streptavidin (Item G), Stop Solution (Item I) and Cell
	Lysate Buffer (Item J) should be stored at 4 °C to avoid repeated freeze-thaw cycles. Return
	unused wells to the pouch containing desiccant pack, reseal along entire edge and store at -20
	°C. Reconstituted Positive Control (Item K) should be stored at -70 °C.

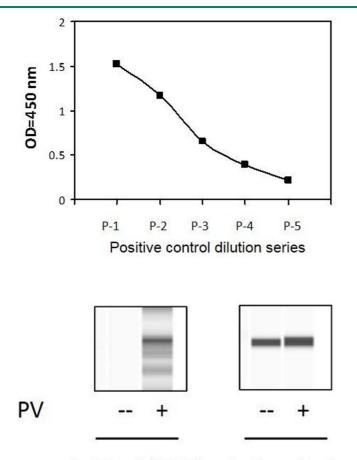
Expiry Date: 6 months

## Validation report #103596 for Western Blotting (WB)



#### **ELISA**

**Image 1.** K562 cells were untreated or treated with PV. Cell lysates were analyzed using this phosphoELISA and Western Blot.



#### ELISA

**Image 2.** K562 cells were treated with Pervanadate (PV). Cells were solubilzed at 4 x 10<sup>^7</sup> cells/ml in Cell Lysate Buffer. Serial dilutions of lysates were analyzed in this ELISA.

### ELISA

**Image 3.** K562 cells were untreated or treated with PV. Cell lysates were analyzed using this phosphoELISA and Western Blot.

Anti Stat2 (Y689)

Anti pan Stat2

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