

## Datasheet for ABIN5068087

# **Histamine ELISA Kit**



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Quantity:	1 kit		
Target:	Histamine (HIS)		
Reactivity:	Chemical		
Method Type:	Competition ELISA		
Application:	ELISA		
Product Details			
Purpose:	This kit is designed for in vitro quantification of histamine in various biological fluids by		
	competitive direct enzyme-linked immunosorbent assay (CD-ELISA). This kit is intended for use		
	in investigative biomedical research only. It is not for human clinical diagnostic use. This		
	Histamine ELISA Kit should not be used for determining histamine levels in scombroid fish.		
Sample Type:	Cell Culture Lysate, Cell Samples, Cell-free Extracts, Tissue Culture Medium		
Analytical Method:	Quantitative		
Detection Method:	Colorimetric		
Sensitivity:	2.5 ng/mL		
Characteristics:	Histamine (HIS) ELISA Kit		
Components:	Wash Buffer (25X), 1x30ml		
	TMB Substrate, 1x20ml		
	PBS Sample Diluent, 1x1Liter		
	Histamine Enzyme (HRP), 1x6ml		
	Standard (0ng/ml), 1x500ul		
	<ul> <li>Standard (2.5ng/ml), 1x500ul</li> </ul>		

- Standard (5ng/ml), 1x500ul
- · Standard (10ng/ml), 1x500ul
- Standard (20ng/ml), 1x500ul
- Standard (50ng/ml), 1x500ul
- · Microtiter Plate, 1x96 wells

#### **Target Details**

Target:	Histamine (HIS)	
Abstract:	HIS Products	
Target Type:	Chemical	

### **Application Details**

#### Protocol:

Assay Precision:

#### Principle of the Assay:

Precision: ≤10% ,Precision: ≤10%

• This Histamine ELISA Kit is a competitive direct ELISA in a microwell format that allows users to obtain exact concentrations of histamine in nanograms per milliliter. The microwells in this assay kit are pre-coated with a monoclonal antibody to histamine. The sample or standard solution is first added to the antibody coated microplate. Next, the enzyme conjugate is added and the mixture is shaken and incubated at RT for 45 minutes. During the incubation, unbound (free) histamine in the samples or standards is allowed to compete with enzyme HRP-labeled histamine (conjugate) for antibody binding sites. The plate is then washed, removing all the unbound material. The bound enzyme conjugate is detected by the addition of a one-component substrate which generates color by horseradish peroxidase. An optimal color is generated after 30 minutes. A microplate reader is then used to take an absorbance reading at 650nm.

Restrictions:	For Research Use only			
Handling				
Storage:	4 °C			
Storage Comment:	4°C			