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Caspase 7 ELISA Kit

Images



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Quantity:	96 tests	
Target:	Caspase 7 (CASP7)	
Binding Specificity:	Cleaved-Asp198, Uncleaved	
Reactivity:	Human, Mouse	
Method Type:	Sandwich ELISA	
Application:	ELISA	
Product Details		
Purpose:	Human/Mouse CASP-7 (D198) ELISA Kit. This ELISA is for measuring cleaved CASP-7 (Asp-198) as well as CASP-7 in human and mouse cell lysates.	
Sample Type:	Cell Lysate, Tissue Lysate	
Analytical Method:	Semi-Quantitative	
Detection Method:	Colorimetric	
Specificity:	The antibody pair provided in this kit recognizes human/mouse cleaved-caspase-7 cleaved at site Aspartic Acid-198 as well as caspase-7.	
Characteristics:	 Simultaneously measure cleaved protein and pan protein in one experiment (for normalization purpose) Screen numerous different cell lysates without performing a Western Blot analysis Minimal hands-on time, convenient, and non-radioactive material 	
Components:	Pre-Coated 96-well Strip MicroplateWash Buffer	

- · Anti-cleaved 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
- · 100 mL and 1 liter graduated cylinders
- Tubes to prepare sample dilutions
- · Protease and Phosphatase inhibitors
- Precision pipettes to deliver 2 µL 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:	Caspase 7 (CASP7)
Alternative Name:	Caspase-7 (CASP7 Products)
UniProt:	P55210
Pathways:	Apoptosis, Caspase Cascade in Apoptosis, Positive Regulation of Endopeptidase Activity

Application Details

Application Details			
Application Notes:	Optimal working dilution should be determined by the investigator.		
Sample Volume:	100 μL		
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.		
	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 μL 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: -20 °C

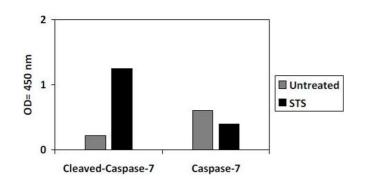
Storage Comment:

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:

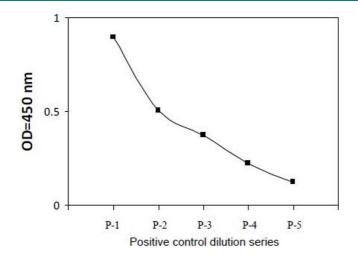
6 months

Images



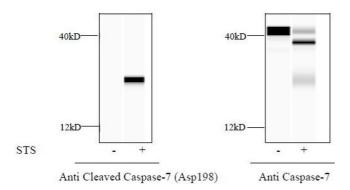
ELISA

Image 1. HeLa cells were treated or untreated with STS. Cell lysates were analyzed using this ELISA and Western Blot.



ELISA

Image 2. HeLa cells were treated with STS. Solubilize cells at 4×10^{7} cells/ml in Cell Lysate Buffer. Serial dilutions of lysates were analyzed in this ELISA.



ELISA

Image 3. HeLa cells were treated or untreated with STS. Cell lysates were analyzed using this ELISA and Western Blot.