



HSP70 ELISA Kit (PLANT)

Catalog number: EK7115

For detection of multiple analytes using one single assay.

This package insert must be read in its entirety before using this product.

For research use only. Not for use in diagnostic procedures.

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Size: 96wells/kit

Sample Type: Cell lysates, Plant extract, Plasma, Serum, Tissue

Sensitivity: 0.18 ng/ml

Assay Range: 1.563 - 100 ng/ml

Storage: Store at 4°C.

Introduction

Boster's ELISA Kit is for the detection of cytoplasmic Plant Hsp70 from plant extracts. But this kit is also compatible with human tissue extracts and lysates and serum and plasma. Each kit contains sufficient components to quantitate the Hsp70 concentration in up to 40 samples, tested in duplicate.

Kit Components

Description	Quantity
Anti-Hsp70 Immunoassay Plate	1 Plate
5X Hsp70 Extraction Reagent	1 vial/10 ml
Recombinant Hsp70 (Alfa) Standard	2 vials
Standard and Sample Diluent	1 vial/ 50 ml
10X Wash Buffer Concentrate	1 vial/100 ml
Anti-Hsp70 Biotinylated Antibody Concentrate	1 vial/150 µl
Anti-Hsp70 Biotinylated Antibody Diluent	1 vial/ 13 ml
Streptavidin: HRP Concentrate	1 vial/150 µl
Streptavidin: HRP Diluent	1 vial/ 13 ml
TMB Substrate	1 vial/ 13 ml
Stop Solution	1 vial/ 13 ml

Materials Required, but Not Provided

1. Ultra pure water.
2. Additional reagents and materials for cell lysate and tissue extract preparation, including protease inhibitors.
3. Precision pipettors, with disposable plastic tips.
4. Polypropylene or polyethylene tubes to prepare samples – do not use polystyrene, polycarbonate or glass tubes.
5. A container to prepare 1X Wash Buffer.

6. A wash bottle or an automated 96-well plate washer.
7. Disposable reagent reservoirs.
8. A standard microtiter plate reader for measuring absorbance at 450 nm.
9. Adhesive plate sealers.

Assay Overview

1. Prepare Standard and samples in Standard and Sample Diluent.
2. Add 100 μ L of Standard to appropriate wells.
3. Cover plate with Plate Sealer and incubate at room temperature for 1 hour.
4. Wash plate four times with Wash Buffer.
5. Add 100 μ L of Detection Antibody Working Solution to each well.
6. Cover plate with Plate Sealer and incubate at room temperature for 1 hour.
7. Wash plate four times with 1X Wash Buffer as described in step 4.
8. Add 100 μ L of Streptavidin-HRP Working Solution to each well.
9. Cover plate with Plate Sealer and incubate at room temperature for 30 minutes.
10. Wash plate four times with 1X Wash Buffer as described in step 4.
11. Add 100 μ L of TMB Substrate to each well.
12. Develop the plate in the dark at room temperature for 30 minutes.
13. Stop reaction by adding 100 μ L of Stop Solution to each well.
14. Measure absorbance on a plate reader at 450 nm.

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