MSD[®] Total ERK1/2 Assay Whole Cell Lysate Kit

For quantitative determination in human, mouse, and rat whole cell lysate samples

Alzheimer's Disease BioProcess Cardiac Cell Signaling

Clinical Immunology Cytokines Hypoxia Immunogenicity Inflammation Metabolic Oncology Toxicology Vascular

Catalog Numbers

Total ERK1/2 Assay Whole

Cell Lysate Kit Kit size

Phospho-ERK1/2 Whole Cell

Lysate Set

Ordering information

MSD Customer Service

Phone: 1-301-947-2085 Fax: 1-301-990-2776 Email: CustomerService@

mesoscale.com

MESO SCALE DISCOVERY®

Meso Scale Diagnostics, LLC. 9238 Gaither Road

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Company Address

A division of

1 plate

5 plates

20 plates

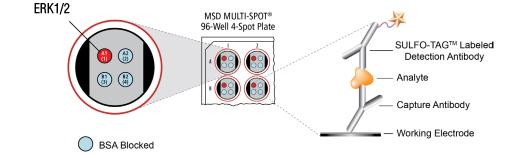
200 µg

K151DXD-1

K151DXD-2

K151DXD-3

C11CM-1



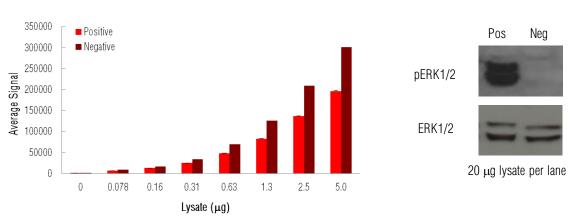
ERK (Extracellular signal Regulated Kinases) 1 and 2, are proline-directed serine/threonine protein kinases, also known as p44 MAPK (Mitogen-Activated Protein Kinase) and p42 MAPK, respectively. These closely-related kinase isoforms, identified by their molecular weight, are activated by various extracellular signals including growth factors, cytokines, hormones, and neurotransmitters. The activation occurs through the phosphorylation of threonine 202 and tyrosine 204 of ERK1 and threonine 185 and tyrosine 187 of ERK2 by the upstream kinases MEK1 and MEK2. Activated ERK1/2 phosphorylates targets in both the cytosol and the nucleus. Cytoplasmic substrates for ERK include SOS, MNK1/2, and the 90 kDa ribosomal protein S6 kinases. Nuclear translocation of activated ERK affects gene expression and DNA replication by the phosphorylation of MSK 1 and 2 and the transcription factors Elk-1, Sap1, and Sap2.

The MSD Total ERK1/2 Assay is available on 96-well 4-Spot plates. This datasheet outlines the performance of the assay.

Typical Data

Representative results for the Total ERK1/2 Assay are illustrated below. The signal and ratio values provided below are example data; individual results may vary depending upon the samples tested. Western blot analyses of each lysate type were performed with phospho-ERK1/2 (Thr202/Tyr204; Thr185/Tyr187) and total ERK1/2 antibodies and are shown below for comparison. Logarithmically growing Jurkat cells were treated with PMA (200 nM; 15 minutes) (positive) or LY294002 (50 µM; 2.5 hours)

(negative). Whole cell lysates were added to MSD MULTI-SPOT[®] 4-Spot plates coated with anti-total ERK1/2 antibody on one of the four spatially distinct electrodes within a well. Total ERK1/2 was detected with anti-total ERK1/2 antibody conjugated with MSD SULFO-TAG[™] reagent.



For Research Use Only. Not for use in diagnostic procedures. Fig. 1: Sample data generated with the MULTI-ARRAY[®] Total ERK1/2 Assay. Increased signals were observed with the titration of both pERK1/2 positive and negative cell lysates. The Total ERK1/2 Assay provides a quantitative measure of the data obtained with the traditional Western blot.

pot the Difference



Total ERK1/2 + PMSF + SDS

Lysate Titration

Data for pERK1/2 positive and negative Jurkat cell lysates using the MULTI-ARRAY Total ERK1/2 Assay are presented below.

Lysate	Positive			Negative			P/N
(µg)	Average Signal	StdDev	%CV	Average Signal	StdDev	%CV	F/N
0	653	15	2.3	709	21	3.0	
0.078	6858	471	6.9	8864	68	0.8	0.8
0.16	13163	962	7.3	17032	530	3.1	0.8
0.31	24988	1479	5.9	34442	1334	3.9	0.7
0.63	48321	2353	4.9	69276	507	0.7	0.7
1.3	82995	3214	3.9	126071	2421	1.9	0.7
2.5	136849	6824	5.0	209256	2675	1.3	0.7
5.0	196096	9104	4.6	300653	3852	1.3	0.7

MSD Advantage

- Multiplexing: Multiple analytes can be measured in one well using typical sample amounts of 25 μg/well or less without compromising speed or performance
- Large dynamic range: Linear range of up to five logs enables the measurement of native levels of biomarkers in normal and diseased samples without multiple dilutions
- Minimal background: The stimulation mechanism (electricity) is decoupled from the signal (light)
- Simple protocols: Only labels near the electrode surface are detected, enabling no-wash assays
- > Flexibility: Labels are stable, non-radioactive, and conveniently conjugated to biological molecules
- > High sensitivity and precision: Multiple excitation cycles of each label enhance light levels and improve sensitivity

For a complete list of products, please visit our website at www.mesoscale.com

References using MSD's platform for the measurement of phosphoproteins

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