

Phospho EGF Receptor MAPmates™

Cat. # 46-603

MILLIPLEX® MAP

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INTRODUCTION

The MILLIPLEX MAP Phospho EGF Receptor MAPmates™ pair is used in conjunction with the MILLIPLEX MAP Cell Signaling Buffer and Detection Kit (Catalog # 48-602) to detect the presence of phosphorylated EGF Receptor in cell lysates using the Luminex® 100™ IS, 200™, or HTS system. Each MAPmates pair is ordered individually and may be combined for simultaneous multiplex analysis of cellular events. The MILLIPLEX MAP Cell Signaling Buffer and Detection Kit is ordered separately and consists of a common set of reagents needed for performing MAPmates assays. The detection assay is a rapid, convenient alternative to Western Blotting and immunoprecipitation procedures. Each kit contains sufficient reagents for 100 individual assays. The MILLIPLEX MAP HeLa Cell Lysate: Unstimulated included in the MILLIPLEX MAP Cell Signaling Buffer and Detection Kit may be utilized as a negative control for this target, while the A431 Cell Lysate: EGF Stimulated included with this MAPmates pair may be utilized as a positive control.

Important note: For a detailed protocol on Cell Signaling Detection Procedures please see the instruction booklet for the MILLIPLEX MAP Cell Signaling Buffer and Detection Kit (#48-602).

REAGENTS SUPPLIED	CATALOG NUMBER	LUMINEX BEAD#	VOLUME	QUANTITY
MILLIPLEX MAP Anti-phospho EGF Receptor Beads (20X)	42-603	34	131 μL	1 tube
MILLIPLEX MAP Anti-phospho EGF Receptor, biotin (20X)	44-603	n/a	131 μL	1 tube
MILLIPLEX MAP A431 Cell Lysate: EGF Stimulated	47-210	n/a		1 vial

STORAGE CONDITIONS UPON RECEIPT

- Recommended storage for kit components is 2 8 °C.
- DO NOT FREEZE Streptavidin-Phycoerythrin.

This kit is for research purposes only.

Please read entire protocol before use.

It is important to use same assay incubation conditions throughout your study.

By purchasing this product, which contains fluorescently labeled microsphere beads authorized by Luminex Corporation ("Luminex"), you, the customer, acquire the right under Luminex's patent rights, if any, to use this product or any portion of this product, including without limitation the microsphere beads contained herein, only with Luminex's laser based fluorescent analytical test instrumentation marketed under the name of Luminex100, 200, HTS.

MATERIALS REQUIRED BUT NOT PROVIDED

Reagents

 MILLIPLEX MAP Cell Signaling Buffer and Detection Kit (Catalog # 48-602) contains necessary reagents and instructions for sample, control, bead and detection antibody preparation

<u>Instrumentation / Materials</u>

- Adjustable Pipettes with Tips capable of delivering 25 μL to 1000 μL
- Multichannel Pipettes capable of delivering 25 μL to 200 μL
- Reagent Reservoirs
- Polypropylene Microfuge Tubes
- Rubber Bands
- Absorbent Pads
- Laboratory Vortex Mixer
- Sonicator
- Titer Plate Shaker
- Vacuum Filtration Unit (Millipore Vacuum Manifold Catalog #MSVMHTS00 or equivalent with Millipore Vacuum Pump Catalog #WP6111560 or equivalent)
- Luminex 100™ IS, 200™, or HTS by Luminex Corporation
- Plate Stand (Millipore Catalog # MX-STAND)

SAFETY PRECAUTIONS

- All tissue components and biological materials should be handled as potentially hazardous. Follow universal precautions as established by the Centers for Disease Control and Prevention and by the Occupational Safety and Health Administration when handling and disposing of infectious agents.
- Sodium azide or Proclin has been added to some reagents as a preservative. Although the concentrations are low, sodium azide and Proclin may react with lead and copper plumbing to form highly explosive metal azides. On disposal, flush with a large volume of water to prevent azide build up.

TECHNICAL CONSIDERATIONS

- The suggested working range of protein concentration for the assay is 1 to 25 μ g of total protein/well (25 μ L/well at 40 to 1000 μ g/mL). A total protein amount of 10 μ g/ well is generally a good starting point for lysates for which target protein expression levels are unknown.
- The following MAPmates should not be multiplexed:
 - a. phosho-specific and total MAPmate pairs
 - b. pTyr and site-specific phospho MAPmates (the pTyr detect may generate false positives on the site-specific MAPmates)
 - c. Phospho MAPmates for a single target (Akt, STAT3, p53)

PREPARATION OF LYOPHILIZED MILLIPLEX MAP CELL LYSATE

MILLIPLEX MAP A431 Cell Lysate: EGF stimulated as a positive control

- Reconstitute the lyophilized cell lysate in 100 μ L of ultrapure water, this will yield 100 μ L of lysate at 2 mg/mL total protein.
- Gently vortex and incubate the reconstituted lysate for 5 min at RT (store on ice).
- Pipette 150 μL MILLIPLEX MAP Assay Buffer 2 to the reconstituted cell lysate vial. The cell lysate is now prepared for use in the MILLIPLEX MAP Cell Signaling Assays.
- Combine prepared lysate (25 μ L/well) with 25 μ L Cell Signaling beads (step 4 of Immunoassay Protocol) and proceed with assay.

IMMUNOASSAY PROTOCOL

- 1. Dilute filtered lysates <u>at least</u> 1:1 in MILLIPLEX MAP Assay Buffer 2. The suggested working range of protein concentration for the assay is 1 to 25 μg of total protein/well (25 μL/well at 40 to 1,000μg/ml).
- Pre-wet filter plate with 25 μL/well of MILLIPLEX MAP Assay Buffer 2. Remove by vacuum filtration by placing the filter plate over a vacuum manifold and gently applying vacuum. Gently blot the bottom of the filter plate on a paper towel to remove excess liquid.
- 3. Vortex the **1X** bead suspension for 10 seconds. Add 25 μ L of 1X bead suspension to each well.
- Add 25 μL of diluted cell lysate to each well and incubate overnight at 4°C (or 2 hours RT) on a plate shaker (600-800rpm) protected from light.
- 5. Remove the lysate by vacuum filtration.
- Add 100 μL/well of MILLIPLEX MAP Assay Buffer 2. Remove buffer by vacuum filtration and gently blot the bottom of the filter plate on a paper towel. Repeat this step again for a total of two washes.
- Add 25 μL/well of 1X MILLIPLEX MAP Biotinylated Detection Antibody.
- 8. Incubate on a plate shaker for 1 hour at room temperature, protected from light.
- 9. Remove Biotinylated Detection Antibody by vacuum filtration and gently blot the bottom of the filter plate on a paper towel.
- 10. Add 25 μL of diluted (1:25) MILLIPLEX MAP Streptavidin-Phycoerythrin (SAPE).
- 11. Incubate on a plate shaker for 15 minutes at room temperature, protected from light.
- 12. **DO NOT REMOVE** SAPE. Add 25 μL of MILLIPLEX MAP Amplification Buffer to each well.

Add 25 μL Assay Buffer 2 per well



Remove buffer by vacuum.

- Add 25 μL 1X beads to wells
- Add 25 μL diluted cell lysate to appropriate wells



Incubate overnight at 4 °C or 2 hour at RT with shaking; dark

Wash 2X with 100 μL Assay Buffer 2. Add 25 μL 1X Biotinylated Detection Antibody.



Incubate 1 hr at RT with shaking; dark

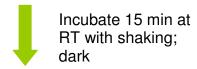
Remove Detection Antibody and add 25 μ L diluted SAPE.



Incubate 15 min at RT with shaking; dark

DO NOT REMOVE SAPE and add 25 μL Amplification Buffer.

- 13. Incubate on a plate shaker for 15 minutes at room temperature, protected from light.
- 14. Remove MILLIPLEX MAP SAPE/Amplification Buffer by vacuum filtration and gently blot the bottom of the filter plate on a paper towel.
- 15. Resuspend beads in 150 μL of MILLIPLEX MAP Assay Buffer 2, and mix on plate shaker for 5 minutes.
- 16. Analyze using the Luminex® system.



Remove Streptavidin-PE/ Amplification buffer and resuspend beads in 150 μL Assay Buffer 2. Read results using appropriate Luminex® instrument.

INSTRUMENT SETTINGS

These specifications are for Luminex_{100/200} instruments. Luminex instruments with other software (e.g. MasterPlex, StarStation, LiquiChip, Bio-Plex, LABScan100) would need to follow instrument instructions for gate settings and additional specifications from the vendors.

Events	50 per bead		
Sample Size	100 μL		
Gate Settings	8,000 to 15,000		
Reporter Gain	Default (Low PMT)		
Time Out	60 seconds		
Bead Set	34		

SINGLE AND MULTI-PLEX ANALYSIS

The recommended lysis and assay buffers for single or multi-plex analysis of Phospho EGF Receptor MAPmates are MILLIPLEX MAP Lysis Buffer (Catalog # 43-040) and MILLIPLEX MAP Assay Buffer 2 (Catalog # 43-041). Both buffers are included in the MILLIPLEX MAP Cell Signaling Buffer and Detection Kit (Catalog # 48-602). For complete cell signaling assay and cell lysis protocols refer to the MILLIPLEX MAP Cell Signaling Buffer and Detection Kit instructions.

MAPmates which are listed in the MILLIPLEX MAP Cell Signaling Buffer and Detection Kit buffer selection chart as "not recommended" for the above buffer conditions must be assayed separately using appropriate buffer conditions.

Note: Phospho and Total MAPmates should not be multiplexed together.

REPRESENTATIVE DATA:

Detection of Phospho EGFR in A431 cells

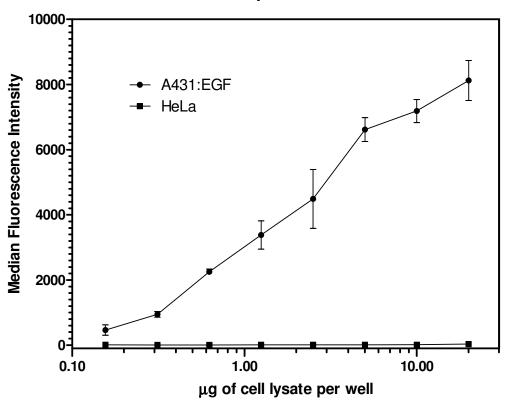
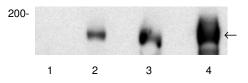


Figure 1. MILLIPLEX MAP detection of phosphorylated EGFR in A431 cell lysate. A431:EGF stimulated (#47-210) and HeLa: unstimulated were prepared according to instructions. Increasing amounts of cell lysate were incubated overnight at 4°C with anti-EGFR capture beads. The beads were washed and probed with biotin labeled anti-phospho EGFR, followed by incubation with streptavidin-PE and amplification buffer. The Median Fluorescent Intensity (MFI) in triplicate wells was measured using the Luminex® Instrument. This graph displays the MFI values obtained utilizing the Phospho EGFR MAPmates.

Figure 2. Western blot detection of phosphorylated EGF receptor in HeLa and A431 cell lysates. HeLa cells and A431 cells were grown to confluence and stimulated with or without 50ng/ml of EGF for 20 minutes. 10μg/well of unstimulated HeLa (lane 1), stimulated HeLa (lane 2), unstimulated A431 (lane 3) or stimulated A431 (lane 4) cell lysate (lysed in MILLIPLEX MAP Lysis Buffer with protease inhibitors) were separated by SDS-PAGE, transferred to nitrocellulose, and probed with mouse monoclonal anti-phosphotyrosine. Blots were incubated with HRP labeled anti-mouse lgG and visualized via chemiluminescence. Arrow indicates phosphorylated EGF receptor (175 kDa).



ORDERING INFORMATION

To place an order:

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Include:

- Your name, telephone and/or fax number
- Customer account number
- Shipping and billing address
- Purchase order number
- Catalog number and description of product
- · Quantity of kits

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6 Research Park Drive

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