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TECHNICAL DATA SHEET



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BETTER TOOLS. REAL DISCOVERIES.

THUNDER™ Phospho-RPS6 (S235/S236) + Total RPS6 TR-FRET Cell Signaling Assay Kit

CATALOG NUMBERS KIT-RPS6APT-500
400 points for phospho-RPS6
and 100 points for total RPS6

Store at -80°C
For research use only.
Not for use in diagnostic procedures.

PRODUCT DESCRIPTION

This assay kit measures intracellular levels of **phospho-RPS6 (S235/S236)** and **total RPS6** protein in cell lysates using a simple, rapid and sensitive immunoassay based on the homogeneous (no-wash) THUNDER™ TR-FRET technology. The kit is compatible with both adherent and suspension cells.

SPECIFICITY

This assay kit contains two specific and selective antibody pairs, one that recognizes **RPS6** phosphorylated at **Ser235** and **Ser236** and another that recognizes **total** (both phosphorylated and unphospho-rylated) **RPS6**.

SPECIES REACTIVITY

Human (Swiss-Prot Acc.: P62753; Entrez-Gene Id: 6194).

Other species should be tested on a case-by-case basis.

TR-FRET ASSAY PRINCIPLE

The **Phospho-RPS6 (S235/S236) + Total RPS6** assay kit is a homogeneous time-resolved Förster resonance energy transfer (TR-FRET) sandwich immunoassay (Figure 1). The THUNDER™ Cell Signaling assay workflow consists of 3 steps (Figure 2). Following cell treatment, cells are first lysed with the specific Lysis Buffer provided in the kit. Then **Phospho-RPS6 (S235/S236)** and **Total RPS6** in the cell lysates are detected in separate wells with two pairs of fluorophore-labeled antibodies in a simple "add-incubate-measure" format (single-step reagent addition; no wash steps). For detection of the phosphorylated protein, one antibody is labeled with a donor fluorophore (Europium chelate; Eu-Ab1) and the second with a far-red acceptor fluorophore (FR-Ab2). The same approach is used for the second antibody pair detecting the total protein (Eu-Ab3 and FR-Ab4). The binding of the two matched labeled antibodies to distinct epitopes on the target protein (either **phospho-RPS6** or **total RPS6**) takes place in solution and brings the two dyes into close proximity. Excitation of the donor Europium chelate molecules with a flash lamp (320 or 340 nm) or a laser (337 nm) triggers a FRET from the donor to the acceptor molecules, which in turn emit a TR-FRET signal at 665 nm. Residual energy from the Eu chelate generates light at 615 nm. The signal at 665 nm is proportional to the concentration of **Phospho-RPS6 (S235/S236)** and **Total RPS6** in the cell lysate. Data can be expressed as either the signal at 665 nm or the 665 nm/615 nm ratio.

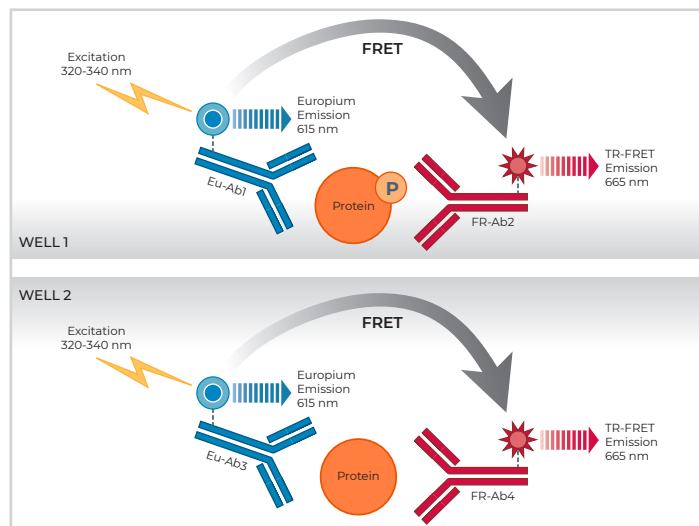


Fig. 1 Schematic representation of the TR-FRET cell signaling assay principle.

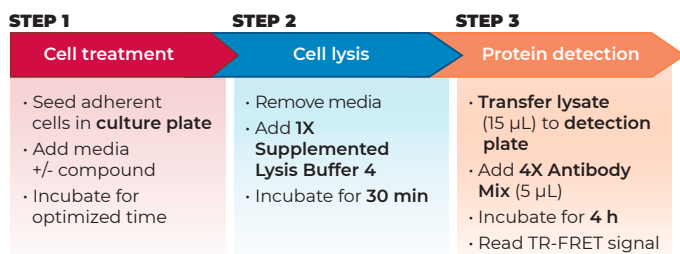


Figure 2 Assay workflow using the 2-plate (transfer) protocol.

KIT COMPONENTS

	500 points*
Eu-labeled phospho-RPS6 (S235/S236) antibody (Eu-Ab1)	20 µL
Acceptor-labeled phospho-RPS6 (S235/S236) antibody (FR-Ab2)	80 µL
Eu-labeled total-RPS6 antibody (Eu-Ab3)	5 µL
Acceptor-labeled total-RPS6 antibody (FR-Ab4)	20 µL
Lysis Buffer 4 (5X)	5 mL
Detection Buffer (10X)	250 µL
Positive control cell lysate	500 µL

* The number of assay points is based on an assay volume of 20 µL in half-area 96-well or low-volume 384-well assay plates using the kit components at the recommended concentrations (refer to the User Manual).

TECHNICAL DATA SHEET

Phospho-RPS6 (S235/S236) + Total RPS6

VALIDATION DATA

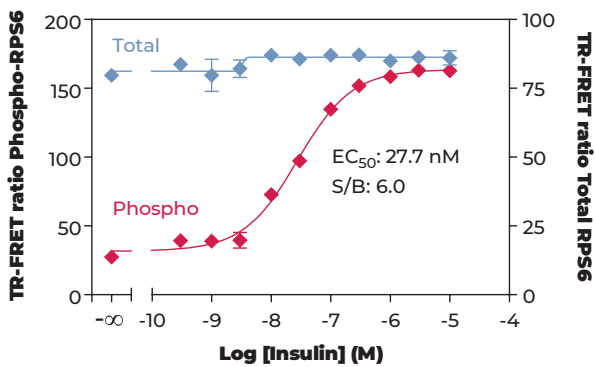
This assay kit has been validated for the relative quantification of phospho-RPS6 (S235/S236) and total RPS6 in MCF7 cell lysates using the 2-plate assay protocol.

- Adherent cells were cultured overnight in a 96-well tissue culture plate (EMEM +10% FBS) and then serum starved for 18 hours.
- Following cell treatment, the media was removed and cells were lysed with the 1X **Lysis Buffer 4** (50 μ L) supplemented with the phosphatase inhibitors sodium fluoride (1 mM) and sodium orthovanadate (2 mM).
- Following a **30-min** incubation at room temperature (RT) on an

orbital shaker (400 rpm), lysates (15 μ L) were then transferred to a 384-well assay plate followed by addition to separate wells of either the labeled antibodies Eu-Ab1 and FR-Ab2 (5 μ L) for detection of phospho-RPS6 (S235/S236) or Eu-Ab3 and FR-Ab4 (5 μ L) for detection of total RPS6.

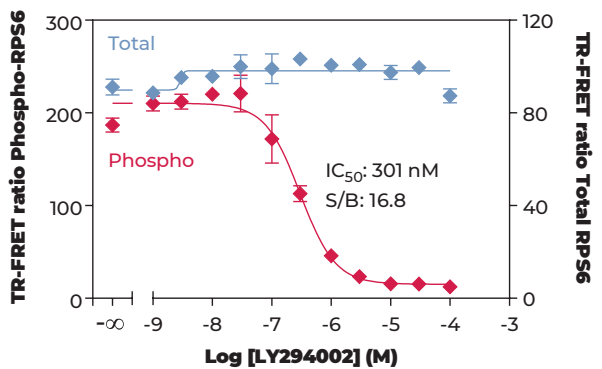
- The plate was incubated at RT for **4 hours** and the TR-FRET signal was recorded at 665 and 615 nm (EnVision[®]; lamp excitation).

STIMULATION OF PHOSPHO-RPS6 (S235/S236) IN MCF7 CELLS



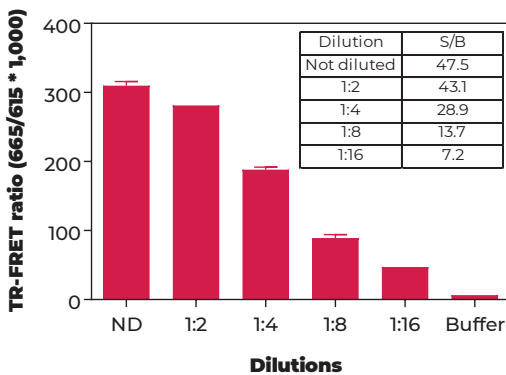
MCF7 cells (75,000 cells/well; in triplicate) were incubated with serial dilutions of insulin for 60 min at RT. Data show that treatment of MCF7 cells with insulin stimulates phosphorylation of RPS6 at S235/S236, but does not affect the levels of total RPS6.

INHIBITION OF PHOSPHO-RPS6 (S235/S236) IN MCF7 CELLS



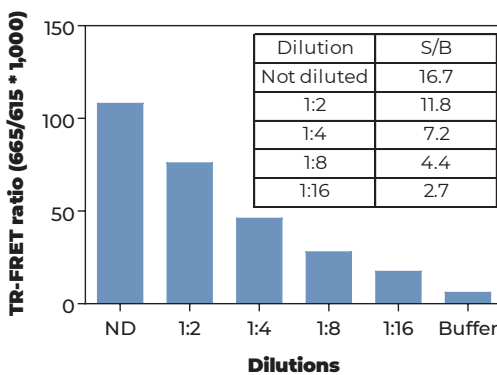
MCF7 cells (75,000 cells/well; in triplicate) were incubated with serial dilutions of the inhibitor LY294002 for 60 min at RT. Cells were then stimulated with 1 μ M of insulin for 60 min at RT. Data show that treatment of MCF7 cells with LY294002 inhibits phosphorylation of RPS6 at S235/S236 by insulin, but does not affect the levels of total RPS6.

MCF7 CONTROL LYSATE TITRATION (QC TEST) PHOSPHO-RPS6 (S235/S236)



Quality Control: the Phospho-RPS6 (S235/S236) + Total RPS6 assay kit is routinely tested against Insulin-treated MCF7 lysates. MCF7 cells were cultured in a T175 flask to 90% confluence and stimulated with 10 μ M of insulin for 60 min at RT. Following cell lysis using 16 mL of 1X Lysis Buffer 4, lysates were serially diluted with 1X Lysis Buffer 4 and tested in triplicate and in separate wells for phospho-RPS6 (S235/S236) and total RPS6. Data show a linear relationship between lysate dilutions and TR-FRET ratio values.

MCF7 CONTROL LYSATE TITRATION (QC TEST) TOTAL RPS6



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FOR MORE INFORMATION ON DEVELOPING AND OPTIMIZING TR-FRET CELL SIGNALING ASSAYS, CONSULT THE USER MANUAL.

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