Human EGFR [pY1068] ELISA Kit

Catalog Number KHR9081 (96 tests)

Pub. No. MAN0014764 **Rev.** 2.0 (30)

CAUTION! This kit contains materials with small quantities of sodium azide. Sodium azide reacts with lead and copper plumbing to form explosive metal azides. Upon disposal, flush drains with a large volume of water to prevent azide accumulation. Avoid ingestion and contact with eyes, skin and mucous membranes. In case of contact, rinse affected area with plenty of water. Observe all federal, state, and local regulations for disposal.

Note: For safety and biohazard guidelines, see the "Safety" appendix in the *ELISA Technical Guide* (Pub. no. MAN0006706). Read the Safety Data Sheets (SDSs) and follow the handling instructions. Wear appropriate protective eyewear, clothing, and gloves.

Product description

The Invitrogen™ Human EGFR [pY1068] ELISA Kit is a solid-phase sandwich Enzyme-Linked Immunosorbent Assay (ELISA). This assay is designed to detect and quantify the level of phosphorylated human EGFR [pY1068] in cell and tissue lysates. The assay recognizes both natural and recombinant human EGFR [pY1068].

The epidermal growth factor receptor (EGFR) belongs to the family of receptor tyrosine kinases (RTKs), which regulate cell growth, survival, proliferation and differentiation. EGFR is a 170 kDa type I transmembrane glycoprotein consisting of an extracellular ligand-binding domain, a single hydrophobic transmembrane region, and an intracellular segment with a highly conserved, tyrosine kinase domain.

For normalizing the EGFR content of the samples, a Human EGFR (Full length) ELISA Kit (Cat. No. KHR9061), is available for detection of EGFR content independent of phosphorylation status.

Contents and storage

Upon receipt, store the kit at 2°C to 8°C.

Contents	Cat. No. KHR9081 (96 tests)
Hu EGFR [pY1068] Standard, lyophilized; contains 0.1% sodium azide	2 vials
Standard Diluent Buffer; contains 0.1% sodium azide	25 mL
Hu EGFR Antibody Coated Wells, 96-well plate	1 plate
Hu EGFR [pY1068] Detection Antibody; contains 0.1% sodium azide	11 mL
Anti-Rabbit IgG HRP (100X)	0.125 mL
HRP Diluent; contains 3.3 mM thymol	25 mL
Wash Buffer Concentrate (25X)	100 mL
Stabilized Chromogen, Tetramethylbenzidine (TMB)	25 mL
Stop Solution	25 mL
Plate Covers, adhesive strips	3

Required materials not supplied

- Distilled or deionized water
- Microtiter plate reader with software capable of measurement at or near 450 nm
- Plate washer–automated or manual (squirt bottle, manifold dispenser, or equivalent)
- Calibrated adjustable precision pipettes and glass or plastic tubes for diluting solutions; beakers, flask and cylinders for preparation of reagents
- Cell Extraction Buffer (Cat. No. FNN0011, or see "Prepare Cell Extraction Buffer")

Before you begin

IMPORTANT! Reagents are lot-specific. Do not mix or interchange different reagent lots from various kit lots.

- Review the **Procedural guidelines** and **Plate washing directions** in the *ELISA Technical Guide* available at **thermofisher.com**.
- Allow reagents to reach room temperature before use. Mix to redissolve any precipitated salts.

Prepare 1X Wash Buffer

1. Dilute 16 mL of Wash Buffer Concentrate (25X) with 384 mL of deionized or distilled water. Label as 1X Wash Buffer.
2. Store the concentrate and 1X Wash Buffer in the refrigerator. Use the diluted buffer within 14 days.

Prepare Cell Extraction Buffer

Note: See the *ELISA Technical Guide* for detailed information on preparing Cell Extraction Buffer.

1. Prepare Cell Extraction Buffer.

   Cell Extraction Buffer consists of 10 mM Tris (pH 7.4), 100 mM NaCl, 1 mM EDTA, 1 mM EGTA, 1 mM NaF, 20 mM $Na_4P_2O_7$, 2 mM Na_3VO_4, 1% Triton™ X-100, 10% glycerol, 0.1% SDS, and 0.5% deoxycholate.
2. Immediately before use, add PMSF (0.3 M stock in DMSO) to 1 mM and 50 µL protease inhibitor cocktail (e.g., Sigma Cat. No. P-2714) for each 1 mL of Cell Extraction Buffer.

Prepare cell lysate

1. Collect cells by centrifugation (non-adherent cells) or scraping from culture flasks (adherent cells), then wash cells twice with cold PBS.
2. Remove and discard the supernatant and collect the cell pellet. The pellet can be stored at −80°C and lysed at a later date if desired.
3. Lyse the cell pellet in Cell Extraction Buffer for 30 minutes, on ice. Vortex at 10-minute intervals.

   Note: The volume of Cell Extraction Buffer used depends on the number of cells in the cell pellet, and expression levels of human EGFR [pY1068]. [Researchers must optimize the extraction procedures for their own applications.]
4. Transfer the lysate into microcentrifuge tubes and centrifuge at 13,000 rpm for 10 minutes at 4°C.
5. Transfer the supernatant into clean microcentrifuge tubes. Samples can be stored at −80°C (avoid multiple freeze-thaw cycles).

Pre-dilute samples

Sample concentrations should be within the range of the standard curve. Because conditions may vary, each investigator should determine the optimal dilution for each application.

- Perform sample dilutions with Standard Diluent Buffer.
- Dilute samples prepared in Cell Extraction Buffer 1:10 or greater in Standard Diluent Buffer (e.g., 10 µL sample into 90 µL buffer).

  This dilution is necessary to reduce the matrix effect of the Cell Extraction Buffer. SDS concentration should be less than 0.01% before adding to the plate. While a 1:10 sample dilution has been found to be satisfactory, higher dilutions such as 1:25 or 1:50 may be optimal.

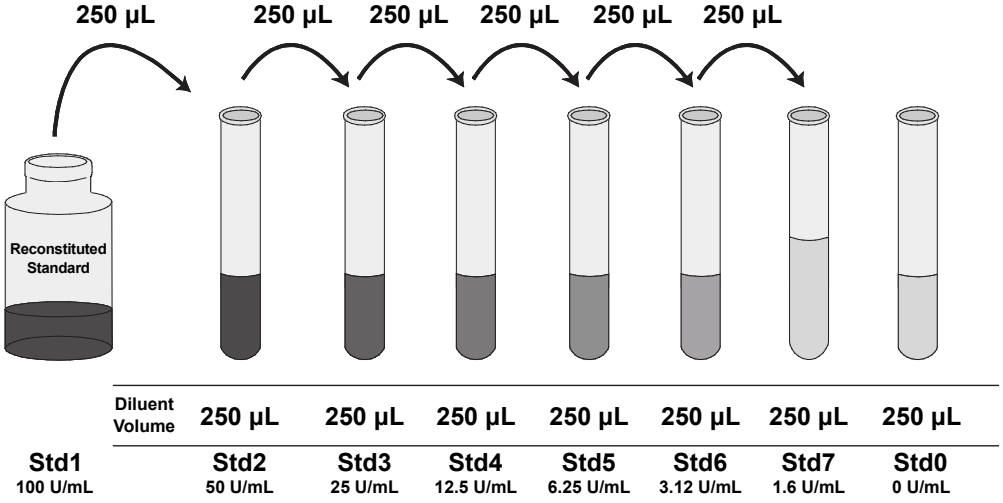
  For 10^8 A431 cells use of 0.1–1 µL of the clarified cell extract diluted to 100 µL in Standard Diluent Buffer for each well.

Dilute standards

Note: Use glass or plastic tubes for diluting standards.

Note: This Hu EGFR [pY1068] Standard was prepared from full length EGFR purified from A431 cells. The purified protein was allowed to autophosphorylate in the presence of 2 mM ATP and 1X Autophosphorylation Buffer (15 mM HEPES, 6 mM $MnCl_2$, 15 mM $MgCl_2$). One unit of standard is equivalent to the amount of EGFR [pY1068] derived from 42 pg of EGFR allowed to autophosphorylate. Subsequent lots of standard will be normalized to this lot of material to allow consistency of human EGFR [pY1068] quantitation.

1. Reconstitute Hu EGFR [pY1068] Standard to 100 Units/mL with Standard Diluent Buffer. Refer to the standard vial label for instructions. Swirl or mix gently and allow the contents to sit for 10 minutes to ensure complete reconstitution. Label as 100 Units/mL human EGFR [pY1068]. **Use the standard within 1 hour of reconstitution.**
2. Add 250 µL Standard Diluent Buffer to each of 7 tubes labeled as follows: 50, 25, 12.5, 6.25, 3.12, 1.6, and 0 Units/mL human EGFR [pY1068].
3. Make serial dilutions of the standard as shown in the following dilution diagram. Mix thoroughly between steps.
4. Discard all remaining reconstituted and diluted standards after completing assay. Return the Standard Diluent Buffer to the refrigerator.



Prepare 1X Anti-Rabbit IgG HRP solution

Note: Prepare 1X Anti-Rabbit IgG HRP solution within 15 minutes of usage.

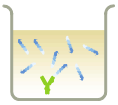
1. For each 8-well strip used in the assay, pipet 10 µL Anti-Rabbit IgG HRP (100X) solution, and dispense the solution into a tube containing 1 mL of HRP Diluent. Mix thoroughly.
2. Return the unused Anti-Rabbit IgG HRP (100X) solution to the refrigerator.

Perform ELISA (Total assay time: 4 hours)

IMPORTANT! Perform a standard curve with each assay.

- Allow all components to reach room temperature before use. Mix all liquid reagents prior to use.
- Determine the number of 8-well strips required for the assay. Insert the strips in the frames for use. Re-bag any unused strips and frames, and store at 2°C to 8°C for future use.

1 Bind antigen



a. Add 100 µL of standards, controls, or samples (see "Pre-dilute samples"on page 2) to the appropriate wells. Leave the wells for chromogen blanks empty.
b. Cover the plate with a plate cover and incubate 2 hours at room temperature. Alternatively, the plate may be incubated overnight at 4 °C.
c. Thoroughly aspirate the solution and wash wells 4 times with 1X Wash Buffer.

2 Add detector antibody

a. Add 100 µL of Hu EGFR [pY1068] Detection Antibody solution into each well except the chromogen blanks.
b. Cover the plate with a plate cover and incubate 1 hour at room temperature.
c. Thoroughly aspirate the solution and wash wells 4 times with 1X Wash Buffer.

3 Add IgG HRP

a. Add 100 µL 1X Anti-Rabbit IgG HRP solution into each well except the chromogen blanks.
b. Cover the plate with plate cover and incubate for 30 minutes at room temperature.
c. Thoroughly aspirate the solution and wash wells 4 times with 1X Wash Buffer.

4 Add Stabilized Chromogen

a. Add 100 µL Stabilized Chromogen to each well. The substrate solution begins to turn blue.
b. Incubate for 30 minutes at room temperature in the dark.

Note: TMB should not touch aluminum foil or other metals.

5 Add Stop Solution

Add 100 µL Stop Solution to each well. Tap the side of the plate to mix. The solution in the wells changes from blue to yellow.

Read the plate and generate the standard curve

1. Read the absorbance at 450 nm. Read the plate within 2 hours after adding the Stop Solution.
2. Use curve-fitting software to generate the standard curve. A 4 parameter algorithm provides the best standard curve fit. Optimally, the background absorbance may be subtracted from all data points, including standards, unknowns and controls, prior to plotting.
3. Read the concentrations for unknown samples and controls from the standard curve. Multiply value(s) obtained for sample(s) by the appropriate factor to correct for the sample dilution.

   Note: Dilute samples producing signals greater than the upper limit of the standard curve in Standard Diluent Buffer and reanalyze. Multiply the concentration by the appropriate dilution factor.

Performance characteristics

Standard curve example

The following data were obtained for the various standards over the range of 0 to 100 Units/mL human EGFR [pY1068].

Standard Human EGFR [pY1068] (U/mL)	Optical Density (450 nm)
100	3.16
50	1.94
25	1.17
12.5	0.76
6.25	0.53
3.12	0.39
1.6	0.32
0	0.21

Recovery

The recovery of Hu EGFR [pY1068] Standard added to Cell Extraction Buffer and diluted 1:10 in Standard Diluent Buffer averaged 103%.

Inter-assay precision

Samples were assayed 48 times in multiple assays to determine precision between assays.

Parameters	Sample 1	Sample 2	Sample 3
Mean (U/mL)	40.55	11.95	2.49
Standard Deviation	1.74	0.68	0.23
% Coefficient of Variation	4.30	5.68	9.27

Intra-assay precision

Samples of known human EGFR [pY1068] concentration were assayed in replicates of 16 to determine precision within an assay.

Parameters	Sample 1	Sample 2	Sample 3
Mean (U/mL)	39.9	12.10	2.58
Standard Deviation	1.40	0.54	0.20
% Coefficient of Variation	3.51	4.49	7.58

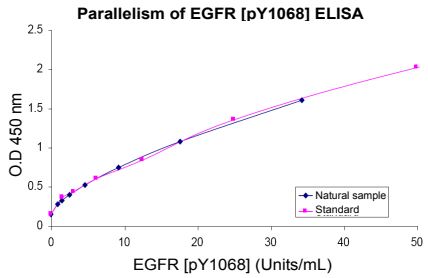
Linearity of dilution

Cell Extraction Buffer spiked with human EGFR [pY1068] was serially diluted in Standard Diluent Buffer over the range of the assay. Linear regression analysis of samples versus the expected concentration yielded a correlation coefficient of 0.99.

Dilution	Measured (U/mL)	Expected (U/mL)	% Expected
Neat	61.68	—	—
1/2	31.11	30.8	101
1/4	15.16	15.4	98
1/8	8.06	7.7	104
1/16	3.92	3.9	102

Parallelism

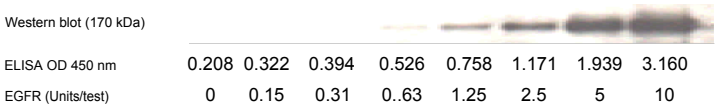
Natural human EGFR [pY1068] from EGF treated-A431 lysate was serially diluted in Standard Diluent Buffer. The optical density of each dilution was plotted against the EGFR [pY1068] standard curve. Parallelism demonstrates that the standard accurately reflects the full EGFR [pY1068] content in samples.



Sensitivity

The analytical sensitivity of this assay is <0.3 Units/mL human EGFR [pY1068] . This was determined by adding two standard deviations to the mean O.D. obtained from 30 assays of the zero standard.

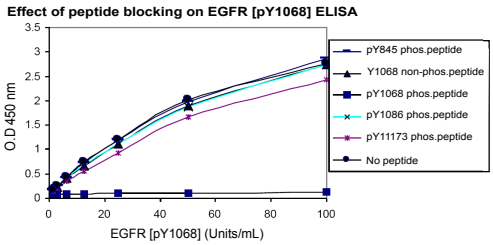
The sensitivity of this ELISA is at least 8-fold greater than that of western blot when tested against known quantities of EGFR [pY1068].



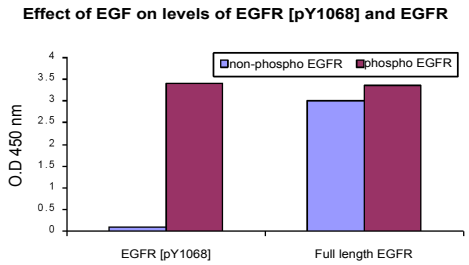
ELISA OD 450 nm	0.208	0.322	0.394	0.526	0.758	1.171	1.939	3.160
EGFR (Units/test)	0	0.15	0.31	0..63	1.25	2.5	5	10

Specificity

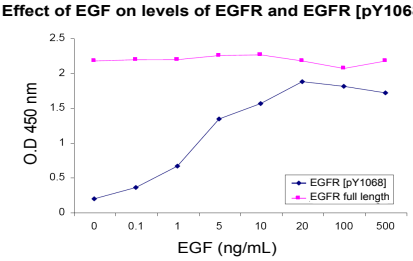
EGFR [pY1068] specificity was confirmed by peptide competition. The assay was performed as usual except that the detection antibody was preincubated with 1 µg/mL of EGFR-derived peptides. The data shows that only the peptide corresponding to the region surrounding tyrosine 1068 blocks the ELISA signal.



EGFR phosphorylation is dependent on EGF stimulation in HeLa cells. Human EGFR [pY1068] is detected at high level in EGF-stimulated HeLa cells, but not in unstimulated controls.



In parallel assays of HeLa cells treated with EGF (0 to 500 ng/mL) for 10 minutes, the amount of full length EGFR remains constant, while the level of EGFR [pY1068] decreases with diminishing EGF dosage.



Limited product warranty

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Product label explanation of symbols and warnings

REF	Catalog Number	LOT	Batch code		Temperature limitation		Use by		Manufacturer		Consult instructions for use		Caution, consult accompanying documents

Manufacturer's address: Bender MedSystems GmbH | Campus Vienna Biocenter 2 | 1030 Vienna, Austria

The information in this guide is subject to change without notice.