

---

## Certificate of Analysis

### Anti-Phosphotyrosine Immunoblotting Kit (4G10), ECL Detection Catalog # 17-153 Lot # 29750

#### Kit Components

**Anti-Phosphotyrosine, clone 4G10**, Catalog # 05-321, Lot # 28818. One vial containing **100µg** of protein G purified mouse monoclonal IgG<sub>2bκ</sub> in **100µl** of 0.1M Tris-glycine, pH 7.4, 0.15M NaCl, 0.05% sodium azide. Protein was determined by a Bradford microtiter protein assay. See page two for more information.

**EGF-Stimulated A431 Cell Lysate (Positive control)**, Catalog # 12-302. Two vials, each vial containing **100µg** in **100µl** of RIPA diluted with non-reducing sample buffer. Concentration: 1µg/µl. See page two for more information.

**Goat Anti-Mouse IgG, HRP-Conjugate**, Catalog # 12-349, Lot # 29574. One vial containing **500µg** of immunoaffinity purified goat IgG conjugated to horseradish peroxidase lyophilized from 0.02M Potassium Phosphate, 0.15M NaCl, pH 7.2, 10mg/ml BSA, and 0.01% gentamicin sulfate. See page two for more information. **Note: This product is light sensitive.**

**Blocking Reagent**, Catalog # 20-200. One vial containing **20g** nonfat dry milk fortified with vitamin A palmitate and vitamin D<sub>2</sub>, containing no preservatives.

**ECL Detection Reagent A**, Catalog # 17-153a, Lot # 29632. One vial containing **20ml**.

**ECL Detection Reagent B**, Catalog # 17-153b, Lot # 29635. One vial containing **20ml**.

**Autorad Orientation Markers**, Catalog # 20-135. One packet containing 5 markers. Use one-half marker per nitrocellulose blot.

**Note:** Markers are light sensitive but must be activated before use by exposing to light for 1-5 minutes.

**FOR RESEARCH USE ONLY  
NOT RECOMMENDED OR INTENDED FOR DIAGNOSIS OF DISEASE IN HUMANS OR ANIMALS  
DO NOT USE IN HUMANS**

---

#### Kit Description

**Quantity:** 10 immunoblots per kit.

**Quality Control:** The reagents in this kit have been matched to optimize the range and sensitivity of detection using EGF-stimulated human A431 cells as a source of phosphotyrosine containing proteins.

**Storage and Stability:** Stable for 6 months at 4°C from date of shipment. **Note:** Upon receipt, aliquot and store Catalog # 12-302, EGF-Stimulated A431 Cell Lysate and Catalog # 12-349, Goat Anti-Mouse IgG, at -20°C for optimal performance. Store Catalog # 05-321 at 4°C for optimal performance.

**Use:** Read the enclosed protocol before use.

## **Technical Information for Kit Components**

### **Anti-Phosphotyrosine, clone 4G10 (monoclonal IgG<sub>2bk</sub>)**

**Immunogen:** Phosphotyramine-KLH.

**Antibody Class:** IgG<sub>2bk</sub> mouse monoclonal antibody produced *in vitro* by mouse-mouse hybridoma 4G10 (FOX-NY [NS-1 derivative] myeloma x spleen cells). Purified by Protein G-agarose chromatography.

**Physical Form:** Liquid.

**Storage and Stability:** Stable for 2 years at 4°C from date of shipment. **NOTE: DO NOT FREEZE.** For maximum recovery of the product, centrifuge the original vial prior to removing the cap. If the product has accidentally been frozen and thawed, spin it at 13,000 x g for 10 minutes at 4°C. Save the supernatant for application.

---

#### **References:**

1. Cohen, B., *et al.*, Proc. Natl. Acad. Sci. USA **87**: 4458-4462, 1990.
2. Druker, B.J., *et al.*, New Eng. J. Med. **321**: 1383-1391, 1989.
3. Kanakura, Y., *et al.*, J. Biol. Chem. **266**: 490-495, 1991.

### **EGF-Stimulated A431 Cell Lysate (human carcinoma cell line)**

**Product Description:** Cellular protein preparation. Cells were cultured for 20 minutes in the presence of 50ng/ml EGF (Catalog # 01-107), lysed in modified RIPA buffer (50mM Tris-HCl, pH 7.4, 1% NP40, 0.25% sodium deoxycholate, 150mM NaCl, 1mM EDTA, 1mM PMSF, 1µg/ml aprotinin, 1µg/ml leupeptin, 1µg/ml pepstatin, 1mM Na<sub>3</sub>VO<sub>4</sub>, 1mM NaF) and **diluted with non-reducing sample buffer** (31mM Tris-HCl, pH 6.8, 5% glycerol, 1% SDS, 0.002% bromophenol blue).

**Physical Form:** Frozen solution.

**Storage and Stability:** Stable for 6 months at -20°C from date of shipment. For maximum recovery of product, centrifuge the original vial after thawing and prior to removing the cap. Aliquot to avoid repeated freezing and thawing.

**Note:** Add 2.5µl of 2-mercaptoethanol/100µl of lysate and boil for 5 minutes to reduce the preparation. Load 20µg of reduced lysate per lane for immunoblot analysis. This preparation may be used as a positive control for some of Upstate's antibodies.

### **Goat Anti-Mouse IgG, HRP-conjugated (goat polyclonal IgG)**

**Immunogen:** Highly purified whole mouse IgG.

**Specificity:** Recognizes mouse IgG, both heavy and light chains.

**Species Cross-reactivity:** Mouse.

**Rehydration:** Add 500µl of 50% glycerol to make a 1mg/ml stock solution. Aliquot to avoid repeated freezing and thawing.

**Storage and Stability:** Lyophilized: Stable for 2 years at 4°C from date of shipment. Rehydrated: Stable for 6 months at -20°C. Aliquot to avoid repeated freezing and thawing.

## Immunoblotting Analysis Protocol

**Safety Precautions:** Safety information for ECL Detection Reagents A and B is provided on the Material Safety Data Sheets (MSDS) enclosed with the kit components. Read carefully prior to use.

### Reagent Preparation:

1. **Blocking Buffer:** Dissolve 0.9g of "Blocking Reagent," Catalog # 20-200 in 30ml of phosphate buffered saline (PBS), pH 7.4.
2. **Primary Antibody Solution:** Add 10-20 $\mu$ g of Anti-Phosphotyrosine Antibody, Catalog # 05-321, to 10ml of **freshly** prepared Blocking Buffer. Mix well and store at 2-8°C.  
**Note: This solution can be reused ONCE within one week.**
3. **Secondary Antibody Solution:** Add 10 $\mu$ l of Goat Anti-Mouse IgG HRP-Conjugate, Catalog # 12-349, to 10ml of Blocking Buffer. Mix well and store in the dark at 2-8°C, or prepare immediately before use.

### Procedure:

#### **Prior to Immunodetection:**

1. Prepare the samples and "Positive Control" (Catalog # 12-302) for electrophoresis and immunoblotting. Load the SDS-PAGE gel with the samples to be tested and 10 $\mu$ l of the Positive Control.  
**Note: A positive control should be included with every analysis.**
2. Perform electrophoresis and blot to nitrocellulose filter paper.
3. Wash the nitrocellulose filter two times with fresh changes of PBS (phosphate buffered saline) or distilled water.

#### **Immunodetection:**

1. Block the blotted nitrocellulose filter by immersing in 10ml of **freshly** prepared Blocking Buffer for 30 minutes at room temperature.
2. Incubate the nitrocellulose filter in the "Primary Antibody Solution" overnight at 4°C. **The solution can be saved for one additional use!!**
3. Wash the nitrocellulose filter two times with fresh changes of PBS or distilled water.
4. Incubate the nitrocellulose filter in the "Secondary Antibody Solution" for one and one-half hours at room temperature.
5. Wash the nitrocellulose filter two times with fresh changes of PBS or distilled water.
6. Wash the nitrocellulose filter in PBS-0.05% Tween 20 for 5 minutes with constant rocking.
7. Rinse the nitrocellulose filter four-five times with fresh changes of PBS or distilled water.
8. Lay the nitrocellulose on a transparency sheet, Blot excess water from nitrocellulose membrane with paper towel being careful not to allow blot to dry out.
9. Combine 2ml ECL Detection Reagent A (Catalog # 17-153a) and 2ml ECL Detection Reagent B (Catalog # 17-153b). Mix thoroughly.

**Immunodetection:**

10. Cover the nitrocellulose, using a pipet, with a uniform layer of the ECL Detection Reagent mixture.
11. Incubate for precisely 1 minute at room temperature.
12. Drain and wick the excess ECL Detection Reagent mixture with a paper towel, making sure that the nitrocellulose does not dry out.
13. Place an Autorad orientation marker (Catalog # 20-135) next to the nitrocellulose.
14. Cover the nitrocellulose with a second transparency sheet being careful to remove all bubbles on the nitrocellulose. **Note:** Expose as soon as possible, may be stored in the dark for up to 30 minutes.
15. Place the transparency covered nitrocellulose in a film cassette, cover entirely with a piece of film and close the cassette securely.
16. Start with a 40 second exposure, remove the film from the cassette and develop. Re-exposure for longer or shorter periods may be necessary depending on intensity of staining.
17. Develop the exposed film for 2 minutes in developer, rinse for 30 seconds in water and fix for 4 minutes in fixer.