## Anti-Phospho-MUC1 (Y1229) Rabbit Polyclonal Antibody



### **Catalog #: U0243**

#### **Aliases**

PUM; Mucin-1; MUC-1; Breast carcinoma-associated antigen DF3; Cancer antigen 15-3; CA 15-3; Carcinoma-associated mucin; Episialin; H23AG; Krebs von den Lungen-6; KL-6; PEMT; Peanut-reactive urinary mucin; PUM; Polymorphic epithelial mucin; PEM; Tumor-associated epithelial membrane antigen; EMA; Tumor-associated mucin; CD227

### **Background**

Gene Name: MUC1 NCBI Gene Entry: 4582 UniProt Entry: P15941

### **Application Information**

Molecular Weight: Predicted, 122 kDa; observed, 170 kDa

Clonality: Rabbit polyclonal antibody Species Reactivity: Human, mouse, rat

Applications Tested: Western blotting (WB), immunohistochemistry (IHC), immunocytochemistry

(IC)

#### **Immunogen**

A synthesized peptide derived from human Phospho-MUC1 (Y1229)

### **Isotype**

Rabbit IgG

### **Storage Buffer**

Supplied in PBS (pH 7.3) containing 30% glycerol, and 0.01% sodium azide.

#### **Storage**

Store at -20 °C for one year.

#### **Recommended Dilutions**

Western Blotting (WB): 1:500-1:1,000 Immunohistochemistry (IHC): 1:50-1:100 Immunocytochemistry (IC): 1:50-1:200

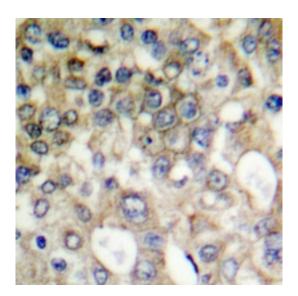
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**Note:** This product is for research use only.

#### **Validation Data**



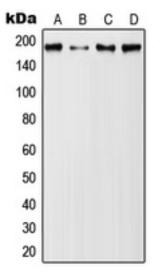
Immunocytochemical analysis of MUC1 (Phospho-Y1229) staining in HepG2 cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a hidified chamber. Cells were washed with PBST and incubated with a AF594-conjugated secondary antibody (red) in PBS at room temperature in the dark.



Immunohistochemical analysis of MUC1 (Phospho-Y1229) staining in human breast cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and

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mounted with DPX.



Western blotting analysis of MUC1 (Phospho-Y1229) expression in MCF7 PMA-treated (A), BT20 (B), NIH3T3 UV-treated (C), H9C2 PMA-treated (D) whole cell lysates.