

## Cytokeratin 7 Recombinant Rabbit Monoclonal Antibody Product Datasheet

Catalog# BX00095

Clone# RR699

**Predicted Molecular Wt:** 51kDa

**Purity:** ProA affinity purified IgG

**Species Cross-reactivity:** Human

**Form:** Liquid

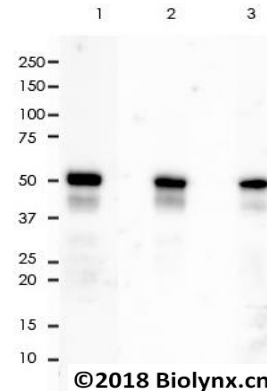
*Species cross-reactivity determined by WB*

**Swissprot ID:** P08729

**Applications:** WB IHC-P IF/ICC FC IP

### Background:

Blocks interferon-dependent interphase and stimulates DNA synthesis in cells. Involved in the translational regulation of the human papillomavirus type 16 E7 mRNA.



All lanes: Anti-Cytokeratin 7 antibody at 1:10,000 dilution

### Immunogen:

A synthetic peptide corresponding to residues aa1-100 of Cytokeratin 7 was used as an immunogen.

Predicted MW: 51 kDa  
 Observed MW: 51 kDa

Lysates at 10 µg per lane  
 2nd Ab:  
 GAR HRP(H+L) 1:5,000

### Storage Buffer:

PBS 59%, Sodium azide 0.01%, Glycerol 40%, BSA 0.05%.

Lane 1: HeLa  
 Lane 2: A549  
 Lane 3: HaCat

Exposure: 60s

### Storage conditions:

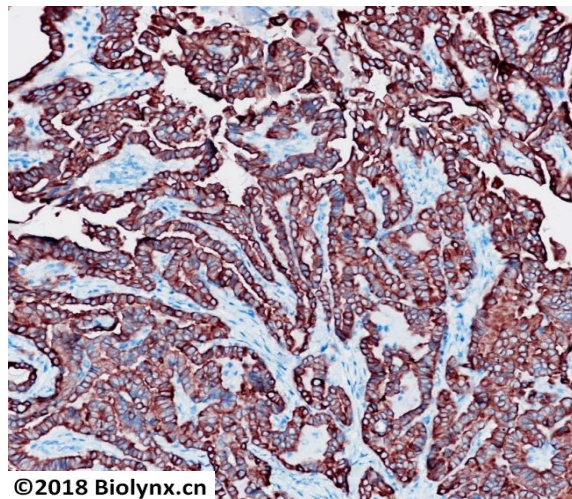
-20°C.

### Storage instructions:

Shipped on blue ice. Upon delivery, aliquot, and store at -20°C. Avoid freeze / thaw cycles.

### Recommended Dilutions:

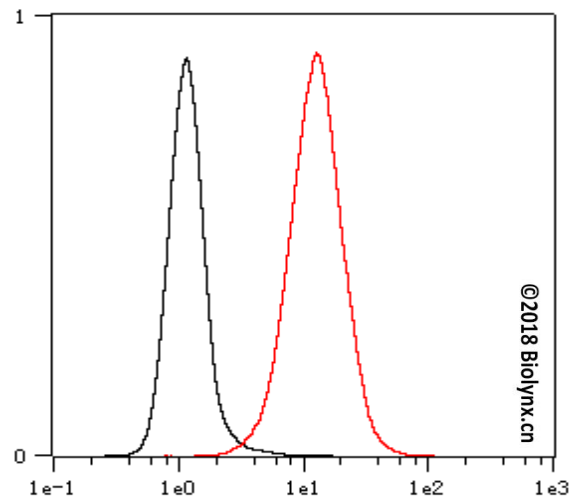
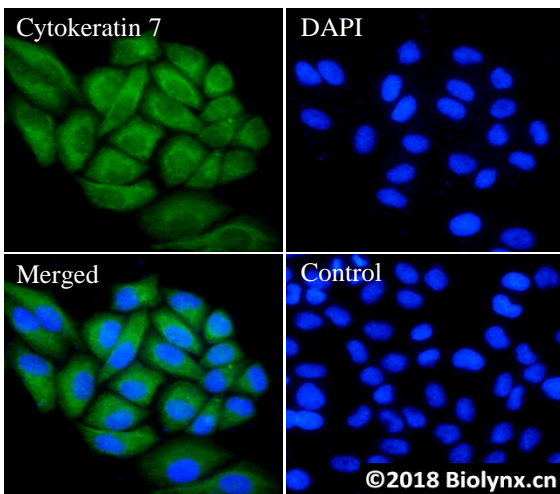
WB: 1:2,000 - 1:5,000  
 IHC-P: 1:1,600 - 1:3,200  
 IF/ICC: 1:800 - 1:2,000  
 FC: 1:200 - 1:800  
 IP: 1:50



### Background References:

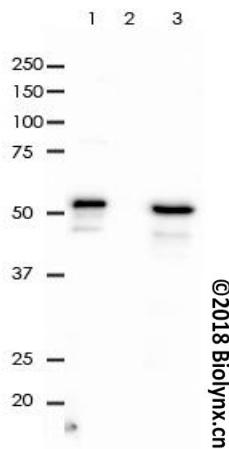
- Ambade A et al. Sci Rep 6:21340 (2016).
- Liu KM et al. Am J Physiol Renal Physiol 309:F318-31 (2015).

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human thyroid cancer tissue labelling Cytokeratin 7 with RR699 at 1:3,200. Heat mediated antigen retrieval was performed using Tris/EDTA buffer pH 9.0.



RR699 staining Cytokeratin 7 in HeLa cells by IF/ICC (immunofluorescence/immunocytochemistry). Cells were fixed with paraformaldehyde, permeabilized with 0.1% Triton X-100 and blocked with 10% goat serum for half an hour at room temperature. Samples were incubated with primary antibody (1:2,000) at 4°C. An Alexa Fluor® 488-conjugated Goat Anti-Rabbit IgG polyclonal was used as the secondary antibody (1:500). DAPI (blue) was used as the nuclear counter stain.  
 Control: PBS and secondary antibody, An Alexa Fluor® 488-conjugated Goat Anti-Rabbit IgG (1:500).

Overlay histogram showing HeLa cells stained with RR699 (Red). The cells were fixed with 4% paraformaldehyde (10 min) and then permeabilized with 0.1% TritonX-100 for 15 min. The cells were then incubated in the antibody (RR699, 1:800 dilution) in 1x PBS/1% BSA for 30 min at room temperature. The secondary antibody used was a Goat Anti-Rabbit Alexa Fluor® 488 (IgG H+L) at 1:2,000 dilution for 20 min at room temperature. Unlabelled sample (Black) was used as a control.



Cytokeratin 7 was immunoprecipitated from 0.4mg of HeLa whole cell lysate with RR699 at 1:50 dilution.  
 2nd Ab:  
 GAR HRP for IP 1:500

Lane 1: RR699 IP in HeLa whole cell lysate  
 Lane 2: PBS instead of RR699 in HeLa whole cell lysate  
 Lane 3: HeLa whole cell lysate, 10 µg (input)

Exposure: 50s

Product QC'd by:



For research use only. Not for use in diagnostic or therapeutic applications.