

## **Product Data Sheet**

## **Anti-Cytokeratin 17 Antibody**

Catalog # Source Reactivity Applications

CPA9804 Mouse H WB, IH

**Description** Mouse monoclonal antibody to Cytokeratin 17

Immunogen KLH-conjugated synthetic peptide encompassing a sequence within human

Cytokeratin 17. The exact sequence is proprietary.

**Purification** The antibody was purified by immunogen affinity chromatography.

**Specificity** Recognizes endogenous levels of Cytokeratin 17 protein.

**Clonality** Monoclonal

**Conjugation** 

Form Mouse IgG1. Liquid in PBS containing 50% glycerol, 0.2% BSA and 0.01% sodium

azide.

**Dilution** WB (1/500 - 1/1000), IH (1/100 - 1/300)

Gene Symbol KRT17

Alternative Names Keratin type I cytoskeletal 17; 39.1; Cytokeratin-17; CK-17; Keratin-17; K17

Entrez Gene 3872 (Human)

SwissProt Q04695 (Human)

Storage/Stability Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid

freeze/thaw cycles.

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

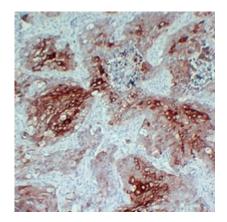
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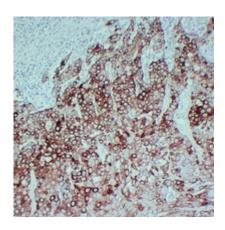




## **Product Data Sheet**



Immunohistochemical analysis of Cytokeratin 17 staining in human lung squamous carcinoma 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 wa



Immunohistochemical analysis of Cytokeratin 17 staining in human oral squamous cell carcinoma 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 mounted with DPX.

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