

## Keratin 17 (Phospho Ser44) rabbit pAb

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| <b>Catalog No :</b>          | YP1377  |
| <b>Reactivity :</b>          | Human;Rat;Mouse;  |
| <b>Applications :</b>        | WB;ELISA;IHC  |
| <b>Target :</b>              | Cytokeratin 17  |
| <b>Fields :</b>              | >>Estrogen signaling pathway;>>Staphylococcus aureus infection                  |
| <b>Gene Name :</b>           | KRT17   |
| <b>Protein Name :</b>        | Keratin 17 (Ser44)  |
| <b>Human Gene Id :</b>       | 3872  |
| <b>Human Swiss Prot No :</b> | Q04695  |
| <b>Mouse Gene Id :</b>       | 16667   |
| <b>Mouse Swiss Prot No :</b> | Q9QWL7  |
| <b>Rat Gene Id :</b>         | 287702  |
| <b>Rat Swiss Prot No :</b>   | Q6IFU8  |
| <b>Immunogen :</b>           | Synthesized phosho peptide around human Keratin 17 (Ser44)                      |
| <b>Specificity :</b>         | This antibody detects endogenous levels of Human Keratin 17 (phospho-Ser44)     |
| <b>Formulation :</b>         | Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.         |
| <b>Source :</b>              | Polyclonal, Rabbit,IgG  |
| <b>Dilution :</b>            | WB 1:500-2000;IHC 1:50-300; ELISA 2000-20000                                    |
| <b>Purification :</b>        | The antibody was affinity-purified from rabbit serum by affinity-chromatography |

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using specific immunogen.

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**Concentration :** 1 mg/ml

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**Storage Stability :** -15°C to -25°C/1 year (Do not lower than -25°C)

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**Observed Band :** 48kD

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**Background :** This gene encodes the type I intermediate filament chain keratin 17, expressed in nail bed, hair follicle, sebaceous glands, and other epidermal appendages. Mutations in this gene lead to Jackson-Lawler type pachyonychia congenita and steatocystoma multiplex. [provided by RefSeq, Aug 2008],

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**Function :** disease:Defects in KRT17 are a cause of pachyonychia congenita type 2 (PC2) [MIM:167210]; also known as pachyonychia congenita Jackson-Lawler type. PC2 is an autosomal dominant ectodermal dysplasia characterized by hypertrophic nail dystrophy resulting in onychogryposis (thickening and increase in curvature of the nail), palmoplantar keratoderma and hyperhidrosis, follicular hyperkeratosis, multiple epidermal cysts, absent/sparse eyebrow and body hair, and by the presence of natal teeth.,disease:Defects in KRT17 are a cause of steatocystoma multiplex (SM) [MIM:184500]. SM is a disease characterized by round or oval cystic tumors widely distributed on the back, anterior trunk, arms, scrotum, and thighs.,disease:KRT16 and KRT17 are coexpressed only in pathological situations such as metaplasias and carcinomas of the uterine cervix and in psoriasis vulgaris.,function:May play a role in the

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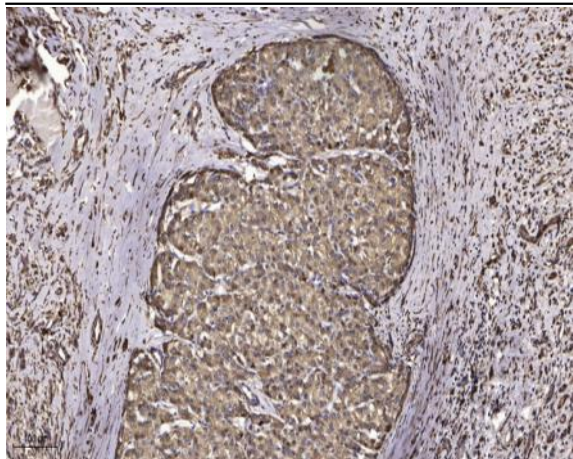
**Subcellular Location :** Cytoplasm .

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**Expression :** Expressed in the outer root sheath and medulla region of hair follicle specifically from eyebrow and beard, digital pulp, nail matrix and nail bed epithelium, mucosal stratified squamous epithelia and in basal cells of oral epithelium, palmoplantar epidermis and sweat and mammary glands. Also expressed in myoepithelium of prostate, basal layer of urinary bladder, cambial cells of sebaceous gland and in exocervix (at protein level).

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## Products Images



Immunohistochemical analysis of paraffin-embedded human liver cancer. 1, Antibody was diluted at 1:200(4° overnight). 2, Tris-EDTA,pH9.0 was used for antigen retrieval. 3,Secondary antibody was diluted at 1:200(room temperature, 45min).