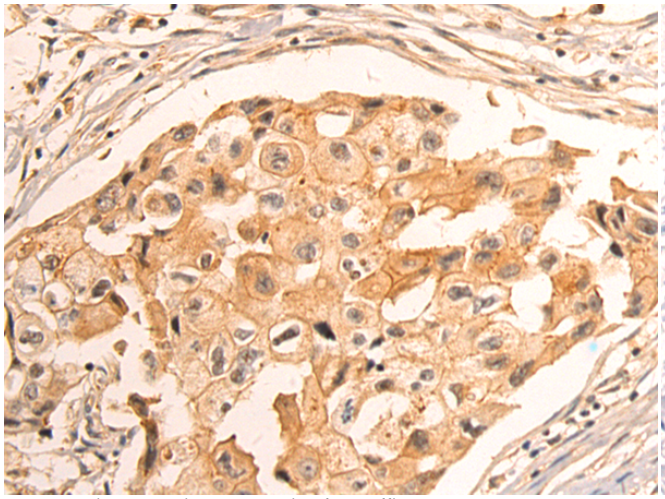
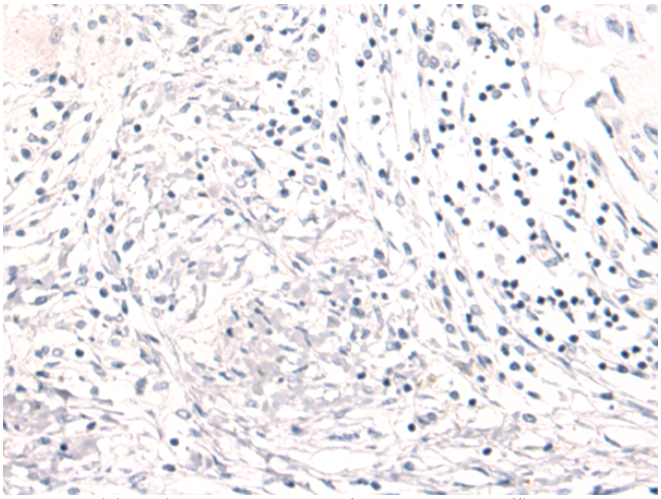
SLC26A4 RABBIT PAB

货号: S221132
产品全名: SLC26A4 兔多抗
基因符号 EVA; PDS; DFNB4; TDH2B
UNIPROT ID: O43511 (Gene Accession – NP_000432)
背景: Mutations in this gene are associated with Pendred syndrome, the most common form of syndromic deafness, an autosomal-recessive disease. It is highly homologous to the SLC26A3 gene; they have similar genomic structures and this gene is located 3' of the SLC26A3 gene. The encoded protein has homology to sulfate transporters.
抗原: Synthetic peptide of human SLC26A4
经过测试的应用: ELISA, IHC
推荐稀释比: IHC: 50-200; ELISA: 5000-10000
种属反应性: Rabbit
克隆性： Rabbit Polyclonal
亚型: Immunogen-specific rabbit IgG
纯化: Antigen affinity purification
种属反应性: Human, Mouse, Rat
成分: PBS (without Mg2+ and Ca2+), pH 7.4, 150 mM NaCl, 0.05% Sodium Azide and 40% glycerol
研究领域: Neuroscience, Signal Transduction, Metabolism
储存和运输: Store at -20°C. Avoid repeated freezing and thawing



Immunohistochemistry analysis of paraffin embedded Human breast cancer tissue using 221132(SLC26A4 Antibody) at a dilution of 1/50(Cell membrane).



In comparision with the IHC on the left, the same paraffin-embedded Human breast cancer tissue is first treated with the synthetic peptide and then with 221132(Anti-SLC26A4 Antibody) at dilution 1/50.