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Anti-Human SLC26A4 Polyclonal Antibody

多克隆抗体

产品货号: PA11034

第三版

描述: 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 SLC25A4

配方:

如何使用: 加1ml超纯水重溶

稳定性: -20 ° C保存条件下，冻干粉,保质期为五年；液体，保质期为两年。

稀释液: PBS (pH7.4) ， 1% BSA

应用: WB 1 ~ 5 μ g/ml.

特异性: High expression in adult thyroid, lower expression in adult and fetal kidney and fetal brain. Not expressed in other tissues.