本公司提供的电子版本说明书仅供参考，实验请以收到的纸质手册为准。

Anti-Human SLC26A4 Polyclonal Antibody

多克隆抗体

产品货号: PA08136

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.