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 Alpha galactosidase A Monoclonal Antibody

单克隆抗体

产品货号: MA01851

第三版

免疫原: Fusion protein of Alpha galactosidase A

亚型: IgG2a

克隆:407

描述:GLA,also named as Melibiase,Agalsidase and Alpha-galactosidase A,belongs to the glycosyl hydrolase 27 family. It hydrolyzes terminal,non-reducing alpha-D-galactose residues in alpha-D-galactosides,including galactose oligosaccharides,galactomannans and galactolipids. Fabry disease is an X-linked lysosomal storage disorder resulting from the deficient activity of GLA. Enzyme replacement therapy (ERT) with GLA is currently the most effective therapeutic strategy for patients with Fabry disease,a lysosomal storage disease.

配方:

浓度: None

保存: +4 ° C 保存 (1-2 周). 长期保存在-20 ° C或者-70 ° C. 避免反复冻融.

应用:ELISA, WB, ICC/IF