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Anti-Human ABHD11 Polyclonal Antibody

多克隆抗体

产品货号: PA02951

第三版

描述: This gene encodes a protein containing an alpha/beta hydrolase fold domain. This gene is deleted in Williams syndrome, a multisystem developmental disorder caused by the deletion of contiguous genes at 7q11.23. ABHD11 (Abhydrolase Domain Containing 11) is a Protein Coding gene. Diseases associated with ABHD11 include Williams-Beuren Syndrome. GO annotations related to this gene include hydrolase activity and hydrolase activity, acting on ester bonds.

抗原: Synthesized peptide derived from the Internal region of human ABHD11

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: Ubiquitously expressed.