Anti-Human ABHD11 Polyclonal Antibody

Polyclonal Antibody

Cat.NO.: PA02951

3th Edition

Description: 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.

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

Form:

How to use:1.0 ml distilled water will be added to the product

Stability: Lyophilized product, 5 years at 2 – 8°C; Solution, 2 years at –20°C

Dilution:PBS (pH7.4) containing 1% BSA

Application: This antibody can be used for western blotting in concentration of 1?5?g/ml.

Specificity: Ubiquitously expressed.