ABHD11 Antibody



PACO22109

Reactivity:

Product Information

Size: Protein Background:

100ul 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. Alternatively spliced transcript variants have

Human been described, but their biological validity has not been determined.

Source: Gene ID:

Rabbit ABHD11

Isotype: Uniprot

IgG Q8NFV4

Applications: Synonyms:

ELISA, WB ABHDB; abhydrolase domain containing 11; EC 3. -. -. -; PP1226; WBSCR21

Recommended dilutions: Immunogen:

ELISA:1:2000-1:10000, WB:1:500-1:3000 Synthesized peptide derived from internal of human ABHD11.

Storage:

Rabbit IgG in phosphate buffered saline (without Mg2+ and Ca2+), pH 7.4, 150mM

NaCl, 0.02% sodium azide and 50% glycerol.

Product Images

