

DNAXPAb

Hard-to-Find
Antibody

ABHD11 DNAXPab

Catalog # H00083451-W01P

Size 200 ug

Specification

Product Description	Rabbit polyclonal antibody raised against a full-length human ABHD11 DNA using DNAX™ Immune technology.
Technology	DNAX™ Immune
Immunogen	Full-length human DNA
Sequence	MRAGQQLASMLRWTRAWRLPREGLGPHGPSFARVPVAPSSSSGGRGGAEPRLPLSYRLLDG EAALPAVVFLHGLFGSKTNFNSIAKILAAQTGRRVLTVDARNHGDSPHSPDMSYEIMSQDLQDLLP QLGLVPCVVVGHSMMGGKTAMLLALQRPELVERLIAVDISPVESTGVSHFATYVAAMRAINADELP RSRARKLADEQLSSVIQDMAVRQHLLTNLVEVDGRFVWRVNLDAQTQHLDKILAFPQRQESYLG P TLFLLGGNSQFVHPSHHPEIMRLFPRMQMTPVNPAGHWIHADRPQDFIAAIRGFLV
Host	Rabbit
Reactivity	Human
Purification	Protein A
Quality Control Testing	Antibody reactive against mammalian transfected lysate.
Storage Buffer	In 1x PBS, pH 7.4
Storage Instruction	Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

Applications

- Western Blot (Transfected lysate)

[Protocol Download](#)

- Immunofluorescence (Transfected cell)
- Flow Cytometry (Transfected cell)

Gene Info — ABHD11

Entrez GeneID [83451](#)**GeneBank Accession#** [NM_148912.2](#)**Protein Accession#** [NP_683710.1](#)**Gene Name** ABHD11**Gene Alias** PP1226, WBSCR21**Gene Description** abhydrolase domain containing 11**Gene Ontology** [Hyperlink](#)

Gene Summary 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 been described, but their biological validity has not been determined. [provided by RefSeq]

Other Designations Williams Beuren syndrome chromosome region 21