

# **Product datasheet for TP325443L**

### OriGene Technologies, Inc.

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## ASPA (NM\_001128085) Human Recombinant Protein

**Product data:** 

**Product Type:** Recombinant Proteins

**Description:** Recombinant protein of human aspartoacylase (Canavan disease) (ASPA), transcript variant 2,

1 mg

Species: Human
Expression Host: HEK293T

**Expression cDNA Clone** >RC225443 protein sequence or AA Sequence: Red=Cloning site Green=Tags(s)

MTSCHIAEEHIQKVAIFGGTHGNELTGVFLVKHWLENGAEIQRTGLEVKPFITNPRAVKKCTRYIDCDLN RIFDLENLGKKMSEDLPYEVRRAQEINHLFGPKDSEDSYDIIFDLHNTTSNMGCTLILEDSRNNFLIQMF HYIKTSLAPLPCYVYLIEHPSLKYATTRSIAKYPVGIEVGPQPQGVLRADILDQMRKMIKHALDFIHHFN EGKEFPPCAIEVYKIIEKVDYPRDENGEIAAIIHPNLQDQDWKPLHPGDPMFLTLDGKTIPLGGDCTVYP

VFVNEAAYYEKKEAFAKTTKLTLNAKSIRCCLH

**TRTRPLEQKLISEEDLAANDILDYKDDDDKV** 

Tag: C-Myc/DDK

Predicted MW: 35.6 kDa

Concentration:  $>0.05 \mu g/\mu L$  as determined by microplate BCA method

**Purity:** > 80% as determined by SDS-PAGE and Coomassie blue staining

**Buffer:** 25 mM Tris-HCl, 100 mM glycine, pH 7.3, 10% glycerol

**Preparation:** Recombinant protein was captured through anti-DDK affinity column followed by

conventional chromatography steps.

**Note:** For testing in cell culture applications, please filter before use. Note that you may experience

some loss of protein during the filtration process.

Storage: Store at -80°C.

Stability: Stable for 12 months from the date of receipt of the product under proper storage and

handling conditions. Avoid repeated freeze-thaw cycles.

**RefSeg:** NP 001121557

Locus ID: 443



#### ASPA (NM\_001128085) Human Recombinant Protein - TP325443L

UniProt ID: <u>P45381</u>, <u>Q6FH48</u>

RefSeq Size: 1368
Cytogenetics: 17p13.2
RefSeq ORF: 939

Synonyms: ACY2; ASP

Summary: This gene encodes an enzyme that catalyzes the conversion of N-acetyl\_L-aspartic acid (NAA)

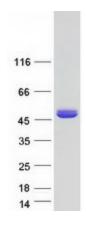
to aspartate and acetate. NAA is abundant in the brain where hydrolysis by aspartoacylase is thought to help maintain white matter. This protein is an NAA scavenger in other tissues. Mutations in this gene cause Canavan disease. Alternatively spliced transcript variants have

been found for this gene. [provided by RefSeq, Jul 2008]

**Protein Families:** Druggable Genome

**Protein Pathways:** Alanine, aspartate and glutamate metabolism, Histidine metabolism

## **Product images:**



Coomassie blue staining of purified ASPA protein (Cat# [TP325443]). The protein was produced from HEK293T cells transfected with ASPA cDNA clone (Cat# [RC225443]) using MegaTran 2.0

(Cat# [TT210002]).