

# **Product datasheet for PH312019**

## OriGene Technologies, Inc.

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#### WIPF1 (NM 001077269) Human Mass Spec Standard

**Product data:** 

**Product Type:** Mass Spec Standards

**Description:** WIPF1 MS Standard C13 and N15-labeled recombinant protein (NP\_001070737)

Species:HumanExpression Host:HEK293

Expression cDNA Clone

RC212019

or AA Sequence:

**Protein Sequence:** 

Predicted MW: 51.1 kDa

Red=Cloning site Green=Tags(s)

>RC212019 representing NM\_001077269

RERGAPPLPPIPR

TRTRPLEQKLISEEDLAANDILDYKDDDDKV

Tag: C-Myc/DDK

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

**Concentration:** >0.05 μg/μL as determined by microplate BCA method

Labeling Method: Labeled with [U- 13C6, 15N4]-L-Arginine and [U- 13C6, 15N2]-L-Lysine

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

Storage: Store at -80°C. Avoid repeated freeze-thaw cycles.

**Stability:** Stable for 3 months from receipt of products under proper storage and handling conditions.

**RefSeq:** NP 001070737

RefSeq Size: 4664 RefSeq ORF: 1509





### WIPF1 (NM\_001077269) Human Mass Spec Standard - PH312019

Synonyms: PRPL-2; WAS2; WASPIP; WIP

**Locus ID:** 7456

UniProt ID: <u>043516</u>, <u>A0A140V|Z9</u>, <u>Q2YDC4</u>

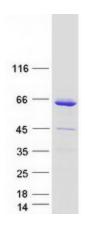
**Cytogenetics:** 2q31.1

**Summary:** This gene encodes a protein that plays an important role in the organization of the actin

cytoskeleton. The encoded protein binds to a region of Wiskott-Aldrich syndrome protein that is frequently mutated in Wiskott-Aldrich syndrome, an X-linked recessive disorder. Impairment of the interaction between these two proteins may contribute to the disease. Two

transcript variants encoding the same protein have been identified for this gene. [provided by RefSeq, Jul 2008]

# **Product images:**



Coomassie blue staining of purified WIPF1 protein (Cat# [TP312019]). The protein was produced from HEK293T cells transfected with WIPF1 cDNA clone (Cat# [RC212019]) using MegaTran 2.0 (Cat# [TT210002]).