

## **Product datasheet for PH309596**

## OriGene Technologies, Inc.

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## RD3 (NM\_183059) Human Mass Spec Standard

**Product data:** 

**Product Type:** Mass Spec Standards

**Description:** RD3 MS Standard C13 and N15-labeled recombinant protein (NP\_898882)

Species:HumanExpression Host:HEK293

Expression cDNA Clone

RC209596

or AA Sequence: Predicted MW:

22.7 kDa

Protein Sequence: >RC209596 protein sequence

Red=Cloning site Green=Tags(s)

MSLISWLRWNEAPSRLSTRSPAEMVLETLMMELTGQMREAERQQRERSNAVRKVCTGVDYSWLASTPRST YDLSPIERLQLEDVCVKIHPSYCGPAILRFRQLLAEQEPEVQEVSQLFRSVLQEVLERMKQEEEAHKLTR

QWSLRPRGSLATFKTRARISPFASDIRTISEDVERDTPPPLRSWSMPEFRAPKAD

TRTRPLEQKLISEEDLAANDILDYKDDDDKV

Tag: C-Myc/DDK

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

Concentration:  $>0.05 \mu g/\mu L$  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 898882

RefSeq Size: 4290 RefSeq ORF: 585

Synonyms: C1orf36; LCA12

 Locus ID:
 343035

 UniProt ID:
 Q7Z3Z2





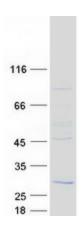
Cytogenetics:

1q32.3

**Summary:** 

This gene encodes a retinal protein that is associated with promyelocytic leukemia-gene product (PML) bodies in the nucleus. Mutations in this gene cause Leber congenital amaurosis type 12, a disease that results in retinal degeneration. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Sep 2009]

## **Product images:**



Coomassie blue staining of purified RD3 protein (Cat# [TP309596]). The protein was produced from HEK293T cells transfected with RD3 cDNA clone (Cat# [RC209596]) using MegaTran 2.0 (Cat# [TT210002]).