

## **Product datasheet for PH300757**

## OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

## SOX2 (NM\_003106) Human Mass Spec Standard

**Product data:** 

**Product Type:** Mass Spec Standards

**Description:** SOX2 MS Standard C13 and N15-labeled recombinant protein (NP\_003097)

Species:HumanExpression Host:HEK293

Expression cDNA Clone

RC200757

or AA Sequence: Predicted MW:

34.3 kDa

Protein Sequence: >RC200757 protein sequence

Red=Cloning site Green=Tags(s)

MYNMMETELKPPGPQQTSGGGGNSTAAAAGGNQKNSPDRVKRPMNAFMVWSRGQRRKMAQENPKMHNSE ISKRLGAEWKLLSETEKRPFIDEAKRLRALHMKEHPDYKYRPRRKTKTLMKKDKYTLPGGLLAPGGNSMA SGVGVGAGLGAGVNQRMDSYAHMNGWSNGSYSMMQDQLGYPQHPGLNAHGAAQMQPMHRYDVSALQYNSM TSSQTYMNGSPTYSMSYSQQGTPGMALGSMGSVVKSEASSSPPVVTSSSHSRAPCQAGDLRDMISMYLPG

AEVPEPAAPSRLHMSQHYQSGPVPGTAINGTLPLSHM

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 003097

RefSeq Size: 2520 RefSeq ORF: 951

**Synonyms:** ANOP3; MCOPS3

**Locus ID:** 6657





UniProt ID: P48431, A0A0U3FYV6

Cytogenetics: 3q26.33

**Summary:** This intronless gene encodes a member of the SRY-related HMG-box (SOX) family of

> transcription factors involved in the regulation of embryonic development and in the determination of cell fate. The product of this gene is required for stem-cell maintenance in the central nervous system, and also regulates gene expression in the stomach. Mutations in

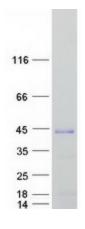
this gene have been associated with optic nerve hypoplasia and with syndromic

microphthalmia, a severe form of structural eye malformation. This gene lies within an intron of another gene called SOX2 overlapping transcript (SOX2OT). [provided by RefSeq, Jul 2008]

**Protein Families:** Adult stem cells, Cancer stem cells, Embryonic stem cells, ES Cell Differentiation/IPS, Induced

pluripotent stem cells, Transcription Factors

## **Product images:**



Coomassie blue staining of purified SOX2 protein (Cat# [TP300757]). The protein was produced from HEK293T cells transfected with SOX2 cDNA clone (Cat# [RC200757]) using MegaTran 2.0 (Cat# [TT210002]).