

Product datasheet for PH315021

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

FOXP2 (NM_014491) Human Mass Spec Standard

Product data:

Product Type: Mass Spec Standards

Description: FOXP2 MS Standard C13 and N15-labeled recombinant protein (NP_055306)

Species:HumanExpression Host:HEK293

Expression cDNA Clone

RC215021

or AA Sequence: Predicted MW:

79.9 kDa

Protein Sequence: >RC215021 representing NM_014491

Red=Cloning site Green=Tags(s)

DDREIEEEPLSEDLE

TRTRPLEOKLISEEDLAANDILDYKDDDDKV

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 055306



FOXP2 (NM_014491) Human Mass Spec Standard - PH315021

RefSeq Size: 6373 RefSeq ORF: 2145

Synonyms: CAGH44; SPCH1; TNRC10

 Locus ID:
 93986

 UniProt ID:
 015409

 Cytogenetics:
 7q31.1

Summary: This gene encodes a member of the forkhead/winged-helix (FOX) family of transcription

factors. It is expressed in fetal and adult brain as well as in several other organs such as the

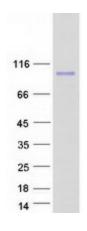
lung and gut. The protein product contains a FOX DNA-binding domain and a large polyglutamine tract and is an evolutionarily conserved transcription factor, which may bind directly to approximately 300 to 400 gene promoters in the human genome to regulate the expression of a variety of genes. This gene is required for proper development of speech and language regions of the brain during embryogenesis, and may be involved in a variety of biological pathways and cascades that may ultimately influence language development. Mutations in this gene cause speech-language disorder 1 (SPCH1), also known as autosomal dominant speech and language disorder with orofacial dyspraxia. Multiple alternative

transcripts encoding different isoforms have been identified in this gene.[provided by RefSeq,

Feb 2010]

Protein Families: Transcription Factors

Product images:



Coomassie blue staining of purified FOXP2 protein (Cat# [TP315021]). The protein was produced from HEK293T cells transfected with FOXP2 cDNA clone (Cat# [RC215021]) using MegaTran 2.0 (Cat# [TT210002]).