

## **Product datasheet for TP762353**

## OriGene Technologies, Inc.

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## FOXC1 (NM\_001453) Human Recombinant Protein

**Product data:** 

**Product Type:** Recombinant Proteins

**Description:** Purified recombinant protein of Human forkhead box C1 (FOXC1), Pro197-Gln306, with N-

terminal His tag, expressed in E.coli, 50ug

Species: Human
Expression Host: E. coli

Expression cDNA Clone

or AA Sequence:

A DNA sequence encoding the region(Pro197-Gln306) of FOXC1

Tag: N-His

**Predicted MW:** 10.7 kDa

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

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

**Buffer:** 25 mM Tris-HCl, pH 8.0, 150 mM NaCl, 10% glycerol

**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 after receiving vials.

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

handling conditions. Avoid repeated freeze-thaw cycles.

RefSeq: NP 001444

**Locus ID:** 2296

UniProt ID: <u>Q12948</u>, <u>W6CJ52</u>

RefSeq Size: 3452 Cytogenetics: 6p25.3 RefSeq ORF: 1659

Synonyms: ARA; ASGD3; FKHL7; FREAC-3; FREAC3; IGDA; IHG1; IRID1; RIEG3





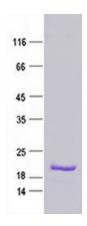
**Summary:** 

This gene belongs to the forkhead family of transcription factors which is characterized by a distinct DNA-binding forkhead domain. The specific function of this gene has not yet been determined; however, it has been shown to play a role in the regulation of embryonic and ocular development. Mutations in this gene cause various glaucoma phenotypes including primary congenital glaucoma, autosomal dominant iridogoniodysgenesis anomaly, and Axenfeld-Rieger anomaly. [provided by RefSeq, Jul 2008]

**Protein Families:** 

Druggable Genome, Transcription Factors

## **Product images:**



Purified recombinant protein FOXC1 was analyzed by SDS-PAGE gel and Coomossie Blue Staining.