



# SZABO SCANDIC

Part of Europa Biosite

## Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

Weitere Information auf den folgenden Seiten!  
See the following pages for more information!



### Lieferung & Zahlungsart

siehe unsere [Liefer- und Versandbedingungen](#)

### Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

### SZABO-SCANDIC HandelsgmbH

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## Datasheet

### FOXC1 (Human) Recombinant Protein (Q01)

**Catalog Number:** H00002296-Q01

**Regulation Status:** For research use only (RUO)

**Product Description:** Human FOXC1 partial ORF ( NP\_001444.1, 74 a.a. - 172 a.a.) recombinant protein with GST-tag at N-terminal.

**Sequence:**

KDMVKPPYSYIALITMAIQNAPDKKITLNGIYQFIMDRFP  
FYRDNKQGWQNSIRHNLSLNECFVKVPRDDKKPGKG  
SYWTLDPDSYNTFENGSLRRRR

**Host:** Wheat Germ (in vitro)

**Theoretical MW (kDa):** 36.63

**Applications:** AP, Array, ELISA, WB-Re  
(See our web site product page for detailed applications information)

**Protocols:** See our web site at  
<http://www.abnova.com/support/protocols.asp> or product page for detailed protocols

**Preparation Method:** [in vitro wheat germ expression system](#)

**Purification:** Glutathione Sepharose 4 Fast Flow

**Storage Buffer:** 50 mM Tris-HCl, 10 mM reduced Glutathione, pH=8.0 in the elution buffer.

**Storage Instruction:** Store at -80°C. Aliquot to avoid repeated freezing and thawing.

**Entrez GeneID:** 2296

**Gene Symbol:** FOXC1

**Gene Alias:** ARA, FKHL7, FREAC-3, FREAC3, IGDA, IHG1, IRID1

**Gene 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]