



# 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

### OLIG2 (Human) Recombinant Protein (P01)

bHLHe19

**Catalog Number:** H00010215-P01**Regulation Status:** For research use only (RUO)**Product Description:** Human OLIG2 full-length ORF (AAH36245.1, 1 a.a. - 323 a.a.) recombinant protein with GST-tag at N-terminal.**Sequence:**

```
MDSASLVSSRPSSPEPDDLFLPARSKGSSGSAFTGG
TVSSSTPSCDPELSAELRGAMGSAGHPVDKLGGS
GFKSSSSSTSSSTSSAAASSTKKDKKQMTPELQQLR
LKINSRERKRMHDLNIAMDGLREVMPYAHGPSVRKLS
KIATLLARNYILMLTNSLEEMKRLVSEIYGGHHAGFHP
SACGGLAHSAPLPAATAHPAAAAHAAHHPAVHHPILP
PAAAAAAAAAAAAVSSASLPGSGLPSVGSIRPPHGLL
KSPSAAAAAPLGGGGGGSGASGGFQHWGGMPCPCS
MCQVPPPHHVSAMGAGSLPRLTSDAK
```

**Host:** Wheat Germ (in vitro)**Theoretical MW (kDa):** 58.8**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:** 10215**Gene Symbol:** OLIG2**Gene Alias:** BHLHB1, OLIGO2, PRKCBP2, RACK17,**Gene Summary:** This gene encodes a basic helix-loop-helix transcription factor which is expressed in oligodendroglial tumors of the brain. The protein is an essential regulator of ventral neuroectodermal progenitor cell fate. The gene is involved in a chromosomal translocation t(14;21)(q11.2;q22) associated with T-cell acute lymphoblastic leukemia. Its chromosomal location is within a region of chromosome 21 which has been suggested to play a role in learning deficits associated with Down syndrome. [provided by RefSeq]