



# SZABO SCANDIC

Part of Europa Biosite

## Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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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

### MYO5A (Human) Recombinant Protein (Q01)

**Catalog Number:** H00004644-Q01

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

**Product Description:** Human MYO5A partial ORF ( NP\_000250.1, 1758 a.a. - 1853 a.a.) recombinant protein with GST-tag at N-terminal.

**Sequence:**

KKTDDDAEAICSMCNALTTAQIVKVLNLYTPVNEFEER  
VSVSFIRTIQMRLRDRKDSPQLLMDAKHIFPVTFPFNP  
SSLALETIQIPASLGLGFIS

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

**Theoretical MW (kDa):** 36.3

**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:** 4644

**Gene Symbol:** MYO5A

**Gene Alias:** GS1, MYH12, MYO5, MYR12

**Gene Summary:** This gene is one of three myosin V heavy-chain genes, belonging to the myosin gene superfamily. Myosin V is a class of actin-based motor proteins involved in cytoplasmic vesicle transport and anchorage, spindle-pole alignment and mRNA

translocation. The protein encoded by this gene is abundant in melanocytes and nerve cells. Mutations in this gene cause Griscelli syndrome type-1 (GS1), Griscelli syndrome type-3 (GS3) and neuroectodermal melanolyosomal disease, or Elejalde disease. Multiple alternatively spliced transcript variants encoding different isoforms have been reported, but the full-length nature of some variants has not been determined. [provided by RefSeq]