



# 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

### AICDA (Human) Recombinant Protein

**Catalog Number:** P9590

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

**Product Description:** Human AICDA (Q9GZX7, 1 a.a. - 198 a.a.) full recombinant protein with His tag at N-terminus expressed in *Escherichia coli*.

**Sequence:**

MGSSHHHHHSSGLVPRGSHMDSLLMNRRKFLYQFK  
NVRWAKGRRETYLCYVVKRRDSATSFSLDFGYLRNK  
NGCHVELLFLRYISDWLDPGRCYRVTWFTSWSPCY  
DCARHVADFLRGNPNLSLRIFTARLYFCEDRKAPEGL  
RRLHRAGVQIAIMTFKDYFCWNTFVENHERTFK  
AWEGLHENSVRLSRQLRRILLPLYEVDDLRFDAFRTLGL

**Host:** *Escherichia coli*

**Theoretical MW (kDa):** 26.1

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

**Form:** Liquid

**Preparation Method:** *Escherichia coli* expression system

**Purity:** > 85.0% by SDS-PAGE

**Recommend Usage:** Biological Activity

SDS-PAGE

The optimal working dilution should be determined by the end user.

**Storage Buffer:** In 20mM Tris-HCl pH 8.0 (0.4 M Urea and 10% glycerol)

**Storage Instruction:** Store at 2°C to 8°C for 1 week.

For long term storage, aliquot and store at -20°C to -80°C.

Aliquot to avoid repeated freezing and thawing.

**Entrez GeneID:** 57379

**Gene Symbol:** AICDA

**Gene Alias:** AID, ARP2, CDA2, HIGM2

**Gene Summary:** This gene encodes a RNA-editing deaminase that is a member of the cytidine deaminase family. The protein is involved in somatic hypermutation, gene conversion, and class-switch recombination of immunoglobulin genes. Defects in this gene are the cause of autosomal recessive hyper-IgM immunodeficiency syndrome type 2 (HIGM2). [provided by RefSeq]