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

Quellenstraße 110, A-1100 Wien

T. +43(0)1 489 3961-0

F. +43(0)1 489 3961-7

mail@szabo-scandic.com

www.szabo-scandic.com

linkedin.com/company/szaboscandic

9F, No. 108, Jhouzih St.,Taipei, Taiwan
Tel: + 886-2-8751-1888 Fax: + 886-2-6602-1218
E-mail: sales@abnova.com

Datasheet

PRODH (Human) Recombinant Protein (Q01)

Catalog Number: H00005625-Q01

Regulation Status: For research use only (RUO)

Product Description: Human PRODH partial ORF (NP_057419, 441 a.a. - 540 a.a.) recombinant protein with GST-tag at N-terminal.

Sequence:
LVRGAYLAQERARAAEIGYEDPINPTYEATNAMYHRCL
DYVLEELKHNAKAKVMVASHNEDTVRFALRRMEELGL
HPADHRVYFGQLLGMCDQISFPLGQ

Host: Wheat Germ (in vitro)

Theoretical MW (kDa): 36.74

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: 5625

Gene Symbol: PRODH

Gene Alias: FLJ33744, HSPOX2, MGC148078, MGC148079, PIG6, POX, PRODH1, PRODH2, SCZD4, TP53I6

Gene Summary: The protein encoded by this gene is a mitochondrial proline dehydrogenase that catalyzes the first step in proline degradation. Defects in this gene are a cause of hyperprolinemia type 1 and possibly susceptibility to schizophrenia 4 (SCZD4). The gene is located on chromosome 22q11.21, a region which has also been associated with the contiguous gene deletion syndromes DiGeorge syndrome and CATCH22 syndrome. [provided by RefSeq]