

# 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 <u>mail@szabo-scandic.com</u> www.szabo-scandic.com

#### SANTA CRUZ BIOTECHNOLOGY, INC.

## PKLR (m): 293 Lysate: sc-179336



#### BACKGROUND

In mammals, four different isoenzymes exist for pyruvate kinase. Based on their tissue distribution, the isoenzymes are designated L-type (for predominant expression in the liver), R-type (for predominant expression in red blood cells), M1-type (for predominant expression in muscle, brain and heart) and M2-type (for predominant expression in fetal tissues). Pyruvate kinases are responsible for catalyzing the final step in glycolysis: the conversion of phosphoenolpyruvate to pyruvate with the coinciding generation of ATP. The PKLR (pyruvate kinase, liver and RBC) gene encodes the L- and R-type isoenzymes through alternative splicing events under the control of different promoters. The R-type isoform, also known as RPK (R-type pyruvate kinase) exists as a tetramer and when functioning improperly, can result in chronic/hereditary nonspherocytic hemolytic anemia (CNSHA/HNSHA) or pyruvate kinase hyperactivity (also called high red cell ATP syndrome). The L-type isoform, alternatively known as PKL (pyruvate kinase L-type), also exists as a tetramer and is upregulated by glucose with implications in maturity-onset diabetes of the young (MODY).

#### REFERENCES

- Tani, K., Fujii, H., Tsutsumi, H., Sukegawa, J., Toyoshima, K., Yoshida, M.C., Noguchi, T., Tanaka, T. and Miwa, S. 1987. Human liver type pyruvate kinase: cDNA cloning and chromosomal assignment. Biochem. Biophys. Res. Commun. 143: 431-438.
- Tani, K., Tsutsumi, H., Takahashi, K., Ogura, H., Kanno, H., Hayasaka, K., Narisawa, K., Nakahata, T., Akabane, T. and Morisaki, T. 1988. Two homozygous cases of erythrocyte pyruvate kinase (PK) deficiency in Japan: PK Sendai and PK Shinshu. Am. J. Hematol. 28: 186-190.
- Nordström, L. and Lerner, S.A. 1991. Single daily dose therapy with aminoglycosides. J. Hosp. Infect. Suppl. A: 117-129.
- Wang, H., Chu, W., Das, S.K., Ren, Q., Hasstedt, S.J. and Elbein, S.C. 2002. Liver pyruvate kinase polymorphisms are associated with type 2 diabetes in northern European Caucasians. Diabetes 51: 2861-2865.
- van Wijk, R., van Solinge, W.W., Nerlov, C., Beutler, E., Gelbart, T., Rijksen, G. and Nielsen, F.C. 2003. Disruption of a novel regulatory element in the erythroid-specific promoter of the human PKLR gene causes severe pyruvate kinase deficiency. Blood 101: 1596-1602.
- Park-Hah, J.O., Kanno, H., Kim, W.D. and Fujii, H. 2005. A novel homozygous mutation of PKLR gene in a pyruvate-kinase-deficient Korean family. Acta Haematol. 113: 208-211.

#### CHROMOSOMAL LOCATION

Genetic locus: Pklr (mouse) mapping to 3 F1.

#### PRODUCT

PKLR (m): 293 Lysate represents a lysate of mouse PKLR transfected 293 cells and is provided as 100  $\mu$ g protein in 200  $\mu$ l SDS-PAGE buffer.

#### **STORAGE**

Store at -20° C. Repeated freezing and thawing should be minimized. Sample vial should be boiled once prior to use. Non-hazardous. No MSDS required.

#### APPLICATIONS

PKLR (m): 293 Lysate is suitable as a Western Blotting positive control for mouse reactive PKLR antibodies. Recommended use: 10-20  $\mu l$  per lane.

Control 293 Lysate: sc-110760 is available as a Western Blotting negative control lysate derived from non-transfected 293 cells.

#### **RESEARCH USE**

For research use only, not for use in diagnostic procedures

#### PROTOCOLS

See our web site at www.scbt.com for detailed protocols and support products.