

GLUT4 (SLC2A4) tetramer transports Glc from extracellular region to cytosol

D'Eustachio, P., He, L.

European Bioinformatics Institute, New York University Langone Medical Center, Ontario Institute for Cancer Research, Oregon Health and Science University.

The contents of this document may be freely copied and distributed in any media, provided the authors, plus the institutions, are credited, as stated under the terms of [Creative Commons Attribution 4.0 International \(CC BY 4.0\) License](https://creativecommons.org/licenses/by/4.0/). For more information see our [license](https://reactome.org/licenses/).

17/05/2024

Introduction

Reactome is open-source, open access, manually curated and peer-reviewed pathway database. Pathway annotations are authored by expert biologists, in collaboration with Reactome editorial staff and cross-referenced to many bioinformatics databases. A system of evidence tracking ensures that all assertions are backed up by the primary literature. Reactome is used by clinicians, geneticists, genomics researchers, and molecular biologists to interpret the results of high-throughput experimental studies, by bioinformaticians seeking to develop novel algorithms for mining knowledge from genomic studies, and by systems biologists building predictive models of normal and disease variant pathways.

The development of Reactome is supported by grants from the US National Institutes of Health (P41 HG003751), University of Toronto (CFREF Medicine by Design), European Union (EU STRP, EMI-CD), and the European Molecular Biology Laboratory (EBI Industry program).

Literature references

Fabregat, A., Sidiropoulos, K., Viteri, G., Forner, O., Marin-Garcia, P., Arnau, V. et al. (2017). Reactome pathway analysis: a high-performance in-memory approach. *BMC bioinformatics*, 18, 142. [↗](#)

Sidiropoulos, K., Viteri, G., Sevilla, C., Jupe, S., Webber, M., Orlic-Milacic, M. et al. (2017). Reactome enhanced pathway visualization. *Bioinformatics*, 33, 3461-3467. [↗](#)

Fabregat, A., Jupe, S., Matthews, L., Sidiropoulos, K., Gillespie, M., Garapati, P. et al. (2018). The Reactome Pathway Knowledgebase. *Nucleic Acids Res*, 46, D649-D655. [↗](#)

Fabregat, A., Korninger, F., Viteri, G., Sidiropoulos, K., Marin-Garcia, P., Ping, P. et al. (2018). Reactome graph database: Efficient access to complex pathway data. *PLoS computational biology*, 14, e1005968. [↗](#)

Reactome database release: 88

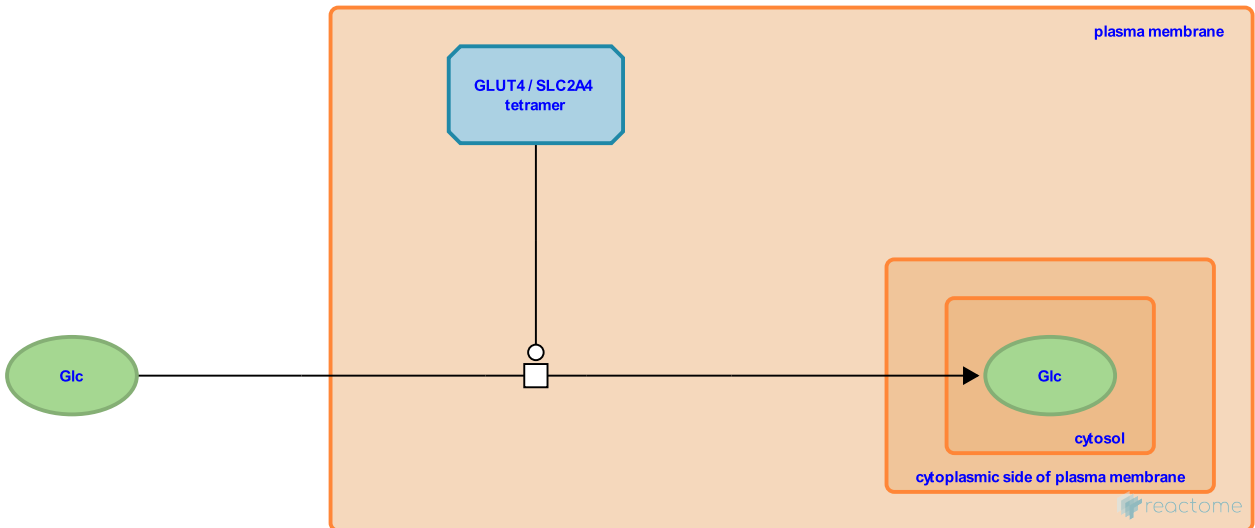
This document contains 1 reaction ([see Table of Contents](#))

GLUT4 (SLC2A4) tetramer transports Glc from extracellular region to cytosol ↗

Stable identifier: R-HSA-8981570

Type: transition

Compartments: cytosol, plasma membrane, extracellular region



Tetrameric GLUT4, the SLC2A4 gene product, associated with the plasma membrane, mediates the facilitated diffusion of glucose (Glc) into cells. GLUT4 is found in heart, skeletal muscle, brain and adipose tissue. GLUT4 molecules are translocated from an intracellular store to the cell surface in response to increased insulin levels, increasing glucose transport 10-20-fold (Bryant et al. 2002; Fukumoto et al. 1989). Defects in SLC2A4 may be a cause of non-insulin-dependent diabetes mellitus (NIDDM) (Kusari et al. 1991; Choi et al. 1991).

Literature references

Flier, JS., Buse, JB., Choi, WH., Morgan, R., Moller, DE., O'Rahilly, S. et al. (1991). Molecular scanning of insulin-responsive glucose transporter (GLUT4) gene in NIDDM subjects. *Diabetes*, 40, 1712-8. ↗

James, DE., Bryant, NJ., Govers, R. (2002). Regulated transport of the glucose transporter GLUT4. *Nat Rev Mol Cell Biol*, 3, 267-77. ↗

Buse, JB., Verma, US., Olefsky, JM., Henry, RR., Kusari, J. (1991). Analysis of the gene sequences of the insulin receptor and the insulin-sensitive glucose transporter (GLUT-4) in patients with common-type non-insulin-dependent diabetes mellitus. *J Clin Invest*, 88, 1323-30. ↗

Bell, GI., Pilch, PF., Seino, S., Kayano, T., Buse, JB., Fukumoto, H. (1989). Cloning and characterization of the major insulin-responsive glucose transporter expressed in human skeletal muscle and other insulin-responsive tissues. *J Biol Chem*, 264, 7776-9. ↗

Editions

2004-06-23	Authored, Edited	D'Eustachio, P.
2009-08-24	Reviewed	He, L.
2009-12-12	Revised	D'Eustachio, P.