

Formation of Ragulator complex

Jupe, S., Zwartkruis, FJ.

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](#). For more information see our [license](#).

29/04/2024

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

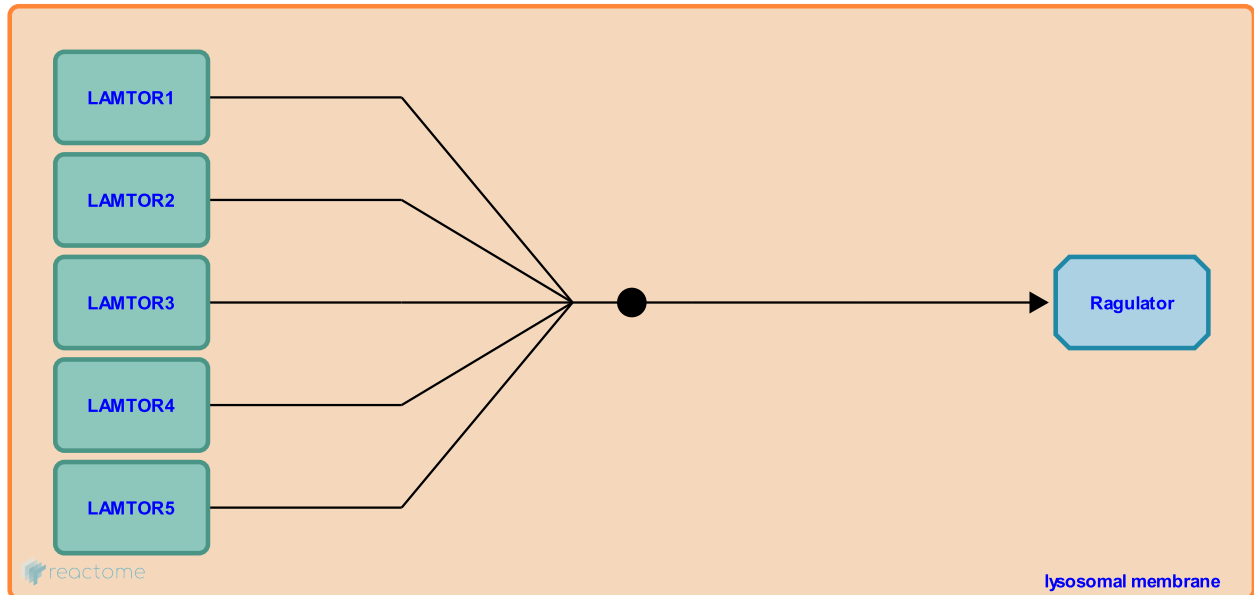
This document contains 1 reaction ([see Table of Contents](#))

Formation of Ragulator complex ↗

Stable identifier: R-HSA-5653936

Type: binding

Compartments: lysosomal membrane



A complex of LAMTOR1 to 5, known as Ragulator, interacts with Rag GTPases recruiting them to lysosomes, an essential step in mTORC1 activation (Sancak et al. 2010). LAMTOR1 is probably myristoylated and/or palmitoylated to enhance its association with the lysosomal surface, acting as a platform for the other members of the complex. Lamtor1 (Nada et al. 2009) and Lamtor2 (Teis et al. 2006) knockout mice exhibit severe growth retardation, severe defects in intracellular organelle organization and embryonic lethality. Partial reduction of LAMTOR2 in humans leads to reduced height (Bohn et al. 2007).

Literature references

Sabatini, DM., Zoncu, R., Sancak, Y., Bar-Peled, L., Nada, S., Markhard, AL. (2010). Ragulator-Rag complex targets mTORC1 to the lysosomal surface and is necessary for its activation by amino acids. *Cell*, 141, 290-303. ↗

Sabatini, DM., Zoncu, R., Bar-Peled, L., Schweitzer, LD. (2012). Ragulator is a GEF for the rag GTPases that signal amino acid levels to mTORC1. *Cell*, 150, 1196-208. ↗

Editions

2015-02-02	Authored	Jupe, S.
2015-04-08	Revised	Jupe, S.
2015-05-13	Edited	Jupe, S.
2015-05-14	Reviewed	Zwartkruis, FJ.