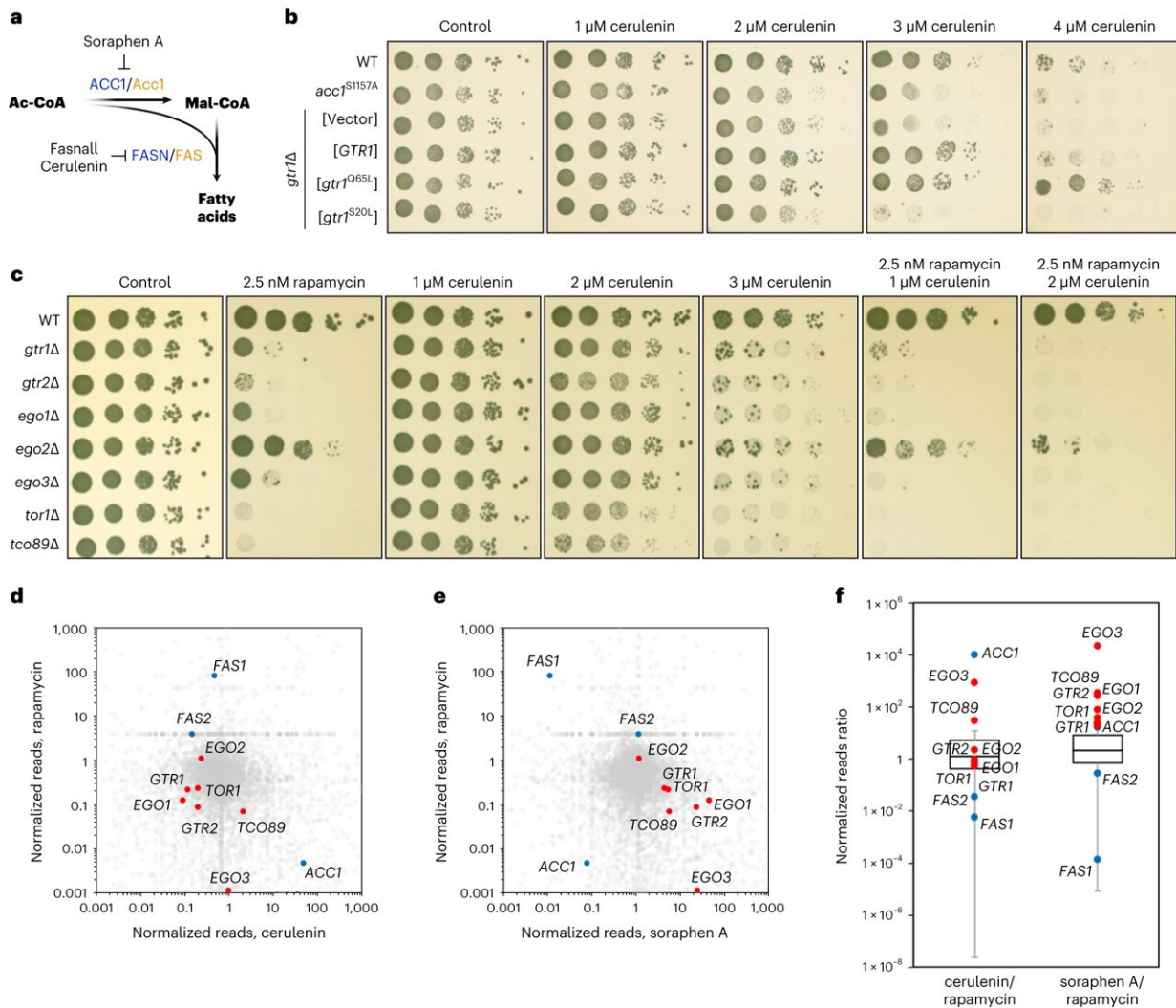


Metabolism meets signaling to fine-tune cell growth

August 10 2023, by Maren Berghoff



Functional pharmacogenetic interactions between TOR-pathway genes, and Acc1 and Fas1 activity in yeast. **a**, Schematic diagram of de novo FA biosynthesis. Yeast and mammalian proteins are shown in orange and blue,

respectively. **b**, Yeast Rag GTPase mutants that impair or promote TORC1 activity are cerulenin-sensitive or cerulenin-resistant, respectively. Tenfold serial dilutions of wild-type (WT), hyperactive $\text{Acc1}^{\text{S1157A}}$ -expressing ($\text{acc1}^{\text{S1157A}}$) and $\text{gtr1}\Delta$ cells expressing plasmid-encoded WT GTR1 , $\text{gtr1}^{\text{Q65L}}$ (TORC1-activating) or $\text{gtr1}^{\text{S20L}}$ (TORC1-inactivating), or containing an empty vector were spotted on plates with the indicated concentrations of cerulenin or vehicle (control) and cultured at 30 °C for 2 d. **c**, TORC1 and EGO C mutants are sensitive to cerulenin and hypersensitive to combined cerulenin and rapamycin treatment. Drop spot assays were performed as in **b** using the indicated cell strains and plates containing the indicated concentrations of rapamycin and/or cerulenin. **d**, Positive correlation between rapamycin- and cerulenin-induced SATAY transposition profiles in TORC1 and EGO C genes. **e**, Negative correlation between rapamycin- and soraphen A-induced SATAY transposition profiles in TORC1 and EGO C genes. **d,e**, The dot plots show the ratio of transposition events (reads) per coding gene in rapamycin- and cerulenin-treated (**d**), and rapamycin- and soraphen A-treated (**e**) versus previously published untreated SATAY libraries^{24,25}. **f**, Summary of the pairwise correlations of the transposon profiles of TORC1, EGO C (red dots) and FA biosynthesis (blue dots) genes shown in **d,e**. Boxplots: central line, median; box, interquartile range (IQR; 25th (Q1)–75th (Q3) percentile); and whiskers, $\text{Q3} + 1.5 \times \text{IQR}$ and $\text{Q1} - 1.5 \times \text{IQR}$. Source numerical data are provided. Credit: *Nature Cell Biology* (2023). DOI: 10.1038/s41556-023-01198-6

Cells use a variety of metabolic pathways to synthesize the building blocks for growth and proliferation. To ensure balanced growth, these biosynthetic processes must be tightly coordinated. Researchers from the Max Planck Institute for Biology of Ageing, together with a team of national and international collaborators, have now identified a molecular machinery that senses the ability of a cell to make lipids to then block or activate all other biosynthetic processes, such as protein synthesis, accordingly.

Cells accumulate mass and grow by using nutrients and energy to build

membranes, proteins, [nucleic acids](#), and other macromolecular structures, through the coordinated action of various metabolic pathways. Therefore, nutrient sensing mechanisms ensure that cells only grow when all conditions are optimal.

The most important nutrient sensor in cells is a [protein complex](#) called mTORC1. When [amino acids](#), the building blocks of proteins, are abundant, mTORC1 is active and promotes [protein synthesis](#), thus linking amino acid availability to the respective cellular function. However, how the ability of cells to make lipids is sensed and whether mTORC1 plays a role in this process has been very poorly understood.

In fatty acid biosynthesis, acetyl-CoA, a metabolite originating mainly from the breakdown of glucose (another important nutrient), is first converted to malonyl-CoA by an enzyme called ACC1, which is then converted to [fatty acids](#) by an enzyme called FASN (fatty acid synthase).

The researchers showed that accumulation of malonyl-CoA, which can occur when FASN levels or activity are low, leads to inhibition of mTORC1 activity and downregulation of other biosynthetic functions that are regulated by mTORC1, such as protein synthesis. Surprisingly, they discovered that malonyl-CoA binds directly to the catalytic pocket of mTOR, competing away ATP, another metabolite that is necessary for mTOR to be active and phosphorylate its substrates.

"These findings highlight malonyl-CoA as the first described endogenous metabolite that acts as an ATP-competitive inhibitor for a signaling kinase in [mammalian cells](#)," says Constantinos Demetriades, research group leader at the Max Planck Institute for Biology of Ageing and head of the study.

"It is noteworthy that this phenomenon is evolutionarily conserved from yeast to [human cells](#), thus it represents an ancient mechanism via which

lipid metabolism talks to the core regulator of all other metabolic processes, mTORC1."

Linking fatty acid biosynthesis to cell growth through mTORC1

At the [molecular level](#), the researchers found that mTORC1 forms physical interactions with ACC1 and FASN, allowing it to directly sense the levels of malonyl-CoA at the site where the metabolite is produced. In this way, the direct binding of malonyl-CoA to mTORC1 switches off all other cellular processes that are controlled downstream of this complex.

As a result, under conditions that a cell cannot produce enough fatty acids, mTORC1 inhibition would also block synthesis of proteins, other anabolic processes and, eventually, cell growth, until conditions are optimal again.

As pharmacological inhibitors of FASN, like those used in this study, are used as anti-cancer drugs in the clinic, these discoveries may also be relevant for cancer treatment in the future. Based on the data described here, FASN inhibitors likely have a two-fold effect in cancer cells: not only do they block fatty acid synthesis, which is required for membrane formation and energy production, but they also inhibit mTORC1 activity, which is dysregulated in most cancers.

The paper is published in the journal *Nature Cell Biology*.

More information: Raffaele Nicastro et al, Malonyl-CoA is a conserved endogenous ATP-competitive mTORC1 inhibitor, *Nature Cell Biology* (2023). [DOI: 10.1038/s41556-023-01198-6](https://doi.org/10.1038/s41556-023-01198-6)

Provided by Max Planck Society

Citation: Metabolism meets signaling to fine-tune cell growth (2023, August 10) retrieved 29 April 2024 from <https://medicalxpress.com/news/2023-08-metabolism-fine-tune-cell-growth.html>

This document is subject to copyright. Apart from any fair dealing for the purpose of private study or research, no part may be reproduced without the written permission. The content is provided for information purposes only.