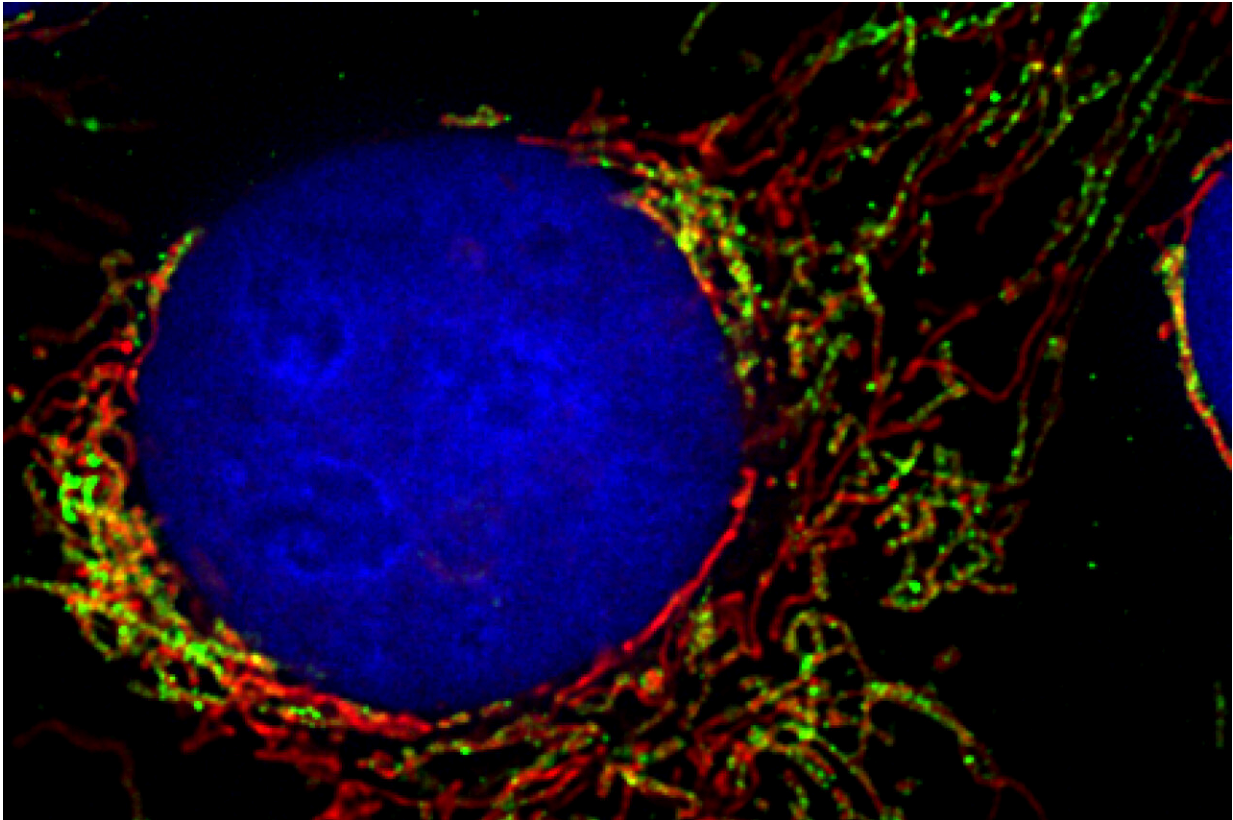


# How cells control mitochondria

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The network of mitochondria runs like a thread through the entire cell (marked in green and red with fluorescent proteins). The cell nucleus is stained in blue.  
Credit: Dr. Pablo Sánchez-Martín/University of Freiburg

Errors in the metabolic processes of mitochondria are responsible for a variety of diseases such as Parkinson's and Alzheimer's. Scientists needed to find out just how the necessary building blocks are imported

into the complex biochemical apparatus of these cell areas. The TOM complex (translocase of the outer mitochondrial membrane) is considered the gateway to the mitochondrion, the proverbial powerhouse of the cell. The working group headed by Professor Chris Meisinger at the Institute of Biochemistry and Molecular Biology at the University of Freiburg has now demonstrated—in human cells—how signaling molecules control this gate. A signaling protein called DYRK1A modifies the molecular machinery of TOM and makes it more permeable for enzymes that are important for the cell metabolism. The group has thus discovered the first signaling protein that directly influences this import process in humans. Their work has been published in the journal *Nature Communications*.

## **Developmental disorders in a new light**

In [neurodevelopmental disorders](#) such as autism, microcephaly and Down's syndrome, DYRK1A is defective. "The connection with mitochondria is new. These results allow us to better understand these disorders and develop treatment strategies," says Dr. Adinarayana Marada, a member of Meisinger's team.

"For a long time, researchers thought that the TOM complex was a rigid structure in the mitochondrial membrane whose doors were always open," Meisinger explains. His team recently demonstrated signaling mechanisms in baker's yeast that alter the subunits of the TOM complex depending on the metabolic state of the cell, or in response to sudden stress. In this way, the cell can specifically control the influx of precursor proteins for building elements of the metabolism, and it can adapt the function of the mitochondria to an altered cellular state. Whether such mechanisms also exist in humans was previously unknown.

## **DYRK1A acts upon the TOM complex**

The first authors of the study, Dr. Corvin Walter and Dr. Adinarayana Marada of Meisinger's research group, developed a systematic approach to track down signaling mechanisms such as those triggered by protein kinases, in humans. Over several years, they tested candidates using cell biological and bioinformatic methods and found what they were looking for—DYRK1A, one such protein kinase, acts on the TOM complex. "With this, we actually found the needle in the haystack," says Walter.

**More information:** Corvin Walter et al, Global kinome profiling reveals DYRK1A as critical activator of the human mitochondrial import machinery, *Nature Communications* (2021). [DOI: 10.1038/s41467-021-24426-9](https://doi.org/10.1038/s41467-021-24426-9)

Provided by Albert Ludwigs University of Freiburg

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