

## Molecular biologists discover an active role of membrane lipids in health and disease

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Both unfolded proteins and aberrant membrane lipid compositions are sensed by Ire1 and can activate the so-called unfolded protein response. This pathway is crucial for the function and survival of secretory cells such as antibody-producing plasma cells. Credit: Robert Ernst



All living cells that grow and divide have a constant demand for producing new proteins and new membrane lipids. Some cells of the human body, however, are specialized to secrete tremendous amounts of proteins. Plasma cells, for example, produce antibodies that ward off bacteria and viruses. Another example are cells from the pancreas that manufacture insulin, which is essential for regulating blood sugar levels. Such cells are known as secretory cells.

The secretory capacity of a cell is controlled by a process known as the unfolded protein response (UPR). A central task of UPR is to detect misfolded proteins, i.e. proteins that do not have the correct shape to do their job. These misfolded proteins are dangerous as they can aggregate with other proteins, form clumps and clog cellular functions. The UPR can stop this vicious circle arising from the accumulation of misfolded or unfolded proteins by helping to degrade the clumps and by promoting protein refolding. The UPR therefore plays a key role in secretory cells but also protects other cells of the body from the stresses associated with an accumulation of unfolded or misfolded proteins.

But there's a catch: Viruses and tumour cells can also exploit the UPR for their own ends, enabling them to grow at a faster rate and to thwart the body's immune response. Moreover, sustained activation of the UPR can drive cells into cell death.

Cells in which the UPR has been activated can produce larger quantities of proteins, but they also become more sensitive – a bit like a finely tuned race car. As Robert Ernst explains: "Whereas a racing car will often fail after completing a hundred super-fast laps because the engine has overheated, a tractor (representing a normal body cell) will continue to drive up and down the field for a lot longer, but also a lot slower." Why these high-performance cells are so much more sensitive was not known up until now.



The team led by Professor Ernst has now solved this puzzle and has described how the UPR senses <u>membrane</u> lipids and responds accordingly. Working in close collaboration with scientists from Goethe University and the Max Planck Institute of Biophysics, both located in Frankfurt, the research team from the Department of Medical Biochemistry and Molecular Biology at Saarland University has identified a novel mechanism that leads to UPR activation and that can trigger long-term stress in cells.

According to this new mechanism, the UPR is activated not only by <u>misfolded proteins</u>, but also by anomalous membrane <u>lipid</u> compositions. Secretory <u>cells</u> are particularly sensitive to these changes, because they have already activated their UPR to produce more proteins and therefore at risk of 'overheating'– just like the racing car engine described above. The study provides a new perspective on the active role of biological membranes may be a game changer for the understanding of a great variety diseases.

Over the last few years, research work in Japan, Great Britain and the U.S. has provided growing evidence of the unexpected role played by membrane lipids. In 2011, Peter Walter and David Ron, pioneer in the field of human UPR, came to regard the relationship between membrane lipids and the UPR as a central unresolved question in the field. Now, it has been solved.

"In order to boost the production and secretion of proteins, the UPR regulates more than 5 percent of all human genes," explains Robert Ernst. "We now have the conceptual framework to understand why secretory cells are hypersensitive to changes of their <u>membrane lipids</u> induced by the diet."

News of the results has spread fast, and the journal *Nature Reviews Molecular Cell Biology* has launched a commentary to the article that has



been published in Molecular Cell.

The study "Activation of the Unfolded Protein Response by Lipid Bilayer Stress" will be published on 17 August 2017 in the journal *Molecular Cell*. The article has been available online since 6 July.

The commentary to this article, "Unfolded protein response: Reacting to membrane stress," by Paulina Strzyz, was published online by *Nature Reviews Molecular Cell Biology* on 21 July 2017.

**More information:** Kristina Halbleib et al. Activation of the Unfolded Protein Response by Lipid Bilayer Stress, *Molecular Cell* (2017). <u>DOI:</u> <u>10.1016/j.molcel.2017.06.012</u>

Paulina Strzyz. Unfolded protein response: Reacting to membrane stress, *Nature Reviews Molecular Cell Biology* (2017). DOI: 10.1038/nrm.2017.74

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