

Molecular culprits of protein aggregation in ALS and FTLD

19 July 2018

The mutated and aggregated protein FUS is implicated in two neurodegenerative diseases: amyotrophic lateral sclerosis (ALS) and frontotemporal lobar degeneration (FTLD). Using a newly developed fruit fly model, researchers led by prof. Ludo Van Den Bosch (VIB-KU Leuven) have focused on the protein structure of FUS to gain more insight into how it causes neuronal toxicity and disease.

ALS and FTLD are adult-onset neurodegenerative disorders that differ in terms of the symptoms they cause and the neurons they affect. In ALS, the neurons that control muscle movement degenerate, resulting in progressive paralysis, while FTLD affects specific brain areas and causes dementia. Still, many patients present with symptoms of both diseases, so scientists and clinicians now believe they are actually on opposite ends of the same disease spectrum.

The overlap between ALS and FTLD is not only apparent in the clinic, but also when looking at the underlying disease mechanisms. FUS, for example, is implicated in both diseases. Mutations in the FUS gene cause familial ALS and aggregation of the FUS [protein](#) is observed in both ALS and FTLD.

From liquid droplets to insoluble aggregates

FUS normally resides in the nucleus of the cell, but relocates to stress granules in the cytoplasm upon cellular stress. Stress granules are basically [liquid droplets](#) in the cell, similar in content to the toxic protein aggregates found in ALS and FTLD, but different in that their assembly is dynamic and reversible.

Could these liquid [droplets](#) serve as stepping stones toward the formation of aggregates that are typical for disease? "We believe so," says Elke Bogaert, one of the researchers working with prof. Van Den Bosch. "Both FUS droplets and hydrogels

have been shown to undergo a switch to irreversible fibrillarization in a test tube, but this process has not been studied in a cellular context".

The team generated a fruit fly model of FUS toxicity to investigate the formation of liquid droplets in more detail. They identified a previously unrecognized synergistic effect between two different protein regions that mediate toxicity. "We found that the FUS protein that is misbehaving in ALS and FTLD can form liquid droplets via specific molecular interactions between hydrophobic and charged amino acids," explains Steven Boeynaems, another researcher involved in the study. One FUS protein domain was generally considered to mediate aggregation, but the new findings indicate that arginine residues in another region of the protein are also required for maturation of FUS in cellular stress granules.

Problems in flies, and humans?

Further experiments immediately hinted at the importance of this protein interaction. Boeynaems says, "We showed that the interactions between exactly these two protein domains could explain toxicity in our fruit fly ALS model, suggesting how the protein may be misregulated in patients as well."

Prof. Ludo Van Den Bosch underscores the importance of figuring out the exact process of this so-called phase separation of droplets, that mature into aggregates: "Deciphering how and why proteins such as FUS start aggregating in the brain will be key to understand neurodegenerative diseases and could lead to novel therapeutic strategies. Our new findings highlight an important role for arginine-rich domains in the pathology of these proteins."

More information: Molecular dissection of FUS points at synergistic effect of low-complexity domains in toxicity, Bogaert, Boeynaems, et al.

2018 *Cell Reports*. [www.cell.com/cell-reports/full ...](http://www.cell.com/cell-reports/full...)
[2211-1247\(18\)30997-5](https://doi.org/10.1016/j.celrep.2018.07.019)

Provided by VIB (the Flanders Institute for
Biotechnology)

APA citation: Molecular culprits of protein aggregation in ALS and FTLD (2018, July 19) retrieved 30
August 2022 from

<https://medicalxpress.com/news/2018-07-molecular-culprits-protein-aggregation-als.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.