Headline: Goodbye ‘stress granules’: Study expands possibilities for treating neurological diseases

(Tokyo, May 31) Cell biologists have deepened understanding of proteins associated with neurodegenerative diseases. The findings could open up new treatment approaches for disorders including amyotrophic lateral sclerosis (ALS), among others.

Researchers in Japan have gained valuable insights into ‘stress granules’ — clumps of RNAs and proteins that form when cells are stressed by factors such as heat, toxins and viruses.

As stress granules are linked to a range of neurodegenerative diseases, understanding how they form, and how they can be reduced, is of great interest to the medical world.

Published in the Journal of Cell Science, their study reveals the important role of two enzymes in disassembling stress granules.

These two enzymes, named USP5 and USP13, belong to a group of nearly 100 known deubiquitylases, which are thought to work by cutting ubiquitin chains inside stress granules.

The study is the culmination of over five years of work by the team, including Masayuki Komada, Toshiaki Fukushima and Shunsuke Matsumoto of Tokyo Institute of Technology (Tokyo Tech). First author Xuan Xie, a PhD student at the laboratory led by Komada, has described how they arrived at their ‘eureka’ moment in an interview with the journal’s First Person series.

As a first step, the researchers demonstrated that USP5 and USP13 are preferentially recruited to heat-induced stress granules. (See Figure 1.)

**Figure 1. Evidence of USP5, USP13 and ubiquitin chains inside stress granules.** Immunostaining and microscopic analysis revealed that cells under heat stress form stress granules containing USP5, USP13 and ubiquitin chains. This gathering together is known as co-localization.
“We found that heat-induced stress granules contain ubiquitin chains, much more so than in stress granules induced by other stressors,” explains Fukushima. “This implied that ubiquitin chains may recruit USP5 and USP13 to stress granules.”

Importantly, as ubiquitin chains are often found in stress granules in neurodegenerative diseases, the heat-induced stress granules provided a good model for further investigation.

Next, the team compared what happens to heat-shocked cells with and without the two enzymes. The cells were exposed to a temperature of 44°C for one hour, and returned to 37°C for one hour. During the recovery period, in cells lacking USP5 and USP13, the team found that the disassembly of stress granules was delayed. (See Figure 2.)

Specifically, in cells containing USP5 and USP13, the percentage of cells with stress granules fell to 14%, whereas this figure was 60% or more in cells without the two enzymes.

Figure 2. Comparison of cells with and without USP5 and USP13
Normal cells (top row) and cells without USP5 and USP13 (bottom row) were exposed to heat stress, and then recovered from heat stress. Cells were stained for USP5 and USP13 (left panels), as well as a stress granule marker (G3BP) and a nucleus marker (DAPI) (right panels). In cells lacking the two enzymes, the disassembly of stress granules after recovery from heat stress was delayed.

The findings suggest that the presence of USP5 and USP13 is critical to the disassembly of stress granules. (See Figure 3.)

Although the exact mechanisms have yet to be determined, the researchers propose that USP5 hydrolyzes or “cuts” unanchored ubiquitin chains, while USP13 cuts protein-conjugated ubiquitin chains.

“We concluded that both reactions are required for the efficient destabilization of stress granules,” says Fukushima.

The study may lead to the development of “artificial deubiquitinating enzymes”, which could have a profound impact on future medical treatments.
Developing such innovative enzymes that “possess high activity and show specific localization to stress granules” is a feat that could be achieved in five years, Fukushima adds.

Figure 3. Illustration of the inner structure of stress granules
USP5 and USP13 “cut” ubiquitin chains in stress granules via a reaction called hydrolysis. This reaction appears to be crucial to the efficient disassembly of stress granules after recovery from heat stress.

Technical terms
1 Ubiquitin chains: Chains of small regulatory proteins called ubiquitin that are found everywhere in eukaryotic cells.

2 Protein-conjugated: Refers to the way that some ubiquitin chains are bound to proteins, as opposed to being unanchored, within the stress granules. One of the next goals of the team is to identify these target proteins.

References
Authors: Xuan Xie1,2, Shunsuke Matsumoto2, Akinori Endo1, Toshiaki Fukushima1,2,*, Hiroyuki Kawahara3, Yasushi Saeki2 and Masayuki Komada1,2
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DOI: 10.1242/jcs.210856
Affiliation: 1. Cell Biology Center, Institute of Innovative Research, Tokyo Institute of Technology
2. School of Life Science and Technology, Tokyo Institute of
Technology
3. Department of Biological Sciences, Tokyo Metropolitan University
4. Laboratory of Protein Metabolism, Tokyo Metropolitan Institute of Medical Science

Corresponding author’s email: tofu@bio.titech.ac.jp

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Contact
Emiko Kawaguchi
Public Relations Section,
Tokyo Institute of Technology
media@jim.titech.ac.jp
+81-3-5734-2975

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