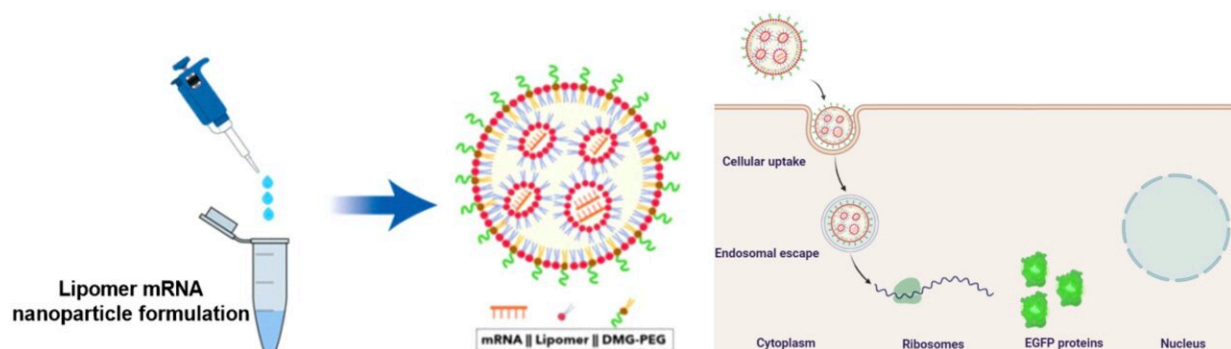


# Lipid polymer enables safe delivery of RNA drugs to the lungs

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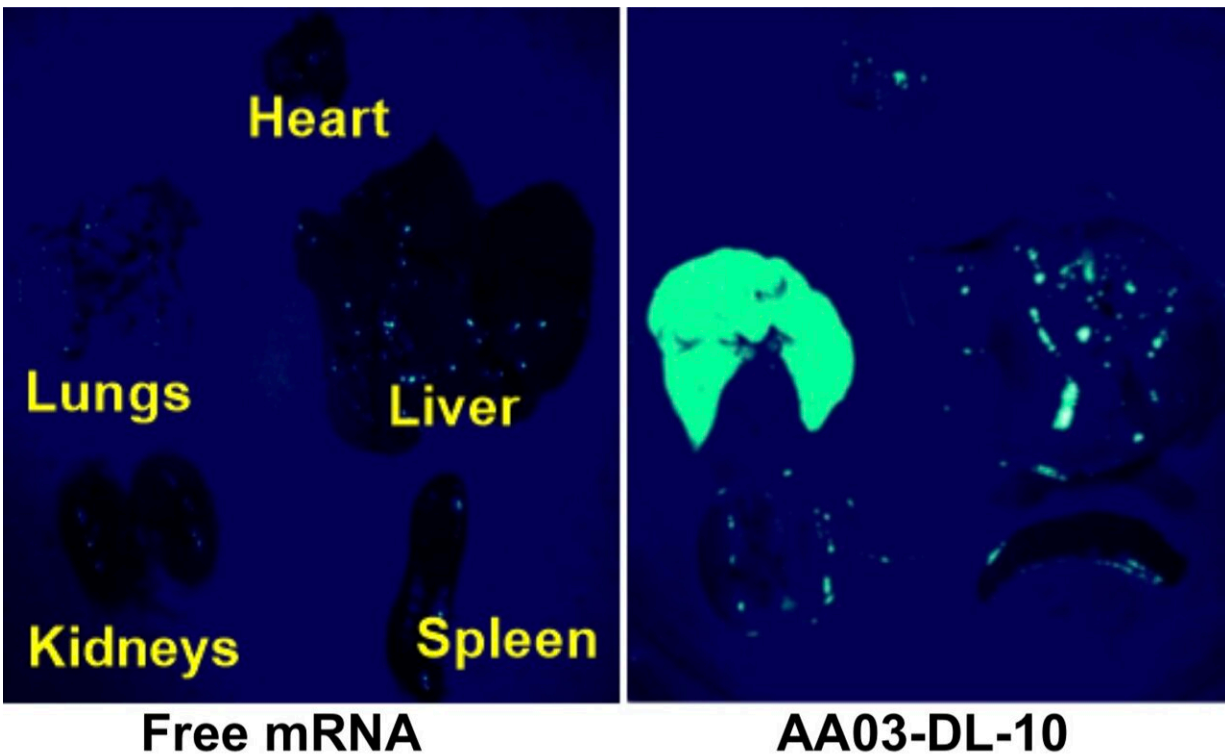
The lipomer is combined with mRNA and DMG-PEG to form mRNA-carrying nanoparticles (NPs). These NPs can bypass the liver to deliver the mRNA directly to lung cells. The scientists used mRNA encoding enhanced green fluorescent protein (EGFP) for visualization of the proteins. Credit: Mahmoud M. Abd Elwakil, et al., *Materials Horizons*. June 7, 2021

Hokkaido University researchers in Japan created and tested a library of lipid-based compounds to find a way to safely and effectively deliver RNA drugs to the lungs. Their analyses, published in the journal *Materials Horizons*, pinpointed a lipid polymer that might in the future be used to treat acute respiratory distress syndrome, pulmonary hypertension and lung cancers.

The COVID-19 pandemic response has made us all familiar with RNA vaccines that carry [genetic code](#) into [cells](#) to prompt the production of

virus proteins that trigger our protective immunity. RNA drugs are showing great potential for treating a large variety of other diseases by similarly directing protein production inside cells, without the need for inserting or deleting DNA. But scientists face several challenges in their safe delivery to targeted cells. One successful but complex approach involves carrying the RNA codes inside nanoparticles covered with compounds, called targeting ligands, that can bind to specific cells. This has worked for targeting [liver cells](#).

Hokkaido University pharmaceutical scientist Hideyoshi Harashima and polymer chemist Toshifumi Satoh led a team of researchers in developing and testing a library of  $\epsilon$ -decalactone-based compounds, lipids that could bypass the liver—which degrades toxins and foreign substances—and specifically deliver RNA code into the lungs. Harashima recently received the Høst-Madsen Medal, the highest scientific honor awarded by The International Pharmaceutical Federation (FIP).



Free mRNA did not target any organ (left), while NPs containing lipomer AA03-DL-10 specifically transported mRNA encoding EGFP to the lungs and produced green fluorescent proteins eight hours after intravenous administration into mice. Credit: Mahmoud M. Abd Elwakil, et al., *Materials Horizons*, June 7, 2021

The scientists worked with two closely related ring-shaped compounds:  $\epsilon$ -caprolactone and  $\epsilon$ -decalactone. Lipid nanoparticles (NPs) containing these lactones were previously shown to accumulate in lungs. They were subjected to ring-opening reactions with one of eleven amino alcohols. The resulting products were further classified on the basis of the molecular weight of each arm. The products were combined with mRNA and another compound called DMG-PEG to form mRNA-carrying NPs. NPs made from  $\epsilon$ -caprolactone were unstable, so the team proceeded solely with the NPs from  $\epsilon$ -decalactone.

The team tested the delivery of RNA-carrying  $\epsilon$ -decalactone NPs first into laboratory cancer cells and then intravenously into mice. They used mRNA encoding enhanced green fluorescence protein (EGFP) to identify the destination of the NPs. Ultimately, they found that  $\epsilon$ -decalactone combined with a linear amino alcohol called AA03 produced the best result. The investigations showed that NPs containing this lipomer were able to largely bypass the liver and carry the RNA material specifically into the lungs. The NPs were engulfed by the cell membrane and the RNA content was released into the cytoplasm of the [lung](#) cells.

"We showed that expanding the chemical space of smart materials could enable the fabrication of nanoparticles for hard-to-reach targets without the need for targeting ligands," says Harashima. "Designing combinatorial libraries that provide diverse  $\epsilon$ -decalactone lipomers could be an easy and scalable strategy for the development of next-generation gene therapies for organs beyond the liver."

**More information:** Mahmoud M. Abd Elwakil et al, Engineered  $\epsilon$ -decalactone lipomers bypass the liver to selectively in vivo deliver mRNA to the lungs without targeting ligands, *Materials Horizons* (2021). [DOI: 10.1039/d1mh00185j](https://doi.org/10.1039/d1mh00185j)

Provided by Hokkaido University

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