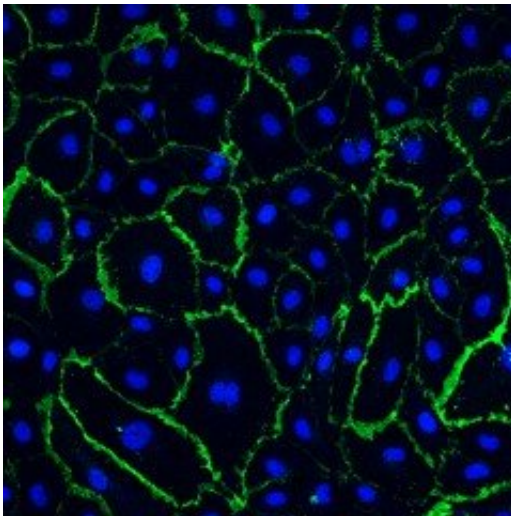


Clinical trial in a dish: A novel strategy for drug development in heart disease

July 30 2020



Healthy control iPSC-derived endothelial cells. Staining: CD31 (green); Dapi for nucleus (blue). Credit: Sayed et al.

Dilated cardiomyopathy (DCM) is a disease of the heart muscle that results in decreased function of the left ventricle, the main pumping chamber. This decreases the heart's ability to pump blood, which can lead to irregular heartbeats (arrhythmias), blood clots, heart failure, or sudden cardiac death. While there are several contributing factors to the development of DCM, it is known that up to one-third inherit it from their parents: familial DCM. Familial DCM is known to be caused by changes in different genes, the most common being variants (mutations) in the gene that encodes what is known as lamin A and C (LMNA);

DCM caused by these variants is usually referred to as cardiolaminopathies. Although LMNA-associated DCM (or cardiolaminopathy) accounts for 6% of all familial cases, there are no targeted drug to prevent onset or progression of the disease, and the mechanisms that result in the cardiolaminopathy are not known.

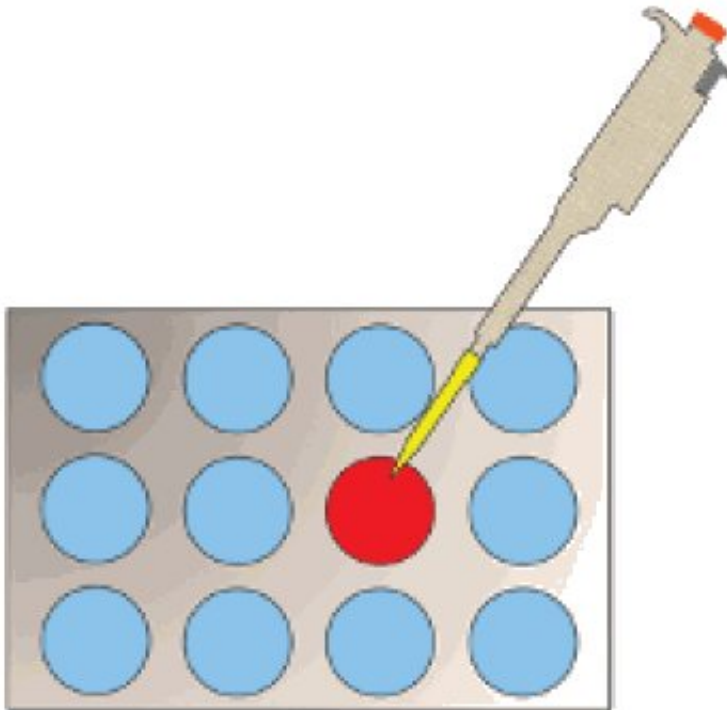
Looking beyond the heart

Stanford Cardiovascular Institute scientists, Nazish Sayed, MD, Ph.D., and Joseph Wu, MD, Ph.D., Director of the Cardiovascular Institute, addressed both important needs in a paper recently published in *Science Translational Medicine*. They reasoned that as lamins are expressed in all [differentiated cells](#), and dysfunction of non-myocytes such as endothelial cells (ECs) can contribute to the progressive failure of the [heart](#) muscle, there is opportunity to understand heart disease by looking beyond the heart muscle. They studied a large family cohort spanning four generations that had LMNA-associated DCM. Induced [pluripotent stem cells](#) (iPSCs), adult [blood](#) cells that were reprogrammed into stem cells, made from this family (members with DCM and healthy controls) allowed researchers to directly address questions of mechanism, especially in the tissue lining blood vessels and heart (endothelium). As iPSCs can be turned into any other kind of cells, the team generated iPSC-derived [endothelial cells](#) (iPSC-ECs) and were able to show and confirm a direct link between the change/mutation in the LMNA and the narrowing of large blood vessels (endothelial dysfunction). Narrowing of blood vessels contributes to decreased ability to pump blood.

Teaching old drugs new tricks

Using next-generation sequencing, Stanford CVI researchers were able to identify a protein, Krüppel-like Factor 2 (KLF2) that plays a crucial role in mediating EC dysfunction in these recruited cardiolaminopathy family members. With the target identified, the team leveraged the iPSC

platform to conduct a "clinical trial in-a dish" to identify drugs that could alleviate their observed clinical EC dysfunction. Importantly, they found that lovastatin, a 1st-generation statin, can help restore endothelial cell function in cells by increasing the expression of KLF2. "This was a significant discovery as it allowed us to repurpose lovastatin, an already FDA-approved drug for cardiometabolic disease and allowed us to move from cells to patients," said Dr. Wu. Two members of the LMNA family were recruited, and the therapeutic effects of lovastatin were tested for up to 18 months. They found that as early as six months after starting oral lovastatin regimen, both the patients showed improvement in their clinical endothelial function.



Clinical trials in-a dish: A revolutionary strategy for drug development, for drug safety, and for precision medicine using human tissues. Credit: Stanford University Medical Center

Fixing the plumbing

Similar to the plumbing in our house, the human body has an intricate piping system that is directly or indirectly connected to the heart, carrying blood to and from the heart. From the time when Leonardo da Vinci sketched the first diagram showing the distribution of the heart and its blood vessels in the fifteenth century, it is well established that the vasculature, lined by the endothelium as a diaphanous film of tissue, make for a vast network of pipes, which when damaged or dysfunctional, can lead to heart problems. "With this analogy in mind, we used our iPSC platform to test whether improving EC function in LMNA iPSC-ECs can improve heart muscle function for these patients," said Dr. Sayed. First, the team tested this in-a dish and found that lovastatin treatment of LMNA endothelium improved function of LMNA heart [cells](#) when cultured together by improving the contractile properties of the heart muscle. Next, they tested this in patients, but found that even though lovastatin improved EC function in these patients, it failed to reverse the already set-in cardiac pathology. However, this does lend the intriguing possibility that improving endothelial cell function with lovastatin at an early stage, prior to cardiac symptoms, could delay onset of LMNA-associated DCM.

This *Science Translational Medicine* paper has important findings for understanding and treating a common form of DCM, and provides a potential therapeutic for delaying progression. Equally important, they demonstrate a workflow for identifying and validating potential drug treatments or patients that includes iPSCs (clinical trial in-a-dish) and patients.

More information: Nazish Sayed et al., Clinical trial in a dish using iPSCs shows lovastatin improves endothelial dysfunction and cellular cross-talk in LMNA cardiomyopathy, *Science Translational Medicine* (2020). [DOI: 10.1126/scitranslmed.aax9276](https://doi.org/10.1126/scitranslmed.aax9276)

Provided by Stanford University Medical Center

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