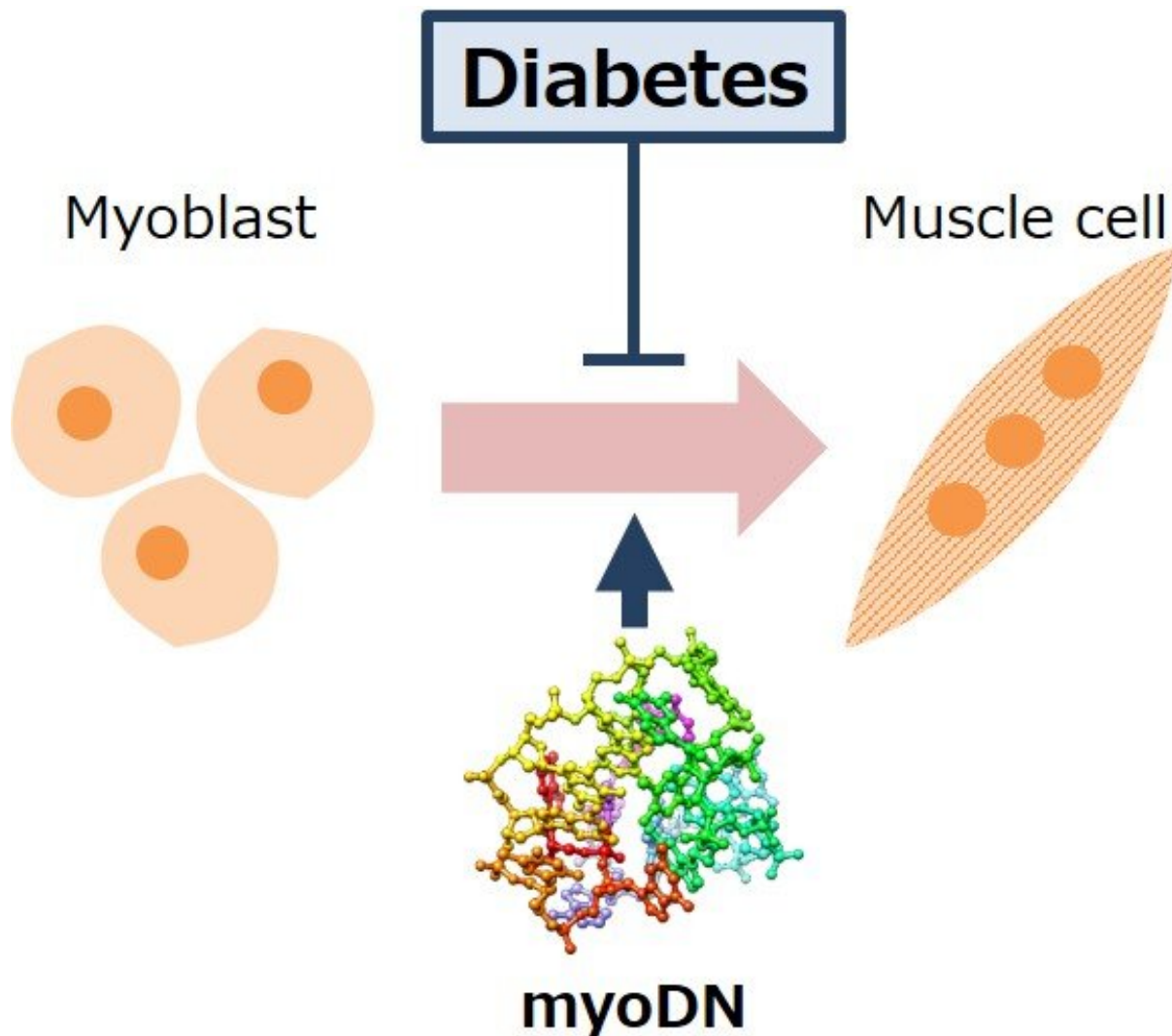


Skeletal muscle loss exacerbated by diabetes improved with oligo DNA

May 26 2021

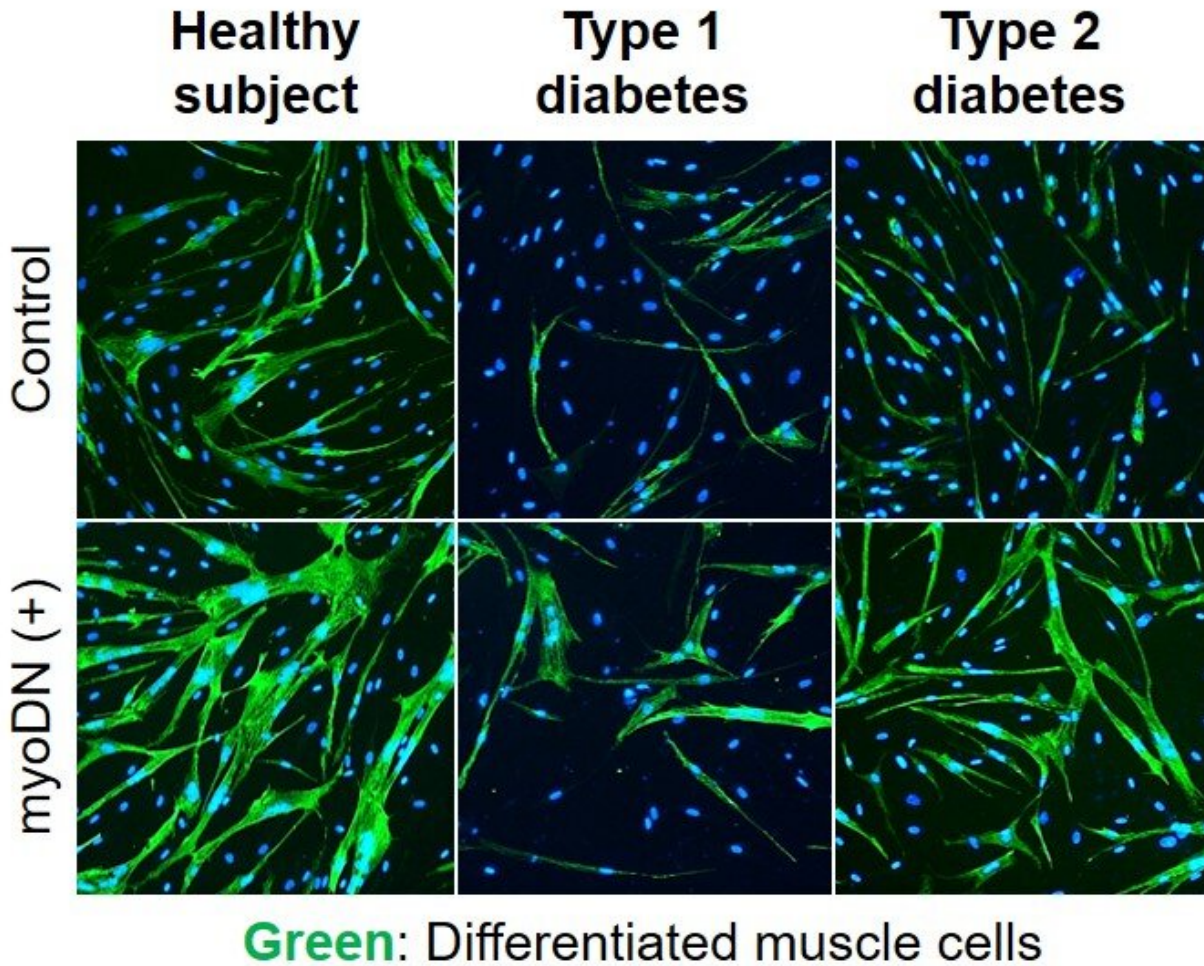


Application of myoDN to improve the muscle differentiation deteriorated by diabetes mellitus. Credit: Tomohide Takaya, Shinshu University

Skeletal muscle is the largest organ in the body. It accounts for 30 to 40% of body weight and is responsible for multiple functions such as energy metabolism and heat production. However, skeletal muscle mass is reduced in some diabetics, and that muscle loss correlates with mortality. It has been reported that the differentiation of myoblasts, which are the muscle precursor cells, is reduced in diabetic patients, and this is thought to be one of the underlying causes of muscle loss.

Assistant Professor Tomohide Takaya of Shinshu University recently reported that oligo DNA derived from the genome sequence of lactic acid bacteria promotes differentiation into [skeletal muscle](#) by binding to the target protein nucleolin in myoblasts. This "muscle-forming oligo DNA" (myoDN) has potential for use in nucleic acid drugs for various muscle diseases. In this study, a group examined whether myoDN improves the differentiation of myoblasts, which is exacerbated by diabetes.

Myoblasts collected from healthy subjects, patients with type 1 diabetes, and patients with type 2 diabetes were used for this experiment. Compared to healthy subjects, diabetic myoblasts had a reduced ability to differentiate into skeletal muscle. Remarkably, administration of myoDN improved skeletal muscle differentiation exacerbated by diabetes. The group also found that glucose and [fatty acids](#), which are increased in the blood of [diabetic patients](#), induce an inflammatory response in myoblasts, and myoDN suppresses these inflammatory responses.



myoDN recovers the impaired muscle differentiation of patients with type 1 and type 2 diabetes. Credit: Modified from Nakamura et al., *Frontiers in Physiology* (2021) 12: 679152

It is thought that myoDN is a nucleic acid molecule that is effective in reducing muscle mass associated with diabetics. Currently, there is no effective therapeutic agent for muscle wasting, and the action of myoDN, which promotes skeletal muscle differentiation, is unique. myoDN is effective for healthy subjects with various backgrounds and type 1 and type 2 patients. On the other hand, analysis using multiple

myoblasts revealed that there are individual differences in the effects of myoDN, and issues for clinical application were also clarified.

In the future, it will be necessary to expand the scope and test in order to clarify the cause of [individual differences](#). Diseases other than diabetes such as cancer and heart failure also cause muscle wasting. The group hopes to verify whether myoDN promotes skeletal muscle differentiation in these diseases as well. The ultimate goal is to apply myoDN as an effective nucleic acid drug for muscle wasting associated with various diseases.

More information: Shunichi Nakamura et al, Myogenetic Oligodeoxynucleotide (myoDN) Recovers the Differentiation of Skeletal Muscle Myoblasts Deteriorated by Diabetes Mellitus, *Frontiers in Physiology* (2021). [DOI: 10.3389/fphys.2021.679152](https://doi.org/10.3389/fphys.2021.679152)

Provided by Shinshu University

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