

# Seminar

## Role of NAMPT and NAD<sup>+</sup> precursors for skeletal muscle health and disease



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**Date:** 4 February 2019

**Time:** 11:15

**Place:** Lykeion

**Host:** Nils J. Færgeman, BMB, SDU.

**Abstract:** NAD<sup>+</sup> is important in energy metabolism and as a substrate for enzymes controlling a wide range of metabolic processes that maintain cellular homeostasis. Over the past years, we have investigated the role of nicotinamide phosphoribosyltransferase (NAMPT) and specific NAD<sup>+</sup> precursors in humans and mice for the ability of cells to adapt to metabolic stress. NAMPT is ubiquitously expressed and it is the enzyme mainly responsible for maintaining NAD<sup>+</sup> levels in mammalian cells. Knockout of *Nampt* in mouse skeletal muscle has recently been shown to result in severe perturbations in NAD<sup>+</sup> levels and to lead to fiber degeneration and reduced muscle function. Moreover, we and others have shown that

NAMPT levels decrease with age in humans, and that exercise training is a potent intervention to circumvent this decline in NAD<sup>+</sup> salvage capacity. These findings highlight the importance of maintaining NAD<sup>+</sup> levels in skeletal muscle, and they suggest that strategies to boost NAD<sup>+</sup> availability in skeletal muscle could be a viable way to alleviate conditions related to skeletal muscle dysfunction. Using novel muscle-specific Nampt knockout mouse models and samples obtained from a clinical trial with the NAD<sup>+</sup> precursor nicotinamide riboside (NR), we are further detailing the role of the NAD<sup>+</sup> salvage systems in skeletal muscle. Data from experiments using these models will be presented and discussed.