The Keap1/Nrf2 Axis Regulates the Quiescence/Activation States of Adult Muscle Stem Cells

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Adult muscle satellite cells (MuSCs) are quiescent under normal homeostatic conditions. Upon muscle injury, MuSCs exit quiescence, re-enter the cell cycle to proliferate, and then differentiate and fuse to drive muscle regeneration. Although several factors and pathways are already known to regulate quiescence maintenance of MuSCs, a comprehensive picture remains to be established. Here, we reveal that loss of Keap1 in adult MuSCs impairs quiescence maintenance in a sex-dependent manner. In female mice, the loss of Keap1 in MuSCs leads to their spontaneous activation and gradual depletion. In male mice, Keap1 deletion in MuSCs drives cells into a GAlert-like state. Detailed mechanistic insights will be presented during my talk. Our study reveals an unexpected role of the Keap1/Nrf2 axis in controlling the quiescence/activation states in adult MuSCs.