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Generating reference values on mitochondrial respiration in permeabilized muscle fibers

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Introduction

Permeabilized muscle fibers (pfi) are widely used to assess mitochondrial (mt) respiratory function in skeletal muscle of various models in different physiological and pathological conditions. Facing the numerous data available for mt-respiration from the literature, it remains challenging to determine what the right values are for a specific respiratory protocol. Moreover, mt-respiration values are highly dependent on pfi preparation, which required good technical skills.

In the frame of COST Action MITOEAGLE, one of the objectives of WG2 is the generation of reference values for mitochondrial respirometry in permeabilized skeletal muscle sample preparations. The idea is that new researchers in the field follow a reference protocol and check if their values are in an acceptable range. This approach could serve to test researchers' technical skills and therefore determine if they are proficient enough to perform their own experiments with confidence.

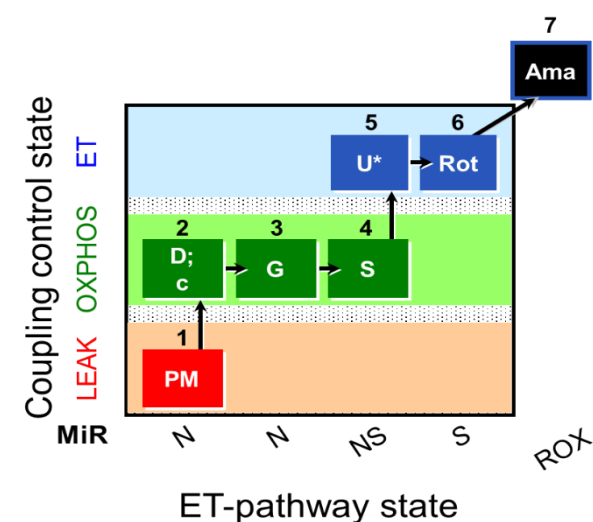
Materials and Methods

Sixteen international research groups participated to this study and received males ($N=4$) and females ($N=4$) C57BL/6J mice aged 14-16 weeks from the same provider. They performed permeabilized fibers from soleus muscle according to a common defined protocol.

Mt-respiration was measured in respiration media MiR05-Kit following the substrate-uncoupler-inhibitor titration (SUIT) protocol SUIT-008_O2_pfi_D014 represented in Figure1 [1]. Chemicals were provided by each group.

Results

Preliminary analyses of oxygen fluxes showed variability between the groups with median values for NS-OXPHOS capacity ranging from 116 to 335 pmol·s⁻¹·mg tissue⁻¹ (Figure 2). Sources of this variability are currently under investigation through the



analysis of respiratory traces to define exclusion criteria as well as a questionnaire gathering technical information and operator skills.

This unique international study has the ambition to significantly contribute to the generation of a mt-database that will help research groups in the investigation of mitochondrial physiology and pathology. Moreover, such database will help in the harmonization of mt-respiratory protocols and the resolution of the reproducibility crisis in mt-research [2].

Figure 1. Substrate-uncoupler-inhibitor titration protocol (SUIT-008 O2 pfi D014). Sequential titrations and respiratory states. **1PM**: NADH-pathway (N-pathway) in the presence of 5 mM pyruvate and 2 mM malate in the N-LEAK state. **2D**: saturating ADP (N-OXPHOS state). **2c**: 10 μ M cytochrome c for evaluating the integrity of the outer mitochondrial membrane. **3G**: 10 mM glutamate as an additional NADH-linked substrate (N-OXPHOS state). **4S**: 10 mM succinate (NS-OXPHOS capacity). **5U**: uncoupler titrations to evaluate the electron transfer- (ET-) capacity (NS-ET capacity). **6Rot**: inhibition of CI by rotenone (S-ET capacity). **7Ama**: inhibition of CIII by antimycin A (residual oxygen consumption, Rox). Oxygen concentration range in the experiment was maintained between 400-250 μ M O₂.

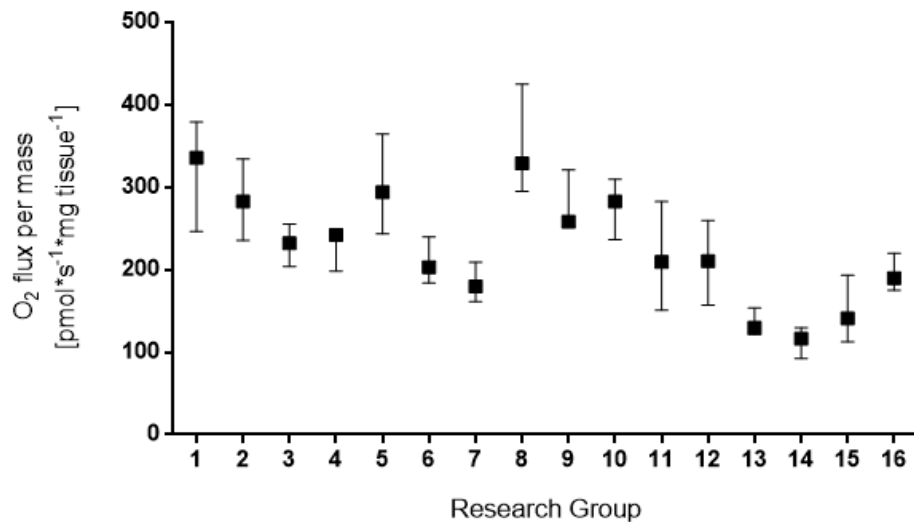


Figure 2. NS-OXPPOS capacity of permeabilized soleus muscle fibers. NS-OXPPOS capacity was measured in permeabilized soleus muscle fibers from male C57BL/6J mice by 16 research groups. Median with interquartile range show results from individual group with muscle fibers obtained from at least four soleus muscles.

References

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