

Mitochondrial Disease Community Registry: First look at the data, perspectives from patients and families

Sophia Zilber,^a and Philip E. Yeske,^{b,*}

^aData Analyst. Newton, MA ^bUnited Mitochondrial Disease Foundation, 8085 Saltsburg Road, Pittsburgh, PA 15239.

*Corresponding author emails: philip.yeske@umdf.org, sophiazilber@gmail.com

ERRATUM

The authors have submitted a correction to this paper.

The correction reads as follow:

“On page e3 of the manuscript, the authors state that “Both the PEER platform and the UMDF survey were IRB-approved by Western Institutional Review Board.” This should be corrected as follows: PEER as a registry platform was IRB-approved by Western Institutional Review Board in 2014 and oversight has since been taken over by the Genetic Alliance’s IRB. The protocol UMDF001 covering the registry and the survey discussed in this paper was under the ethical oversight of Genetic Alliance IRB from 2014 to 2019 at which time the survey was closed, and no further data collected”.

Citation: Sophia Zilber and Philip E. Yeske (2020) Mitochondrial Disease Community Registry: First look at the data, perspectives from patients and families. Mitochondrial and Metabolic Medicine 2: doi:10.9777/mmm.2020.10001

