Association of mitochondrial DNA variants with myalgic encephalomyelitis/chronic fatigue syndrome (ME/CFS) symptoms

Maureen R. Hanson1*, Zhenglong Gu2, Alon Keinan3, Kaixiong Ye3, Arnaud Germain1 and Paul Billing-Ross2

Abstract

Earlier this year, we described an analysis of mitochondrial DNA (mtDNA) variants in myalgic encephalomyelitis (ME)/chronic fatigue syndrome (CFS) patients and healthy controls. We reported that there was no significant association of haplogroups or singe nucleotide polymorphisms (SNPs) with disease status. Nevertheless, a commentary about our paper appeared (Finsterer and Zarrouk-Mahjoub. J Transl Med14:182, 2016) that criticized the association of mtDNA haplogroups with ME/CFS, a conclusion that was absent from our paper. The aforementioned commentary also demanded experiments that were outside of the scope of our study, ones that we had suggested as follow-up studies. Because they failed to consult a published and cited report describing the cohorts we studied, the authors also cast aspersions on the method of selection of cases for inclusion. We reiterate that we observed statistically significant association of mtDNA variants with particular symptoms and their severity, though we observed no association with disease status.

Keywords: Myalgic encephalomyelitis (ME), Chronic fatigue syndrome (CFS), Next-generation sequencing, Mitochondrial DNA, mtDNA, Heteroplasmy, Association, SNPs, Haplogroup, Variants

Following the publication of our paper [1] concerning mitochondrial DNA variation in ME/CFS patients and controls, Finsterer and Zarrouk-Mahjoub [2] published a commentary on our paper entitled “Is chronic fatigue syndrome truly associated with haplogroups or mtDNA single nucleotide polymorphisms?” Anyone reading this title might assume that our paper claimed a correlation of mtDNA SNPs with disease status. In fact, in describing our paper they wrote that “the study showed that CFS is associated with mtDNA haplogroups J, U and H.” However, we concluded exactly the opposite. We reported that we did not detect any statistically significant association of mtDNA variation with patient vs. control status. Instead, we reported that individuals of certain haplogroups or carrying certain mtDNA SNPs were more likely to report experiencing particular symptoms or certain severity levels of particular symptoms [1]. It is evident from the title of the commentary, as well as its content, that Finsterer and Zarrouk-Mahjoub [2] did not read our paper carefully.

The authors criticized our study for examining mtDNA variation only, rather than variation in nuclear genes encoding proteins involved in mitochondrial functions. Because the funding available allowed us only mtDNA analysis, we made no claims in the paper that all genes affecting mitochondrial performance had been analyzed. We stated clearly that we were analyzing mtDNA variation, not all genetic variation that could affect mitochondrial function. Moreover, the last sentence of our discussion sets forth the need for “further analysis of the mitochondrial and nuclear genomes.”

The authors also expressed concern that we might have missed some heteroplasmic mutations because of the use of DNA from lymphocytes, rather than from other tissues. We reported very low levels of heteroplasmy in our samples, which were comprised of DNA from whole blood of ME/CFS patients and controls. Whole blood DNA was available to us; DNA from other tissues was not. Although we were aware that tissue specificity...
of mtDNA heteroplasm exists [3], it is highly unlikely that any subjects harbored physiologically significant amounts of a heteroplasmic mutation despite our use of blood DNA, which is suitable for identifying mutations in most mtDNA-encoded diseases, with one exception. It is known that individuals with MELAS, especially as age increases, sometimes harbor far fewer abnormal mtDNAs in their blood cells than in other cell types [4, 5]. Someone with severe MELAS was unlikely to remain undetected, given the number of differential symptoms that would have allowed the ME/CFS experts, who provided the samples, to identify any such individual victim. Furthermore, while in a study of 34 Dutch families, some individuals harbored undetectable levels of the MELAS mutations in blood but significant amounts in other cell types, the majority of MELAS carriers in that study did have detectable heteroplasm in blood [6].

The authors then pointed out that we did not carry out “immune-histological nor biochemical studies of muscle biopsies” and that such studies would be valuable to identifying dysfunctional mitochondrial pathways. We do not disagree with this point, but again we made no claim that we had detected all possible such dysfunctions, given that such experiments were outside the scope of our project. While such studies might be of future interest, obtaining muscle biopsies from a patient population that usually exhibits high sensitivity to pain poses ethical considerations.

Finsterer and Zarrouk-Mahjoub [2] then ask how other causes of fatigue were distinguished from ME/CFS and then imply that our subject cohort could have included individuals with a number of other diseases. Had the authors consulted the paper we cited [7], which provides a thorough discussion of how ME/CFS subjects were selected for inclusion by a thorough discussion of how ME/CFS subjects were authors consulted the paper we cited [7], which provides a number of other diseases. Had the authors consulted this foundational paper describing the cohort [7] instead of making unwarranted negative assumptions about its quality.

**Abbreviations**

ME/CFS: myalgic encephalomyelitis/chronic fatigue syndrome; mtDNA: mitochondrial DNA; SNP: single nucleotide polymorphism.

---

**Authors’ contributions**

MRH drafted this rebuttal, which received editorial suggestions and/or approval from all authors. All authors read and approved the final manuscript.

**Author details**

1. Department of Molecular Biology and Genetics, Cornell University, Ithaca, NY 14853, USA. 2. Division of Nutritional Sciences, Cornell University, Ithaca, NY 14853, USA. 3. Department of Biological Statistics and Computational Biology, Cornell University, Ithaca, NY 14853, USA.

**Acknowledgements**

Not applicable.

**Competing interests**

The authors declare that they have no competing interests.

**Availability of data and materials**

Data described in this commentary can be found in the original publication [1]. Contact the corresponding author for any other data requests.

**Consent to publish**

Not applicable; no individually identifiable results are included.

**Ethics approval**

Our original study [1] and consent forms for participants were approved by the Cornell University Institutional Review Board.

**Funding**

This work has been funded by internal funds of Cornell University.

Received: 19 September 2016 Accepted: 4 December 2016
Published online: 20 December 2016

**References**

1. Billing-Ross P, Germain A, Ye K, Keinan A, Gu Z, Hanson MR. Mitochondrial DNA variants correlate with symptoms in myalgic encephalomyelitis/chronic fatigue syndrome. J Transl Med. 2016;14:19.
2. Finsterer J, Zarrouk-Mahjoub S. Is chronic fatigue syndrome truly associated with haplogroups or mtDNA single nucleotide polymorphisms? J Transl Med. 2016;14:182.
3. Naue J, Horer S, Sanger T, Strobl C, Hatzer-Grubwieser P, Parson W, Lutz-Bone Engel S. Evidence for frequent and tissue-specific sequence heteroplasmy in human mitochondrial DNA. Mitochondrion. 2015;20:92–94.
4. Rahman S, Poulton J, Marchington D, Suomalainen A. Decrease of 3243A→G mtDNA mutation from blood in MELAS syndrome: a longitudinal study. Am J Hum Genet. 2000;68:238–40.
5. Shanis S, Pancrudu J, Kauffmann P, Engelstad K, Jhung S, Lu J, Naini A, DiMauro S, De Vivo DC. Varying loads of the mitochondrial DNA A3243G mutation in different tissues: implications for diagnosis. Am J Med Genet A. 2004;130A:134–7.
6. de Laat P, Koene S, van den Heuvel LP, Rodenburg RJ, Janssen MC, Smeitink JA. Clinical features and heteroplasmies in blood, urine and saliva in 34 Dutch families carrying the m.3243A > G mutation. J Inherit Metab Dis. 2012;35:1059–69.
7. Klimas NG, Ironson G, Carter A, Balbin E, Bateman L, Felsenstein D, Levine S, Peterson D, Chiu K, Allen A, et al. Findings from a clinical and laboratory database developed for discovery of pathogenic mechanisms in myalgic encephalomyelitis/chronic fatigue syndrome. Fatigue Biomed Health Behav. 2013;3:75–96.
8. Horning M, Montoya JG, Klimas NG, Levine S, Felsenstein D, Bateman L, Peterson DL, Gottschalk CG, Schultz AF, Che X, et al. Distinct plasma immune signatures in ME/CFS are present early in the course of illness. Sci Adv. 2015;1:e1400121.