#### References

- Anderson S, Bankier AT, Barrell BG, de Bruijn MH, Coulson AR, Drouin J, Eperon IC, Nierlich DP, Roe BA, Sanger F, Schreier PH, Smith AJ, Staden R, Young IG (1981) Sequence and organization of the human mitochondrial genome. Nature 290:457–465
- Andrews RM, Kubacka I, Chinnery PF, Lightowlers RN, Turnbull DM, Howell N (1999) Reanalysis and revision of the Cambridge reference sequence for human mitochondrial DNA. Nat Genet 23:147
- Bandelt H-J, Lahermo P, Richards M, Macaulay V (2001) Detecting errors in mtDNA data by phylogenetic analysis. Int J Legal Med 115:64–69
- Bandelt H-J, Quintana-Murci L, Salas A, Macaulay V (2002) The fingerprint of phantom mutations in mitochondrial DNA data. Am J Hum Genet 71:1150–1160
- Derbeneva OA, Sukernik RI, Volodko NV, Hosseini SH, Lott MT, Wallace DC (2002) Analysis of mitochondrial DNA diversity in the Aleuts of the Commander Islands and its implications for the genetic history of Beringia. Am J Hum Genet 71:415–421
- Finnilä S, Lehtonen MS, Majamaa K (2001) Phylogenetic network for European mtDNA. Am J Hum Genet 68:1475–1484
- Herrnstadt C, Elson JL, Fahy E, Preston G, Turnbull DM, Anderson C, Ghosh SS, Olefsky JM, Beal MF, Davis RE, Howell N (2002) Reduced-median-network analysis of complete mitochondrial DNA coding-region sequences for the major African, Asian, and European haplogroups. Am J Hum Genet 70:1152–1171; 71:448–449 (erratum)
- Ingman M, Kaessmann H, Pääbo S, Gyllensten U (2000) Mitochondrial genome variation and the origin of modern humans. Nature 408:708–713
- Kivisild T, Tolk H-V, Parik J, Wang Y, Papiha SS, Bandelt H-J, Villems R (2002) The emerging limbs and twigs of the East Asian mtDNA tree. Mol Biol Evol 19:1737–1751
- Maca-Meyer N, González AM, Larruga JM, Flores C, Cabrera VC (2001) Major genomic mitochondrial lineages delineate early human expansions. BMC Genetics 2:13
- Macaulay V, Richards M, Sykes B (1999) Mitochondrial DNA recombination: no need to panic. Proc R Soc Lond B 266: 2037–2039
- Richards M, Macaulay V (2001) The mitochondrial gene tree comes of age. Am J Hum Genet 68:1315–1320
- Silva WA Jr, Bonatto SL, Holanda AJ, Ribeiro-dos-Santos AK, Paixão BM, Goldman GH, Abe-Sandes K, Rodriguez-Delfin L, Barbosa M, Paçó-Larson ML, Petzl-Erler ML, Valente V, Santos SEB, Zago MA (2002) Mitochondrial genome diversity of Native Americans supports a single early entry of founder populations into America. Am J Hum Genet 71:187– 192
- Torroni A, Rengo C, Guida V, Cruciani F, Sellitto D, Coppa A, Luna Calderon F, Simionati B, Valle G, Richards M, Macaulay V, Scozzari R (2001) Do the four clades of the mtDNA haplogroup L2 evolve at different rates? Am J Hum Genet 69:1348–1356

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Am. J. Hum. Genet. 72:1346-1348, 2003

# Correction: Mitochondrial DNA Variation in Amerindians

To the Editor:

We thank Yao et al. (2003 [in this issue]) for calling our attention to inconsistencies in our data reporting mitochondrial DNA variations in Amerindians (Silva et al. 2002). We reviewed the original chromatograms and resequenced all the samples (forward and reverse). On the basis of the reanalysis of the initial data and sequencing that has been repeated, we conclude that most criticisms of Yao et al. are correct. We identified two sources of problems: (a) alignment with a variant CRS (Macaulay et al. 1999) and (b) mutations missed at regions of lowquality chromatograms in one (forward or reverse) of the first sequencing. Elimination of these two problems, by a second (and, in a few cases, a third) sequencing, careful manual checking of the chromatograms, and use of the correct rCRS reference sequence (MITOMAP) eliminated the discrepancies. A summary of all 40 corrected sequences is presented in figure 1, and the general pattern is similar to that recently reported by Herrnstadt et al. (2002). The presence of a private mutation in more than one individual or the absence of a basal mutation probably represent examples of homoplasy or of reverse mutations. Extensive homoplasy within the coding region of mtDNA has been documented (Eyre-Walker et al. 1999; Herrnstadt et al. 2002) and will probably be found more often as the number of mtDNA samples sequenced increases. For instance, the group C basal mutation 9545G was found in one individual from the haplogroup A, whereas private mutation 14460G was found in two individuals who belong to haplogroups A and D, and 15670C is present in one individual who belongs to haplogroup A and two who belong to haplogroup C (Herrnstadt et al. 2002). The finding of two similar private mutations (12406A) in two individuals of the same tribe (TYR0004 and TYR0016) is probably the consequence of a single mutational event, as is the occurrence of the reverse mutation 8584 in two individuals of another tribe (YAN0669 and YAN0650).

Recalculation of the age estimates for the four founder haplogroups on the basis of the reviewed data continues

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Data matrix showing the corrected informative nucleotide positions for the 8.8-kb mtDNA segment for 40 individuals sequenced by us Figure 1

#### Table 1

Nucleotide Diversity and Age Estimates for mtDNA Belonging to the Four Founder Haplogroups of New World Natives

Haplogroup	No. of Sequences	Genetic Diversity <sup>a</sup> (SE)	Mean Age in Years <sup>b</sup> (95% CI)
А	10	0.73 (0.15)	15,398 (12,052–18,744)
В	11	0.75 (0.14)	15,819 (12,659–18,970)
С	9	0.64 (0.13)	13,520 (10,616-17,425)
D	5	0.86 (0.18)	18,144 (14,137-22,151)
Weighted mean		0.75 (0.15)	15,720 (12,366–19,074)

<sup>a</sup>  $\pi(\times 10^{-3}).$ 

<sup>b</sup> Calculated as in Silva et al. (2002).

to show similarities between the four haplogroups and does not differ significantly from the previously published values (table 1). This supports our primary conclusion in favor of a single migration wave, with a mean age for the four haplogroups of 12,366–19,074 years before present.

The revised versions of the sequences have been submitted to GenBank.

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#### **Electronic-Database Information**

The URL for data presented herein is as follows:

MITOMAP, http://www.mitomap.org (for a human mitochondrial genome database)

### References

- Eyre-Walker A, Smith NH, Maynard Smith J (1999) Reply to Macaulay et al (1999): mitochondrial DNA recombination: reasons to panic. Proc R Soc Lond B 266:2041–2042
- Herrnstadt C, Elson JL, Fahy E, Preston G, Turnbull DM, Anderson C, Ghosh SS, Olefsky JM, Beal MF, Davis RE, Howell N (2002) Reduced-median-network analysis of complete mitochondrial DNA coding-region sequences for the major African, Asian, and European haplogroups. Am J Hum Genet 70:1152–1171
- Macaulay V, Richards M, Sykes B (1999) Mitochondrial DNA recombination: no need to panic. Proc R Soc Lond B 266: 2037–2039
- Silva WA Jr, Bonatto SL, Holanda AJ, Ribeiro-dos-Santos AK, Paixão BM, Goldman GH, Abe-Sandes K, Rodriguez-Delfin L, Barbosa M, Paçó-Larson ML, Petzl-Erler ML, Valente V, Santos SEB, Zago MA (2002) Mitochondrial genome diversity of Native Americans supports a single early entry of founder populations into America. Am J Hum Genet 71:187– 192
- Yao Y-G, Macaulay V, Kivisild T, Zhang Y-P, Bandelt H-J (2003) To trust or not to trust an idiosyncratic mitochondrial data set. Am J Hum Genet 72:1341–1346 (in this issue)

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Am. J. Hum. Genet. 72:1348-1349, 2003

## Reply to Silva et al.

## To the Editor:

Silva et al. (2003 [in this issue]) have certainly improved their data by eliminating many of the errors in the current version of the data matrix, and they have admitted most of their innocent mistakes. Their efforts and atti-