

Maternal age effect and severe germ-line bottleneck in the inheritance of human mitochondrial DNA

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The manifestation of mitochondrial DNA (mtDNA) diseases depends on the frequency of heteroplasmy (the presence of several alleles in an individual), yet its transmission across generations cannot be readily predicted owing to a lack of data on the size of the mtDNA bottleneck during oogenesis. For deleterious heteroplasmy, a severe bottleneck may abruptly transform a benign (low) frequency in a mother into a disease-causing (high) frequency in her child. Here we present a high-resolution study of heteroplasmy transmission conducted on blood and buccal mtDNA of 39 healthy mother–child pairs of European ancestry (a total of 156 samples, each sequenced at ~20,000× per site). On average, each individual carried one heteroplasmy, and one in eight individuals carried a disease-associated heteroplasmy, with minor allele frequency ≥1%. We observed frequent drastic heteroplasmy frequency shifts between generations and estimated the effective size of the germ-line mtDNA bottleneck at only ~30–35 (interquartile range from 9 to 141). Accounting for heteroplasmy, we estimated the mtDNA germ-line mutation rate at 1.3×10^{-8} (interquartile range from 4.2×10^{-9} to 4.1×10^{-8}) mutations per site per year, an order of magnitude higher than for nuclear DNA. Notably, we found a positive association between the number of heteroplasmy in a child and maternal age at fertilization, likely attributable to oocyte aging. This study also took advantage of droplet digital PCR (ddPCR) to validate heteroplasmy and confirm a de novo mutation. Our results can be used to predict the transmission of disease-causing mtDNA variants and illuminate evolutionary dynamics of the mitochondrial genome.

mitochondria | heteroplasmy

The centerpiece of cellular metabolic machinery—the mitochondrion—harbors a 16.5-kb genome, mitochondrial DNA (mtDNA). Mutations in mtDNA cause over 200 diseases and contribute to diabetes, cancer, male infertility, Parkinson's and Alzheimer's diseases (1). In mammals, mtDNA mutates at high rates and is maternally inherited, making it a popular marker in evolutionary genetics (2). Despite its importance, mtDNA has drifted away from the spotlight eclipsed by nuclear DNA studies (3), and there are still gaps in our understanding of the basic aspects of human mtDNA biology. The lack of cures for diseases caused by mtDNA mutations makes it critical to understand how these mutations arise and are transmitted between generations.

Heteroplasmy, the presence of more than one mtDNA variant in a cell or a tissue, is the result of a de novo mtDNA mutation occurring in an individual or inherited through the maternal lineage. Currently there is no consensus about how prevalent mtDNA heteroplasmy is in human populations (4, 5). Such knowledge is crucial for assessing the load of mtDNA pathogenic mutations, formulating prognoses for patients with mtDNA diseases, and preimplantation diagnostics after mtDNA replacement in oocytes. Most mtDNA diseases are heteroplasmic and their phenotype depends on the allele frequency of the pathogenic variant (1).

Heteroplasmy levels can change dramatically between generations owing to genetic drift during the germ-line bottleneck—a

reduction in the number of mtDNA segregating units during oogenesis (6–8). The size of the bottleneck for mice has been evaluated to be 185 (9), yet for humans this size is difficult to obtain experimentally. Published estimates of the human bottleneck size are too broad [1–200 (10, 11)] to be useful in predicting the transmission of disease variants. Genetic drift theory predicts that a small bottleneck size will result in drastic shifts in heteroplasmy levels from a mother to her child, potentially reaching nondisease levels or levels with higher disease severity. After fertilization, mtDNA variants are distributed among cells owing to mitotic segregation—the random partitioning of mitochondria during cell divisions (12). We also lack an accurate estimate of the germ-line mtDNA mutation rate in humans, with pedigree and phylogenetic studies producing conflicting results (13, 14).

To conduct a population study of heteroplasmy transmission, we analyzed full-length mtDNA in 39 mother–child pairs using the MiSeq platform. Accounting for PCR and sequencing errors, we were able to accurately score heteroplasmy with allele frequency above 1%. With these data, we addressed (i) how common heteroplasmy is in a human population, (ii) how heteroplasmy frequency changes between tissues of the same individual and between generations, and (iii) whether maternal age at conception influences heteroplasmy occurrence in a child. We also estimated the size of the germ-line mtDNA bottleneck and the germ-line mutation rate via population genetics modeling of heteroplasmy. Focused on

Significance

The frequency of intraindividual mitochondrial DNA (mtDNA) polymorphisms—heteroplasmy—can change dramatically from mother to child owing to the mitochondrial bottleneck at oogenesis. For deleterious heteroplasmy, a change may transform alleles that are benign at low frequency in a mother into disease-causing alleles when at a high frequency in her child. Our study estimates the mtDNA germ-line bottleneck to be small (30–35) and documents a positive association between the number of child heteroplasmy and maternal age at fertilization, enabling prediction of transmission of disease-causing variants and informing mtDNA evolution.

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heteroplasmy inheritance in healthy individuals, our data serve as a valuable baseline to study disease associations for mtDNA and provide important insights into mtDNA evolution.

Results

Samples, mtDNA Enrichment, and Sequencing. We studied the prevalence of mtDNA heteroplasmy in blood and buccal cells from 39 mother–child pairs residing in central Pennsylvania, analyzing 156 samples (39 mothers \times 2 tissues + 39 children \times 2 tissues) grouped in sets of four (two tissues from a mother and two tissues from her child). Total genomic DNA was isolated from each sample. Haplogroup analysis conducted via Sanger sequencing of the D-loop indicated European ancestry for all families (Dataset S1, Table S1). For each sample, we amplified mtDNA from total DNA in two overlapping 9-kb fragments and sequenced them with paired-end 250-bp reads on a MiSeq instrument (Materials and Methods). This enriched for mtDNA and minimized the presence of numts (Fig. S1 and SI Materials and Methods), the majority of which are short (15). Multiplexing 12 samples per run resulted in $\sim 10^6$ read pairs per sample. We confirmed the efficacy of our approach for mtDNA enrichment by applying it to Rho0 cells not harboring mitochondria (Fig. S2). To minimize potential contamination among samples, we followed previously devised guidelines (16) and used pUC18 and PhiX174 as spike-ins (Materials and Methods).

Heteroplasmy Discovery. The sequencing read pairs were mapped to human mtDNA and nuclear genomes. On average, 85% of the reads per sample mapped to mtDNA (97% for samples without a spike-in; Fig. S3). We then applied our heteroplasmy discovery pipeline (Materials and Methods and Fig. S4). We required both reads of the pair to map uniquely, and in a proper orientation, to the reference mtDNA (Fig. S5). To compute minor allele frequency (MAF) at each site for each sample, we used bases with sequencing quality ≥ 30 from reads with mapping quality ≥ 20 (other thresholds led to almost identical results; Fig. S6). The mean sequencing depth per sample (averaged across sites) was $19,789 \times \pm 770 \times$ (mean \pm SEM). Tabulating depth on a per-site basis, 90% of sites in the mitochondrial genome were sequenced at $\geq 7,858 \times$ per sample (Fig. S7). The proportion of spike-in reads aligning to their respective references was as anticipated, suggesting absence of contamination among adjacent samples (Fig. S3).

In the search for heteroplasms, we first identified sites with MAF $\geq 1\%$ in individual samples. The sequencing depth per site required to detect true heteroplasms with MAF $\geq 1\%$ over the base quality error (0.1% for Phred score 30) with 99% power is $839 \times$ per site (one-sided power calculation for one-sample proportion test). Conservatively, we rounded up the depth requirement to $1,000 \times$. A detection limit of MAF $\geq 1\%$ allows detection of inherited and de novo variants that pass through the bottleneck if its size is < 100 . Mutations with lower frequency are accounted for with population genetics modeling (discussed below). After filtering for potential sequencing artifacts (Dataset S1, Table S2 and Materials and Methods), we identified 174 point heteroplasms distributed among 100 quartets—groups of site-specific heteroplasmy frequencies from two tissues of a mother and two tissues from her child (Dataset S1, Table S3). These heteroplasms were found in 31 families (eight families had no heteroplasms).

Statistical Validation of Point Heteroplasms. To validate heteroplasms, we used a novel statistical method that identifies heteroplasmy sites via a likelihood function accounting for instrument sequencing and mapping errors (SI Materials and Methods). With this method, all 174 point heteroplasms were significant ($P < 0.0003$ for each site; Dataset S1, Table S4). Additionally, the allele counts for all 174 heteroplasms tested were significant ($P < 0.0003$ for 172 sites, and $P < 0.03$ for the remaining two sites; Table S4) based on the variability observed for the same position among all samples (17).

Experimental Validation of Point Heteroplasms. We used Sanger sequencing to test all point heteroplasms with MiSeq MAF $\geq 10\%$ (Sanger method detection limit, Fig. S84 and Dataset S1, Table S5)

in at least one sample per family and the corresponding sites from the other samples from the same family (we always sequenced newly amplified fragments). In total, we examined 21 sites \times 4 samples = 84 sites, 44 of which had MiSeq MAF $\geq 10\%$ (Dataset S1, Table S6). The presence of heteroplasmy was successfully validated in all these 44 cases. Thus, our false-positive rate for detecting heteroplasms with MAF $\geq 10\%$ is below 0.023 (1/44). The MAFs from the MiSeq and Sanger methods were well correlated ($R^2 = 75\%$; Fig. S9A).

A set of point heteroplasms with MiSeq MAF $< 10\%$ was analyzed with droplet digital PCR (ddPCR) (18), which can detect heteroplasms with MAF $> 0.2\%$ (Fig. S8B and C and Dataset S1, Table S7). Here we analyzed point heteroplasms with MiSeq MAF between 1% and 10% in at least one sample per family and the corresponding sites from the other samples of the same family, a total of 10 sites \times 4 samples = 40 sites, 18 of which had MiSeq MAF $\geq 1\%$ (Fig. S9B and Dataset S1, Table S8). When we assayed the original amplicons used for MiSeq sequencing, the presence of heteroplasmy was confirmed in all these 18 instances. However, when we reamplified mtDNA from these 18 samples, in two instances (site 11,616 in M203C5-ch and site 11,825 in M210-bl) ddPCR did not confirm the presence of heteroplasmy. Repeating amplification and ddPCR for a third time again did not detect heteroplasmy (Dataset S1, Table S8), suggesting PCR errors in the amplicons sequenced with MiSeq. Thus, our false-positive rate for detecting heteroplasms with MAF between 1% and 10% is 0.11 (2/18). Overall, the MAFs from the MiSeq and ddPCR methods were well correlated for the sequenced and newly amplified amplicons ($R^2 = 95\%$ and 79% , respectively; Fig. S9B).

Distribution of Point Heteroplasms. After removing two sites that failed to validate with ddPCR (discussed above), we retained 172 point heteroplasms in 98 quartets (Dataset S1, Table S3). We assumed that these 98 point mutations arose independently in the families analyzed (or in their maternal ancestors). Point heteroplasms were found at 87 unique mtDNA positions (Fig. S10). Six positions (185, 189, 214, 215, 16,093, and 16,183) were heteroplasmy in multiple families (four, three, three, two, three, and two families, respectively; Dataset S1, Table S3), likely owing to high mutation rate at the D-loop (13). Each mother on average carried 1.13 ± 0.04 heteroplasms in her blood. This value was similar for maternal buccal tissue and for buccal and blood tissues of children (Fig. S11). Among the 98 point heteroplasms 96 were transversions, resulting in a transition-to-transversion ratio of 48 (Fig. S10 and Dataset S1, Table S3).

There were significantly more and significantly fewer heteroplasms in the D-loop and protein-coding regions, respectively, than expected based on their length and assuming equal propensity to harbor a heteroplasmy along mtDNA (Table 1). A high mutation rate for the D-loop had been documented previously (13). The nonsynonymous-to-synonymous rate ratio (d_N/d_S) at (concatenated) protein-coding genes was significantly lower than 1 ($P = 5 \times 10^{-3}$; Fisher's exact test; Dataset S1, Table S9), suggesting purifying selection (19). Most nonsynonymous mutations were predicted to affect protein function (Dataset S1, Table S10).

Disease-Associated Mutations and Mutation Burden. Eight families harbored eight point heteroplasms (one per family) that can cause disease when present at high allele frequencies (Table 2). Among 39 mothers, 5 (or 1 in 8) were carriers of disease-associated mtDNA mutations in at least one of the two tissues analyzed. Mutations at four of the eight sites are associated with disease when homoplasmic for the mutant allele (20–23); however, in our data these were heteroplasmic (Table 2). For the other four of the eight sites above, disease can develop even when mutant alleles are heteroplasmic—with disease severity depending on the allele frequency. For A1555G, G13708A, and G3242A mutations, the allele frequencies were much lower than disease-associated frequencies (Table 2) (24–26), suggesting lack of symptoms. Mutations at tRNA-Leu sites 3,242 and 3,243 contribute to several mitochondrial diseases (25, 27); notably, allele frequencies observed at site

Table 1. The distribution of point heteroplasmies among mtDNA regions

Region	bp	Observed	Random	Neutral
D-loop	1,122	34	6.6*	7.9*
tRNA	1,508	6	8.9	10.6
rRNA	2,513	13	14.9	17.7
Protein S	2,834	20	16.8	20.0
Protein N	8,533	25	50.5*	60.2*
Intergenic	88	0	0.5	0.6
Total	16,569 [†]	98	N/A	197

The observed numbers of heteroplasmies and the numbers expected under random and neutral (based on the frequency at synonymous sites) expectations are shown. The numbers of synonymous (S) and nonsynonymous (N) sites were calculated with the Nei–Gojobori method. N/A, not applicable. *Significantly different from observed ($P < 0.05$, test comparing two proportions). [†]Owing to overlapping annotations and exclusion of stop codons, the sum of base pairs in regions does not sum up to the overall length of mtDNA.

3243 in the child of family M512 (Table 2) were comparable to those observed in mitochondrial encephalomyopathy, lactic acidosis, and stroke-like episodes patients (27).

Transmission of Heteroplasmies. Considering heteroplasmy-containing quartets (Dataset S1, Table S3), we used the presence of heteroplasmy with MAF $\geq 1\%$ in at least one sample from a family as a “prior” to support the existence of heteroplasmy at the same position for other samples of the same family if their MAF was $\geq 0.2\%$ (greater than or equal to twice the value of the allowed sequencing quality error of 0.1%—Phred score 30). We classified 98 quartets into five categories (Table 3 and Dataset S1, Table S3) based on whether heteroplasmies were present in (i) both tissues of a mother and both tissues of her child (category “all,” which included dramatic shifts in allele frequency from mother to child, suggesting the germ-line bottleneck); (ii) both tissues of a mother, but absent from both tissues of her child (category “mother,” suggesting loss of a variant in the child owing to the germ-line bottleneck); (iii) both tissues of a child, but absent from both tissues of a mother (category “child” with candidate germ-line de novo mutations); (iv) both tissues of a mother and one tissue of a child, or in one tissue of a mother and both tissues of a child (category “somatic loss,” suggestive of a change in MAF in tissues owing to mitotic segregation) (12); and (v) one tissue of one individual of a family (category “somatic gain” with candidate somatic de novo mutations).

Site 4191 in family M500 seemed to harbor a de novo mutation in the child. ddPCR confirmed complete absence of the mutant allele in both maternal tissues but presence in both tissues of the child (with MAF of 4.7% and 5.6% in buccal and blood tissues,

respectively; Table 3 and Dataset S1, Table S8). Examination of hair from the same individuals indicated homoplasmy in the mother and MAF of 1.2% in the child (Dataset S1, Table S11), confirming emergence of a novel allele.

The changes in allele frequencies between tissues of an individual, or between two generations, tabulated for our 98 quartets (Dataset S1, Table S3) followed an approximately normal distribution with mean zero (Fig. S12), corroborating the action of genetic drift as the major force affecting heteroplasmy allele frequencies (28). A decrease in allele frequency for a variant from mother to child will be indicative of purifying selection (29). When we plotted the relative change in allele frequency between mothers and children (Fig. S13), such a decrease was significant for nonsynonymous sites ($P = 9.54 \times 10^{-7}$, one-tailed nonparametric sign test), suggestive of purifying selection. Consistent with selection operating against transmission of nonsynonymous mutations, we observed a significantly lower proportion of these mutations among transmitted heteroplasmies (5 out of 43, or 12%, in “all” and “somatic loss” categories) compared with untransmitted heteroplasmies (8 out of 22, or 36%, in “mother” category; $P = 0.025$, Fisher’s exact test).

Comparing MAFs Between Tissues and Generations. Compared with mtDNA in maternal tissues, mtDNA in child tissues underwent fewer mitotic segregations and replications and was exposed to mutagens for a shorter time. Therefore, we expect heteroplasmy allele frequency at a site to diverge less in the tissues of a child than in those of a mother. Indeed, the allele frequencies for the sites tabulated as quartets (Dataset S1, Table S3) were more strongly correlated between the two tissues for children ($R^2 = 92\%$, Fig. 1A) than between the two tissues for mothers ($R^2 = 49\%$, Fig. 1B). Stronger correlation for allele frequencies was observed between two tissues of a mother or of a child (discussed above) than between a mother and a child for the same tissue ($R^2 = 13\%$ for buccal, Fig. 1C; $R^2 = 29\%$ for blood, Fig. 1D), likely owing to the stronger action of the mtDNA germ-line bottleneck relative to mitotic segregation.

Maternal Age Effect. We explored the relationship between age and the total number of point heteroplasmies for each individual. No association was found for children. For mothers we found a significant positive association ($P = 0.039$, 0.049, and 0.055 for combined, buccal, and blood heteroplasmies, respectively, Poisson regression; Fig. 2 and Fig. S14). Thus, older mothers accumulate more mutations in their somatic tissues, with the number of point heteroplasmies tripling over 30 y of life. Intriguingly, a positive association exists between the number of heteroplasmies in children and maternal age at fertilization ($P = 0.010$, 0.005, and 0.006 for combined, buccal, and blood heteroplasmies, respectively; Fig. 2 and Fig. S14). This suggests that older mothers accumulate more mutations in their germ-line tissues. In our dataset, there was a correlation between maternal age at fertilization and maternal age

Table 2. Disease-causing heteroplasmies

Site	Region	Wild type	Mutant	Amino acid change	Family	Mother cheek	Mother blood	Child cheek	Child blood	Diseases caused by the mutant allele	Disease allele frequency (ref.)
195	D-loop	T	C	—	M494	0.084	0.012	0.002	0.001	Bipolar disorder	1.0 (20)
1,391	12S	T	C	—	M513	0.040	0.027	0.001	0.000	HCM	1.0 (21)
1,555	12S	A	G	—	M520	0.002	0.001	0.014	0.014	Deafness	>0.52 (24)
2,352	16S	C	T	—	SC16	0.429	0.437	0.247	0.245	LVNC	1.0 (22)
3,242	Leu	G	A	—	M242	0.001	0.001	0.008	0.016	RTD	>0.49 (25)
3,243	Leu	A	G	—	M512	0.335	0.144	0.686	0.611	MELAS, MIDD; MERRF; CPEO	>0.5 (27)
12,634	ND5	A	G	I to V	M203	0.001	0.025	0.001	0.001	Thyroid cancer (cell line)	1.0 (23)
13,708	ND5	G	A	A to T	SC8	0.001	0.000	0.022	0.016	LHON	>0.92 (26)

The allele frequencies for the mutant allele in maternal and child tissues are indicated. CPEO, chronic progressive external ophthalmoplegia; HCM, hypertrophic cardiomyopathy; LHON, Leber’s hereditary optic neuropathy; LVNC, left ventricular noncompaction; MELAS, mitochondrial encephalomyopathy, lactic acidosis, and stroke-like episodes; MERRF, myoclonic epilepsy with ragged-red fibers; MIDD, maternally inherited diabetes deafness; RTD, renal tubular dysfunction.

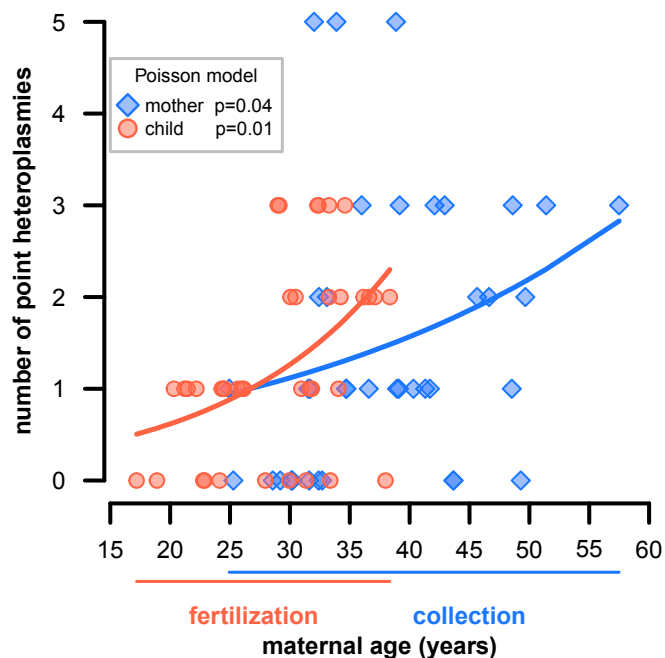


Fig. 2. Maternal age effect. The dependence of the total number of heteroplasmies (blue) in mothers on their age at collection and (red) in children on maternal age at fertilization. Poisson generalized linear model fitted curves are indicated.

pointing to the ubiquitous occurrence of heteroplasmy (4) and are in remarkable agreement with a recent analysis of the 1,000 Genomes Project data (19) as well as several smaller-scale studies (5, 11, 32–34) (Dataset S1, Table S14).

Maternal Age Effect. A positive association between an individual's age and the number of heteroplasmies in postmitotic somatic tissues had already been demonstrated (e.g., refs. 35 and 36). Here, we found evidence for it in the dividing tissues as well. Kennedy et al. (35) found that the frequency of point mutations in brain increases fivefold during 80 y of life. In our data, the number of heteroplasmies in the maternal buccal and blood tissues triples over 30 y. Likewise, with high transition-to-transversion ratio in our data, we do not find transversion-causing oxidative damage (37) to be the major driver of such mutation accumulation.

The positive association we found between maternal age at conception and the number of heteroplasmies in her child has important medical implications. The frequencies of large mitochondrial deletions (38) and the T414G mutation (39) were shown to increase in oocytes as a function of age—consistent with altered mitochondrial cytochemistry and a mutagenic environment with increased glycation and carbonyl stress in aging oocytes (40). However, the number of oocytes with defective mitochondria is significantly reduced during oogenesis (41). Our results suggest that, despite this process, some oocytes with suboptimal mitochondria (e.g., with negatively selected amino acid changes), which are more likely to occur in older women, do proceed to fertilization. This predicts an increase in mtDNA diseases in children born to older mothers—a prediction not evaluated to date—and could be one of the reasons for a lower success rate of assisted reproduction in older women (42).

Germ-Line Bottleneck Size. Our results support a severe germ-line bottleneck—with effective size of only 30–35—and are particularly striking given ~100,000 mtDNAs in mature human oocytes (12). Our findings corroborate strong shifts in heteroplasmy frequency observed in Holstein cows (43, 44) but are more robust because they are based on more accurate estimation of MAF changes at many sites, in two tissues, and for

a large number of transmissions from multiple families. Moreover, results obtained for other species are not directly applicable to humans, especially for the purposes of genetic counseling. Most previous human studies analyzed one or two sites, some of which were disease-associated, and usually a small number of transmissions (Dataset S1, Table S15). Our estimate is comparable to those in some earlier studies [e.g., 36–180 (45)], higher than in some other studies [e.g., 1–5 (10)], but substantially lower than the recently proposed estimate of 200 (11).

The number 30–35 is obtained by taking medians over effective bottleneck sizes estimated for individual transmission sites, which show a broad variation (Fig. S15). Some of this variation is random, because only one transmission was examined for each site. Selection acting at some sites might have contributed to this variation as well; however, the median bottleneck size remained very similar when nonsynonymous sites were removed. Another contributor to the observed variation in bottleneck size might be the variation among women (46, 47). Future studies examining multiple offspring per mother will allow one to evaluate the differential contribution of these factors to the variability in the bottleneck size in more detail.

Germ-Line Mutation Rate. The germ-line mutation rate estimated here for mtDNA is an order of magnitude higher than that for the human nuclear genome [1.2×10^{-8} mutations per site per generation (48)], in agreement with previous studies (1). It is similar to estimates obtained in phylogenetic studies (e.g., refs. 13 and 14) and an order of magnitude lower than estimates in most pedigree studies (13, 49–51) (Dataset S1, Table S16). In part this is due to the fact that analyzing two tissues allowed us to identify germ-line (and discard somatic) heteroplasmies (51). However, we also had to perform strict filtering of candidate heteroplasmic sites to minimize sequencing artifacts when estimating the bottleneck size—which may have led to the removal of some real heteroplasmies. Our mtDNA mutation rate estimate should therefore be seen as a “lower bound” (our bottleneck size estimate is not affected by this potential limitation). In agreement with this, our estimate is only two- and fourfold higher than estimates from mutation accumulation cell lines for *Caenorhabditis elegans* and *Drosophila melanogaster* mtDNA (9.7×10^{-8} and 6.2×10^{-8} mutations per site per generation, respectively) (52, 53); human mutation rates were shown to be approximately fivefold the rates in these species (54, 55).

Disease-Causing Mutations. The high prevalence of mtDNA disease-associated mutations found here—with one carrier in eight individuals—is similar to that reported from the 1,000 Genomes Project data (19) and has important practical implications. Indeed, as we demonstrated, the severe germ-line bottleneck can lead to drastic changes in allele frequencies between generations, potentially affecting the manifestation of 200 diseases caused by mtDNA mutations. In one instance we found a disease-associated mutation present at high allele frequencies in child tissues. Genetic background (both mtDNA and nuclear), known to significantly modulate mtDNA disease manifestation (56), may be preventing symptoms in this individual.

Materials and Methods

Sample Collection, DNA Isolation, and Sequencing. DNA from buccal and blood cells (collected under IRB 30432EP) was isolated as described (32). To determine mtDNA haplogroup, mtDNA was amplified and sequenced using the Sanger method (Dataset S1, Table S17). Before MiSeq sequencing, mtDNA was amplified in two ~9-kb amplicons (16) that were mixed at an equimolar ratio and spiked with 5% (wt/wt) of pUC18 or PhiX174 DNA, or with no spike-in. Sequencing libraries were prepared according to the customized Nextera XT protocol (57).

Experimental Validation of Heteroplasmic Sites. The primers used for heteroplasmy validation with Sanger sequencing are listed in Dataset S1, Table S17. For ddPCR, we followed the manufacturer's protocol. TaqMan probes are listed in Dataset S1, Table S18. All experiments were performed in duplicates. To assess the detection limit for ddPCR and Sanger sequencing, we

examined artificially mixed variant alleles at predetermined frequencies (SI Materials and Methods).

Preprocessing of Next-Generation Sequencing Data. Parameters and versions of all tools are listed in Dataset S1, Table S19. The sequencing read pairs were mapped to chrM and hg19 (Fig. S4). For the pair to be retained, we required both reads to (i) map to chrM, (ii) map properly in a pair, (iii) have read length ≥ 100 bp, and (iv) not form a chimeric alignment.

Identification of Point Heteroplasmic Sites. The tools Naive Variant Caller and Variant Annotator implemented in Galaxy (16) were used to extract the counts of each nucleotide per position in each strand. We selected sites with MAF $\geq 1\%$ and depth $\geq 1,000\times$. We discarded sites with MAF $< 1\%$ on one strand or with strand bias > 1 (58), low complexity regions as annotated in ref. 32, sites at

positions 3106–3107, and sites with the proportion of reads supporting an alternative base within the first and last 25 bp $> 85\%$.

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