

Mitochondrial replacement in human oocytes carrying pathogenic mitochondrial DNA mutations

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Maternally inherited mitochondrial (mt)DNA mutations can cause fatal or severely debilitating syndromes in children^{1–3}, with disease severity dependent on the specific gene mutation and the ratio of mutant to wild-type mtDNA (heteroplasmy) in each cell and tissue⁴. Pathogenic mtDNA mutations are relatively common, with an estimated 778 affected children born each year in the United States⁵. Mitochondrial replacement therapies or techniques (MRT) circumventing mother-to-child mtDNA disease transmission involve replacement of oocyte maternal mtDNA^{6–8}. Here we report MRT outcomes in several families with common mtDNA syndromes. The mother's oocytes were of normal quality and mutation levels correlated with those in existing children. Efficient replacement of oocyte mutant mtDNA was performed by spindle transfer⁸, resulting in embryos containing >99% donor mtDNA. Donor mtDNA was stably maintained in embryonic stem cells (ES cells) derived from most embryos. However, some ES cell lines demonstrated gradual loss of donor mtDNA and reversal to the maternal haplotype. In evaluating donor-to-maternal mtDNA interactions, it seems that compatibility relates to mtDNA replication efficiency rather than to mismatch or oxidative phosphorylation dysfunction. We identify a polymorphism within the conserved sequence box II region of the D-loop as a plausible cause of preferential replication of specific mtDNA haplotypes. In addition, some haplotypes confer proliferative and growth advantages to cells. Hence, we propose a matching paradigm for selecting compatible donor mtDNA for MRT.

Because mitochondrial disease can be attributed to gene mutations in mtDNA and/or nuclear DNA⁹, an important clinical challenge is to first confirm pathogenic mtDNA mutations in families eligible for MRT. We recruited four families diagnosed with Leigh syndrome and one with MELAS (mitochondrial encephalomyopathy with lactic acidosis and stroke-like episodes) and conducted genetic testing to confirm maternal inheritance (Extended Data Fig. 1a). We collected DNA from blood, skin fibroblasts and/or urine from children and mothers, and performed whole mtDNA sequencing. The first Leigh syndrome family had an affected 2-year-old child with a homoplasmic T8993G substitution encoding the ATPase 6 gene (*MT-ATP*) of mtDNA in both blood and skin fibroblast samples while her 22-year-old mother had the same mutation with 70% heteroplasmy in blood and 100% in skin fibroblasts (Fig. 1a, left). In the second related Leigh syndrome family, an affected 2.5-year-old child carried the same T8993G mutation at 95%

heteroplasmy in blood and 100% in skin fibroblasts, while in a second 1-year-old asymptomatic sibling, the mutation load was 50% in blood and 62% in skin fibroblasts. Their 23-year-old mother, who is the elder sister of the subject from the first family, carried this mutation at 13% in blood and 16% in skin fibroblasts (Fig. 1a, left). In the third Leigh syndrome family, the affected 12-year-old boy had a G13513A substitution affecting the *MT-ND5* gene at 56%, 86% and 97% heteroplasmy in blood, skin fibroblasts and urine, respectively. His asymptomatic 19-year-old brother carried the same mutation at 10%, 14% and 23% heteroplasmy levels in blood, skin fibroblasts and urine, respectively. Their 36-year-old mother also harboured the mutation at 3%, 98% and 39% levels in blood, skin fibroblasts and urine, respectively (Fig. 1a, middle). A fourth family presented with a 1-year-old child diagnosed with Leigh syndrome. However, genetic screening did not reveal any pathogenic mtDNA mutations in the child or mother. The last family was from a large, well-studied MELAS pedigree carrying a pathogenic A3243G mutation in the *MT-TL1* encoding tRNA leucine¹⁰ (Extended Data Fig. 1b). The 32-year-old mother carried the mutation at 14%, 47% and 35% in blood, skin fibroblasts and urine, respectively. All three of her children inherited this mutation (Fig. 1a, right). Thus, pathogenic mtDNA mutations were implicated in four of the five families, highlighting the importance of genetic testing for maternally inherited mtDNA diseases before MRT. Moreover, because heteroplasmy levels may vary among different tissues, it is critical to sample and test blood, skin and urine, in both mothers and children¹¹.

We also screened 11 healthy volunteer oocyte donors and confirmed that they did not carry inherited pathogenic mtDNA mutations. The mtDNA sequences and corresponding haplotypes were used for matched MRT combinations (Supplementary Table 1).

Women with mtDNA disease display live birth rates comparable to the general population⁵, suggesting normal fertility. We studied ovarian response to gonadotropin stimulation and oocyte recovery in women carrying pathogenic mtDNA mutations (carriers). Anti-Müllerian hormone levels, a measure of ovarian reserve, were lower in carriers than in healthy donors (1.1 compared to 4.8 ng ml⁻¹). Antral follicle count (AFC) was also lower in carriers compared to the healthy donors (10.3 compared to 22.3). In addition, the duration of controlled ovarian stimulation was about one day longer in the carriers and their peak blood oestradiol (E2) level before human chorionic gonadotropin (hCG) administration also trended lower. Finally, the total number of oocytes and the number of mature metaphase II oocytes retrieved were

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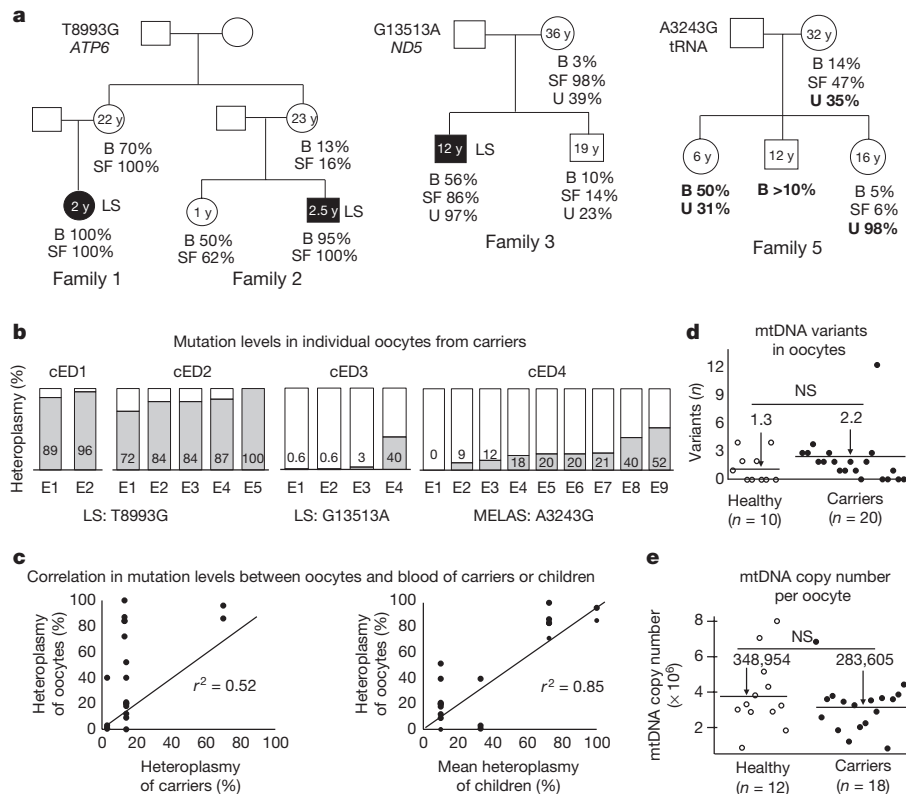


Figure 1 | Characteristics of families and retrieved oocytes in women carrying pathogenic mtDNA mutations (carriers) or wild-type mtDNA (healthy). **a**, Somatic tissue heteroplasmy levels in families (see also Extended Data Fig. 1b). Heteroplasmy levels obtained from previous reports¹⁰ are shown in bold. In all families, the heteroplasmy levels for mtDNA mutations were diverse among tissues and individuals, and associated with clinical disease. See also Supplementary Table 2. Black filled, diagnosed with a mitochondrial disease. LS, Leigh syndrome. y, years-old. Square, male; circle, female. B, blood; SF, skin fibroblasts; U, urine. **b**, Individual oocytes harboured different heteroplasmy levels. cED, carrier egg donor. **c**, The mean heteroplasmy in oocytes correlated

also lower in carriers (5.8 versus 16.6 and 3.8 versus 13.2, respectively) (Extended Data Fig. 2a–g). Of note, one carrier (cED 2) exhibited premature luteinization, as evidenced by increased progesterone levels before hCG administration (Extended Data Fig. 2h). Therefore, we only performed mutation analyses on retrieved atretic oocytes. Although the numbers in our cohort were low¹², the results suggest that mtDNA mutations could lead to a reduced ovarian response. Older age and a higher body mass index of carriers could also have affected the outcome^{13,14}. Another potential contributing factor may be the long-term hormonal contraception these women were on before ovarian stimulation.

After metaphase II spindle (nuclear genome) removal, cytoplasts (mtDNA) were used for sequencing. In the first carrier, two recovered oocytes displayed a T8993G mutation at 89% and 96% levels. In the second sibling carrier, heteroplasmy levels in five atretic oocytes ranged from 72% to 100%. In four oocytes from the third Leigh syndrome carrier, G13513A mutation levels in three eggs were low (0.6–3%) while the fourth was 40%. Finally, in nine oocytes from the A3243G MELAS carrier, no mutation was detected in one egg while the others carried 9–52% (Fig. 1b). Mean heteroplasmy levels in the blood of sibling children correlated highly with those in oocytes^{4,15} ($r^2 = 0.85$ compared to $r^2 = 0.52$ between mutation levels in the carrier's blood and oocytes, Fig. 1c). Next, we analysed whole mtDNA sequences in carrier oocytes to screen for secondary mtDNA mutations and compared to oocytes from healthy donors. Carrier oocytes contained secondary heteroplasmic mtDNA variants, however the average number per oocyte was not

with levels in existing children rather than with maternal levels. Heteroplasmy levels of carriers and children were measured in blood. r^2 , Pearson's r values. **d**, Comparable mtDNA variants between oocytes with healthy mtDNA and those carrying mutations ($P > 0.05$). **e**, Mean mtDNA copy number per oocyte in healthy controls was comparable to those carrying pathogenic mtDNA ($P > 0.05$). Karyoplasts were isolated from carrier oocytes for spindle transfer, except oocytes recovered from cED2 who exhibited premature luteinization. Oocytes from healthy egg donors were mature metaphase II. n , the number of oocytes; the number in each group is a mean; NS, not significant (**d** and **e**; analysed by t -test). Source data files are available for **c–e**.

significantly different from healthy controls (2.2 versus 1.3; $P > 0.05$) (Fig. 1d). Some of these variants were present in several oocytes from the same carrier and were also found in the carrier's blood, skin or in sibling children, suggesting recurring germline mutations. However, these variants were non-coding D-loop mutations (Supplementary Table 2). Others were found only in one oocyte within the group indicative of *de novo* mutations (45%). The majority (83%) of these secondary variants were low-level heteroplasmic (<15%) mutations (Extended Data Fig. 2i). We also measured mtDNA copy number as an indirect measure of oocyte quality and found no significant differences between healthy and carrier oocytes (mean = 348,954 compared to 283,605, respectively) (Fig. 1e). A low mtDNA copy number may play a role in diminished ovarian reserve¹⁶ but we found no correlation between oocyte copy number and anti-Müllerian hormone levels (Extended Data Fig. 2j).

Meiotic spindles (karyoplasts containing carryover maternal mtDNA) recovered from carrier oocytes ($n = 13$) were transferred into enucleated donor oocytes (cytoplasts; donor mtDNA) (Fig. 2a) while controls involved spindle transfer (ST) between healthy oocytes ($n = 36$) with preselected mtDNA haplotypes (Extended Data Fig. 3a). Spindle transfer oocytes, along with intact (non-manipulated) controls, were then fertilized by intracytoplasmic sperm injection and cultured to blastocysts. High fertilization rates, comparable to controls, were observed in both spindle transfer groups (Fig. 2a and Extended Data Fig. 3b, c). Subsequent development of diploid carrier spindle transfer zygotes to the blastocyst stage was 75%, similar to controls. Similar

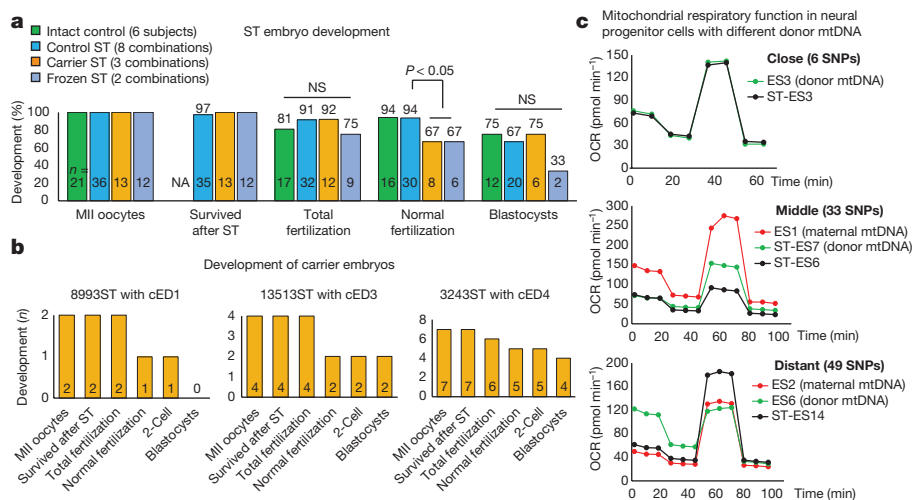


Figure 2 | Spindle transfer outcomes, embryo development and mitochondrial function. **a**, Spindle transfer (ST) embryo development following various experimental manipulations. Similar developmental outcomes were observed among groups, except normal fertilization, which was significantly lower in carrier and frozen spindle transfer groups ($P < 0.05$; one-way ANOVA). Numbers inside bars, number of oocytes or embryos; numbers on the top of bars, percentages. Combinations, mixtures of different mtDNA haplotypes between karyoplast and cytoplasm; intact control, unmanipulated embryos, fertilized by intracytoplasmic sperm injection (ICSI). NA, not applicable. **b**, Carrier spindle transfer embryos

developmental patterns were observed among carriers with different mutation types (8993ST, 13513ST and 3243ST) (Fig. 2b). To address controlled ovarian stimulation asynchrony, we conducted spindle transfer between fresh and vitrified oocytes (frozen spindle transfer). The overall fertilization rate using vitrified oocytes was comparable to fresh spindle transfer, while formation of normal diploid zygotes was reduced (Fig. 2a). No differences in fertilization were noted between combinations involving vitrified karyoplasts versus vitrified cytoplasm (Extended Data Fig. 3d).

Since miscommunication between nuclear and mitochondrial genomes may induce secondary metabolic dysfunction¹⁷, we analysed spindle transfer embryo development as a function of donor mtDNA sequence distance, as measured indirectly by the number of single nucleotide polymorphism (SNP) differences. Spindle transfer oocytes and embryos were grouped based on the SNP differences between the original (maternal) and donor oocyte mtDNA, ranging from close (6 SNPs) to middle (33 SNPs) to distant (57 SNPs) (Extended Data Fig. 3a and Supplementary Table 1). Fertilization and blastulation rates were similar among all three spindle transfer groups, suggesting that embryo development was not compromised (Extended Data Fig. 3e). Next, we established ES cell lines from spindle transfer blastocysts and differentiated them into neural progenitor cells (NPCs), cardiomyocytes and teratomas. Comparable differentiation efficiencies, mitochondrial respiratory chain enzyme activity and oxygen consumption rates were found between groups (Fig. 2c and Extended Data Fig. 4, 5). Thus, MRT in carrier oocytes supported fertilization and blastocyst development similarly to controls. Also, ES cell derivation and differentiation were unaffected by donor mtDNA genetic backgrounds. However, it is still possible that miscommunication between certain combinations of nuclear and mitochondrial genomes may occur, independent of SNP differences.

We biopsied and analysed expanded blastocysts with a comparative genome hybridization array. Aneuploidy rates were similar between spindle transfer and intact controls (Extended Data Fig. 6a). G-banding karyotype analysis in spindle transfer and control ES cells also did not reveal any significant differences (Extended Data Fig. 6b). Finally, we examined copy number variations (CNVs) in selected spindle transfer ES cell lines for subchromosomal abnormalities. *De novo* CNVs were

detected in both spindle transfer and intact controls but were deemed of uncertain clinical significance¹⁸ (Extended Data Table 1). In summary, a total of six spindle transfer blastocysts were produced from four ovarian stimulation cycles in carriers (Extended Data Table 2). These outcomes were comparable to controls. A small amount of maternal mtDNA carry-over is common during spindle transfer, resulting in low heteroplasmy in human embryos and in non-human primate offspring (typically below 2%)^{19,20}. Here, all examined spindle transfer zygotes and cleaving embryos ($n = 22$) contained more than 99% donor mtDNA (Table 1). Similar outcomes were observed in 13 out of 15 ES cell lines (87%) established from control spindle transfer blastocysts, regardless of donor mtDNA (Supplementary Table 3). However, two sibling ES cell lines (ST-ES7 and ST-ES8), generated by a combination of maternal U5a and donor H1b mtDNA (33 SNPs), displayed high levels of maternal mtDNA (81% and 94%, respectively) (Table 1). Extended passaging resulted in a complete loss of donor mtDNA and a return to the original maternal mtDNA haplotype. By contrast, another sibling ES cell line generated from maternal U5a, but with donor V3 (ST-ES9; 33 SNPs) maintained donor mtDNA.

Among the carrier spindle transfer ES cell lines, 3243ST-ES1 also demonstrated 20% maternal H49 mtDNA (Table 1). However, this maternal haplotype did not include the A3243G mutation (Supplementary Table 3). Maternal mtDNA levels in this cell line gradually increased during extended culture to 90% at passage 8 and homoplasmy at passage 10 (Fig. 3c). However, another sibling 3243ST-ES2 generated by the same maternal and donor mtDNA combinations did not show a reversal (Table 1 and Supplementary Table 3).

We expanded our study to include eight ES cell lines derived by somatic cell nuclear transfer (SCNT, NT-ES cells) that also carry donor oocyte mtDNA^{21–23} (Table 1 and Supplementary Table 4). We identified that NT-ES8 displayed a gradual increase in maternal (somatic) mtDNA from 19% at passage 2 to 100% at passage 10 (Table 1 and Extended Data Fig. 7a). We also investigated whether the reversal is specific to undifferentiated ES cells or could occur during differentiation. Several reversed and non-reversed ES cells were differentiated *in vitro* into NPCs and cardiomyocytes, and *in vivo* into teratomas (Extended Data Fig. 5). Maternal mtDNA levels were measured before and after differentiation.

Table 1 | Donor mtDNA in pre-implantation MRT embryos and ES cell lines

Treatment	Carryover maternal mtDNA %					Maternal haplotype	Donor haplotype	SNP differences			
	Pre-implantation embryos		ES cells		Total			Coding*	D-loop		
	Embryo stage	%	Lines	Passage 2–3						Passage 9–10	
Control ST	NT		ST-ES1†	≤1	≤1	H56	H2a	6	4	1	
			ST-ES2†	≤1	≤1	H2a	H56	6	4	1	
			ST-ES3	≤1	≤1	H2a	H56	6	4	1	
			ST-ES4†	≤1	≤1	H44a	H13a	25	7	10	
			ST-ES5	≤1	≤1	H1b	U5a	33	11	9	
			ST-ES6	≤1	≤1	H1b	U5a	33	11	9	
		4-cell	0.7								
		8-cell	0.9	ST-ES7	81±9	100	U5a	H1b	33	11	9
		10-cell	0.4								
		10-cell	0.4	ST-ES8	94±4	100	U5a	H1b	33	11	9
		1-cell	0.4								
		2-cell	0.5	ST-ES9	≤1	≤1	U5a	V3	33	11	9
		8-cell	0.5								
		Morula	0.4	ST-ES10	≤1	≤1	V3	U5a	33	11	9
		1-cell	0.5†	ST-ES11†	≤1	≤1	H1e	D1f	44	15	14
			0.9†								
			1.6†	ST-ES12†	≤1	≤1	H1e	D1f	44	15	14
		NA	0.0†	ST-ES13†	≤1	≤1	H1e	D1f	44	15	14
		1-cell	0.5								
		Morula	0.2								
	Morula	0.2	ST-ES14	≤1	≤1	D4a	A2g	49	18	16	
	1-cell	0.2									
	1-cell	0.5									
	1-cell	0.5	ST-ES15	≤1	≤1	A2g	D4a	49	18	16	
	1-cell	0.5									
Carrier ST	1-cell	0.8									
	3-cell	0.5	NA			H1b	H6a	21	6	11	
	3-cell	0.6									
	NT		13513ST-ES	≤1	≤1	T2b	T2	22	3	5	
	1-cell	0.5									
	8-cell	0.8	3243ST-ES1	20±13	100	H49	B2k	32	7	14	
12-cell	0.5										
	Morula	0.5	3243ST-ES2	≤1	≤1	H49	B2k	32	7	14	
SCNT	NT		NT-ES1		≤1	H1b	H56	13	5	7	
			NT-ES2		≤1	H1b	H56	13	5	7	
			NT-ES3		≤1	H1b	H56	13	5	7	
			NT-ES4		≤1	H1b	H56	13	5	7	
			NT-ES5		≤1	F1a	D1a	47	16	11	
			NT-ES6		≤1	F1a	H1b	38	13	13	
			NT-ES7	≤1	≤1	X2c	U5a	40	9	16	
			NT-ES8	19±9	100	X2c	D4a	39	12	13	

*SNPs resulting in amino acid change in protein-coding genes and residing in RNA genes. NT, not tested. NA, not applicable. ST, spindle transfer; SCNT, somatic cell nuclear transfer. †Ref. 19.

We observed that maternal mtDNA levels in the reversed 3243ST-ES1 reduced to undetectable levels in some tissues but were slightly elevated in other cell types compared to the initial 4% in undifferentiated ES cells (Fig. 3a). The maternal mtDNA was undetectable during *in vivo* and *in vitro* differentiation of the non-reversed sibling cell line 3243ST-ES2 (Supplementary Table 3). By contrast, maternal mtDNA levels increased in all tested differentiated tissues in the

reversed ST-ES7 and NT-ES8 (Fig. 3a and Supplementary Tables 3, 4). These results demonstrate that the mtDNA reversion phenomenon is not unique to ES cells and occurs during differentiation, raising the possibility that reversal to the mutant mtDNA may occur in some MRT children.

Based on our observations that specific haplotype combinations led to a reversal of the maternal mtDNA, two mechanisms were

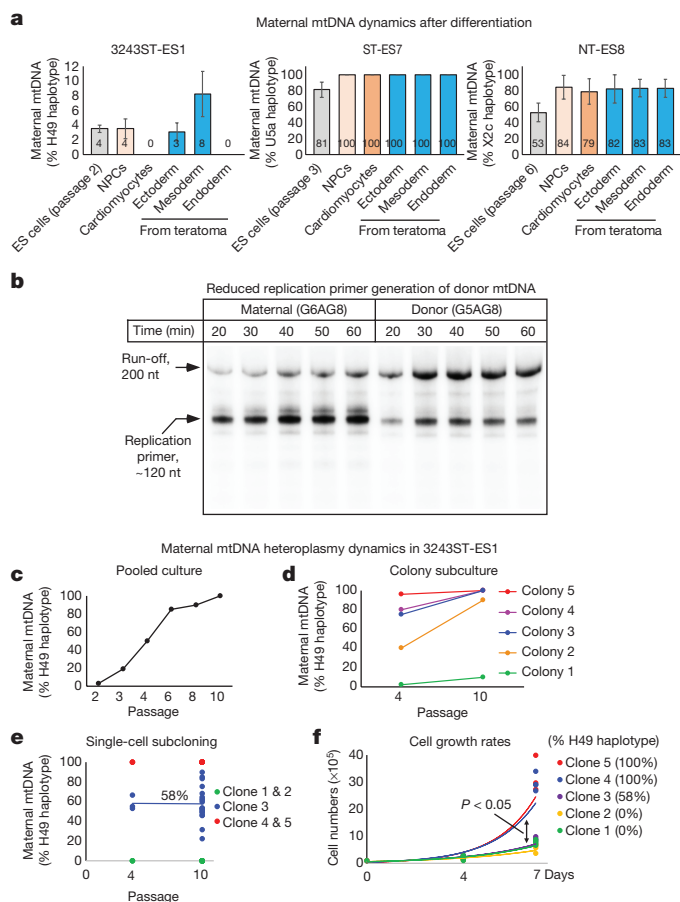


Figure 3 | Mechanisms affecting mtDNA reversal. **a**, Maternal mtDNA heteroplasmy changes in reversed 3243ST-ES1, ST-ES7 and NT-ES8 during *in vitro* and *in vivo* differentiation ($n = 1$). Bars represent a mean heteroplasmy of maternal mtDNA based on all SNP positions. Data are shown as mean \pm s.d. See also 3243ST-ES1, ST-ES7 and NT-ES8 in Supplementary Tables 3 and 4. **b**, Polymorphism in the CSBII region results in decreased efficiency of replication primer generation by mitochondrial RNA polymerase. The 311–315CCCC polymorphism is a shortening of the first G stretch in the G quadruplex region of mtDNA (G5 versus G6), responsible for transcription termination and replication primer generation. **c**, Heteroplasmy rates for the maternal mtDNA (H49 haplotype) gradually increased during extended *in vitro* culture of 3243ST-ES1 and reached homoplasmy ($n = 1$). See also Supplementary Table 3. **d**, The heteroplasmy levels also increased during ES cell colony subculture ($n = 1$). See also Supplementary Table 3. **e**, Maintenance of stable heteroplasmy in progeny of individually cultured cells. Heteroplasmy levels for maternal mtDNA were measured by amplification refractory mutation system–qPCR ($n = 3$ technical replicates) and whole mtDNA sequencing ($n = 1$). Data points at passage 10 are maternal mtDNA at individual SNPs. See also Supplementary Table 3. **f**, Faster cell growth and proliferation rates in clones containing higher maternal mtDNA heteroplasmy. $P < 0.05$; one-way ANOVA; $n = 4$ technical replicates. Source data files are available for **e** and **f**.

investigated: preferential replication of specific haplotypes and the capability of certain mtDNA endow cells with growth advantages^{24,25}. We first focused on the highly polymorphic D-loop region, called conserved sequence box II (CSBII)²⁵. It has been shown that a polymorphism in CSBII (G5AG7) affects efficiency of mitochondrial transcription termination and replication primer generation²⁵. When we analysed the donor and maternal mtDNA in a total of 26 ES cell lines comprised of 18 different haplotype combinations, we noted that two reversed sibling lines, ST-ES7 and ST-ES8 carried donor mtDNA with a G5AG8 polymorphism while the maternal haplotype was G6AG8 (Supplementary Table 5). Using *in vitro* transcription assays, we

analysed whether synthesis of the replication primer by mitochondrial RNA polymerase was affected and found that the deletion of a single guanosine residue (G5AG8 versus G6AG8) in donor mtDNA resulted in a fourfold reduction of replication primer synthesis (Fig. 3b). These results suggest that certain mtDNA haplotypes varying in the CSBII sequence provide more efficient synthesis of the replication primer that may, subsequently, confer replicative advantage. Sequence analysis did not find CSBII SNP differences in the remaining two reversed lines, 3243ST-ES1 and NT-ES8. However, a number of other D-loop polymorphisms were observed in the coreTAS region, which is also implicated in the regulation of mtDNA replication²⁶. Whereas the exact mechanisms by which these polymorphisms affect replication remain unclear, we speculate that, similar to the CSBII, they may contribute to replication bias of a particular mtDNA haplotype.

Because other replication-independent mechanisms may affect reversal, we also examined 3243ST-ES1 and NT-ES8 in whole cultures versus individual cell clones. Starting maternal mtDNA levels increased with passaging in whole cultures (Fig. 3c, d, Extended Data Fig. 7a and Supplementary Tables 3, 4). However, when we dissociated ES cells into individual cells and cultured isolated single-cell clones, the starting mtDNA levels varied in different clones but heteroplasmy did not change with passaging²³ (Fig. 3e and Supplementary Table 3). Notably, we observed substantial differences in cell proliferation and growth rates among different clones. Those with higher maternal mtDNA levels exhibited significantly faster growth rates ($P < 0.05$) than those with lower maternal mtDNA (Fig. 3f). These results suggest that certain mtDNA haplotypes confer ES cells with faster growth and proliferative advantages. Thus, in mixed cultures consisting of cells with varying degrees of heteroplasmy, a sub-population of cells with higher maternal mtDNA may overgrow cells with donor mtDNA. This possibility is independent of detectable mitochondrial activity, as complex I and complex IV enzyme activities were similar (Extended Data Fig. 7b).

In summary, among factors that might contribute to mtDNA reversal, several appear to be unlikely, including the MRT method (spindle transfer, pronuclear transfer or somatic cell nuclear transfer)^{27,28}, the presence of pathogenic mtDNA mutations and the genetic distance between donor and maternal mtDNA measured in total SNPs²⁸. A possible mechanism could be bias towards preferential amplification of mtDNA haplotypes with specific D-loop polymorphisms. Therefore, we propose selecting compatible donors harbouring CSBII or other D-loop sequences similar to the maternal mtDNA. While the number of tested combinations in our study was small, a proposed order of replication advantage in mtDNA is: H56>H1b, U5a>H1b>F1a, U5a>X2c>D4a and H49>B2k. However, stochastic or bottleneck mtDNA amplification during early embryo development may also coexist^{29,30}.

We show that some mtDNA haplotypes could affect cell growth and proliferation thereby providing selective advantage for cells with maternal mtDNA. Future applications of MRT will require additional studies evaluating compatible donor mtDNA haplotypes to avoid reversion to mutant mtDNA.

Online Content Methods, along with any additional Extended Data display items and Source Data, are available in the online version of the paper; references unique to these sections appear only in the online paper.

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Supplementary Information is available in the online version of the paper.

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Author Contributions E.K., N.M.G., P.A. and S.M. conceived the study and designed the experiments. T.Hu., P.A. and A.K. evaluated clinical genetic results and coordinated recruitment of study participants. P.A. and J.J. performed skin biopsies and blood collections. P.A., D.B., D.L. and D.W. performed ovarian stimulations and oocyte retrievals. N.M.G., R.T.H. and S.M. conducted spindle transfer, intracytoplasmic sperm injection, embryo culture and establishment of ES cell lines. E.K., R.T.H., H.M., Y.Le., Y.Li, R.A., T.Ha. and N.M.G. cultured ES cells. E.K., C.V.D., Y.Le. and T.Ha. performed teratoma studies. E.K., C.V.D., Y.Li and D.J. performed mitochondrial complex assays. J.W., P.M.R. and A.P.L. performed *in vitro* differentiation and Seahorse assays. K.A. and D.T. performed mtDNA transcription experiments. E.K., C.V.D., R.T.H. and A.K. prepared mtDNA and performed MiSeq assays. E.K. and Y.Le. performed NGS data analysis and interpretation. E.K., C.V.D. validated mtDNA mutations by Sanger sequencing. H.M. and R.T.H. performed ARMS-qPCR. X.W., S.L. and T.Hu. performed CNV data analysis. R.K. and C.C. performed the comparative genome hybridization array. S.O. performed G-banding karyotypes. E.K., J.W., D.P.W., J.C.I.B., P.A. and S.M. analysed data and wrote the paper.

Author Information Reprints and permissions information is available at www.nature.com/reprints. The authors declare no competing financial interests. Readers are welcome to comment on the online version of the paper. Correspondence and requests for materials should be addressed to S.M. (mitalipo@ohsu.edu) and P.A. (amatop@ohsu.edu).

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METHODS

Study oversight. Initial study protocol approval was granted in 2011 by the Oregon Health & Science University (OHSU) Institutional Review Board (IRB) and strictly followed the guidelines set forth by the Oregon Stem Cell Research Oversight Committee (OSCR). OSCRO guidelines permit recruitment of research gamete donors and subsequent fertilization and generation of pre-implantation embryos for research purposes. The protocol allows testing various MRT approaches prior to and after fertilization and pre-implantation. MRT embryo development can be terminated or used for ES cell derivation as a critical step in testing the safety and efficacy of novel cell and gene therapy treatments before clinical applications. The study protocol has undergone annual continuing reviews and extensions by IRB which includes biannual monitoring of study-related procedures involving human gametes, embryos and ES cells. No federal funds were used for this study in accordance with federal policies.

Participants. Healthy oocyte and sperm donors were recruited locally, via print- and web-based advertising, through OHSU Women's Health Research Unit. Oocyte donors were required to meet strict research inclusion criteria to minimize participant risk. Prior to consenting, eligible participants attended an information session, describing study goals and related clinical procedures. Carriers of mtDNA mutations were identified nationwide through print- and web-based advertising (UMDF and OHSU websites). Five families with existing children diagnosed with mitochondrial disease through MitoClinic in Cincinnati Children's Hospital Medical Center and OHSU Doernbecher Children's Hospital were identified and recruited. All families were out of state and travelled with children to OHSU for all study-related screening and procedures. Carrier oocyte donors underwent additional screening for study inclusion; strict inclusion criteria were met to minimize participant risk. Additionally, children from carrier donors provided samples for confirmation of pathogenic transmission of mtDNA mutations.

Informed consent. Written informed consent was obtained from gamete donors before any study-related procedures after a full review of study goals and clinical procedures. Consent forms for all gamete donors state that oocytes will be fertilized with sperm to generate pre-implantation embryos strictly for research purposes to develop and test novel MRT approaches. Consent forms also state that donated gametes and embryos eventually will be used for *in vitro* analyses or generation of ES cells but not for reproductive purposes. Written informed consent was obtained from parent before skin and blood donations from their underage children. As part of the consenting process, all participants provided written acknowledgement of the potential for incidental findings. If the participant requested to be informed, they were provided with some research findings by a physician collaborator and appropriate referrals were made. Patients undergoing standard IVF treatments were also contacted to donate immature oocytes that are routinely discarded as part of the IVF process. These participants were contacted before their scheduled oocyte retrievals. Written informed consent was obtained before egg retrieval.

Compensation. Research gamete donors were compensated for their time, effort, discomfort and inconvenience associated with the donation process at rates similar to gamete donors for reproductive purposes. Travel and lodging reimbursements were provided for mtDNA families travelling from out of state.

Controlled ovarian stimulation. Baseline screening was conducted including a medical history and physical exam, assessment of body mass index, anti-Müllerian hormone level and antral follicle count (AFC). Subjects underwent ovarian stimulation and oocyte retrieval employing standard IVF protocols and procedures¹⁹. All study-related procedures for oocyte donation took place at OHSU.

Spindle transfer. Spindle transfer was performed as previously described¹⁹. Briefly, meiotic metaphase II (MII) spindles were visualized under polarized microscopy and isolated with minimal cytoplasm (karyoplast, maternal mtDNA). The karyoplast was then placed in the perivitelline space of an enucleated oocyte (cytoplast, donor mtDNA) and fused using HVJ-E (haemagglutinating virus of Japan-envelope).

Oocyte vitrification. Oocyte vitrification was performed as previously described using commercially available human oocyte vitrification and thawing kits (Vitrification kit and Vitrification warming kit, Life Global).

Derivation of ES cells and culture. Spindle transfer blastocysts were freed from their zona pellucida and plated on mouse embryonic fibroblast (MEF) feeder layers on 4-well culture dishes for 5–7 days at 37°C, 3% CO₂, 5% O₂ and 92% N₂ in medium (DMEM/F12 with 10% fetal bovine serum (FBS), 10% knockout serum replacement (KSR), 0.1 mM non-essential amino acids, 1 mM L-glutamine, 0.1 mM β-mercaptoethanol, 5 ng ml⁻¹ basic fibroblast growth factor (bFGF), 10 μM ROCK inhibitor) as previously described¹⁹. Attached blastocyst outgrowths were manually minced into small pieces, re-plated onto fresh plates and maintained in knockout DMEM medium (Invitrogen) supplemented with 20% KSR, 0.1 mM non-essential amino acids, 1 mM L-glutamine, 0.1 mM β-mercaptoethanol, penicillin–streptomycin and 4 ng ml⁻¹ bFGF for further propagation and analysis. All cell lines were negative for mycoplasma contamination.

mtDNA transcription assays. *In vitro* transcription anti-termination assays were performed as described previously²⁵ using PCR-amplified templates containing region 202–481 of donor (H1b) or maternal (U5a) mtDNA. The products of the transcription reactions were resolved by 20% PAGE containing 6 M urea and visualized by PhosphorImager (GE Health).

Cell growth and proliferation assays. ES cells were adapted to grow under feeder-free conditions on Matrigel matrix in mTeSR1 medium (STEMCELL technologies)³¹. The cells were dissociated with Accutase (Life Technologies) for 2 min and approximately 10⁵ cells were seeded into each 60-mm dish. The cells were periodically harvested and counted using a Countess Automated Cell Counter (Invitrogen).

NPC differentiation and culture. A previously published protocol with minor modifications was used for NPC differentiation²². In brief, ES cells were maintained on MEFs in CDF12 medium before NPC differentiation. CDF12 medium contained DMEM/F12 (Life Technology), 20% KSR (Life Technologies), 2 mM Glutamax (Life Technologies), 0.1 mM non-essential amino acids (Life Technology), 0.1 mM β-mercaptoethanol (Life Technologies) and 4 ng ml⁻¹ FGF2 (Peprotech). ES cells were disaggregated with Collagenase IV (Life Technologies) and allowed to grow to about 40% confluence. To initiate neural induction, cells were washed twice with 1 × DPBS without calcium and magnesium (Corning Cellgro) and the medium was then changed to Neural Induction Medium 1 (NIM-1: 50% Advanced DMEM/F12 (Invitrogen), 50% Neurobasal (Invitrogen), 1 × B27 (Invitrogen), 1 × N2 (Invitrogen), 2 mM Glutamax and 10 ng ml⁻¹ hLIF (Peprotech), 4 μM CHIR99021 (Selleckchem), 3 μM SB431542 (Selleckchem), 2 μM dorsomorphin (Sigma) and 0.1 μM Compound E (EMD Chemicals Inc.)). After 2 days in culture in NIM-1 medium, cells were switched to Neural Induction Medium 2 (NIM-2: 50% Advanced DMEM/F12, 50% Neurobasal, 1 × N2, 1 × B27, 2 mM GlutaMAX and 10 ng ml⁻¹ hLIF, 4 μM CHIR99021, 3 μM SB431542 and 0.1 μM Compound E). Cells were further cultured in NIM-2 for 5 days with daily medium change. On the last day in NIM-2 medium, cells were treated overnight with 10 μM Y27632 (Selleckchem) and 20–30 'dome'-shaped colonies were manually picked and digested with Accumax (Innovative Cell Technologies) for 10 min at 37°C. After Accumax treatment cells were gently disaggregated into single cells and re-plated onto Matrigel-coated 6-well plates at a density of about 3.5 × 10⁴ per cm² in Neural Progenitor cell Maintenance Medium (NPM: 50% Advanced DMEM/F12, 50% Neurobasal, 1 × B27, 1 × N2, 2 mM GlutaMAX, 10 ng ml⁻¹ hLIF, 3 μM CHIR99021 and 2 μM SB431542) supplemented with 10 μM Y27632. NPCs were maintained on Matrigel-coated dishes in NPM. NPCs were passaged when they reached around 70% to 80% confluence using Accumax and replated at a density of 3 × 10⁴ per cm² with daily medium change. For the initial six passages, NPCs were pre-treated with 10 μM Y27632 overnight before and during cell splitting. The study was randomized, and the investigators were blinded to treatment allocations in collaborators' laboratories.

Cardiomyocyte differentiation. Cardiomyocyte differentiation was performed based on a GiWi (GSK3 inhibitor and Wnt inhibitor) protocol described previously³². Briefly, ES cells were grown on Matrigel-coated plates in mTeSR1 medium to 80–90% confluence before passaging with Accutase and then re-seeded at 0.5–1.5 million cells per well in 12-well plates in 1 ml of mTeSR1 plus 10 μM Y27632. On day 5, cells were incubated with 6–12 μM CHIR99021 (Selleckchem) for 16–24 h after which the medium was replaced with 2 ml of RPMI/B27 (without insulin) and cultured for two days. On day 8, 2 ml combined medium was prepared by mixing 1 ml of medium collected from the 12-well plate with 1 ml fresh RPMI/B27 (without insulin) medium. Medium on the cells was then replaced with 2 ml combined medium with 5 μM IWP2 (Tocris) and cells were cultured for 48 h. At day 10, 2 ml of fresh RPMI/B27 (without insulin) was added to each well of the 12-well plate. Starting from day 12, medium was changed every three days with 2 ml per well of RPMI/B27 (with insulin). Contracting cardiomyocytes were observed as early as day 17 from the initial passaging of ES cells.

Differentiated cell types were characterized by immunocytochemistry as described previously²².

Teratoma assay. ES cells were injected into the femoral region of 6-week-old, male SCID mice (Charles River). Mice with tumours were euthanized and teratomas were isolated, sectioned and histologically characterized for the presence of representative tissues as described previously²².

Assessment of mitochondrial function. Mitochondrial respiratory chain enzymatic activities (complex I and IV) were measured by spectrophotometry, as previously described^{23,33}. In brief, intact mitochondria were isolated from fibroblasts and treated with and without 3 mM rotenone at 340 nm wavelength for 5 min. Differences of absorbance per minute were obtained, and the specificity of COMI activity was estimated by the percentage of rotenone inhibition.

For live cell oxygen consumption, a XF96 Extracellular Flux Analyzer (Seahorse Biosciences) was used to measure oxygen consumption rates (OCR) and extracellular acidification rates as described previously²². In brief, NPCs were seeded at a density of 30,000 cells per well of a XF96 cell-culture microplate and incubated

for 24 h to allow cells to attach. Prior to the assay, NPCs were equilibrated for 1 h in unbuffered XF-assay medium supplemented with 25 mM glucose, 1 mM sodium pyruvate, 2 mM glutamax, 1 × non-essential amino acids and 1% (v/v) FBS in a non-CO₂ incubator. Sequential compound injections of oligomycin (0.5 μg ml⁻¹), carbonyl cyanide 4-(trifluoromethoxy) phenylhydrazone (FCCP, 1 μM), rotenone (0.5 μM) and antimycin A (1 μM) were used to measure the parameters of mitochondrial respiration. Indices of mitochondrial function were calculated as basal respiration rate (baseline OCR—rotenone or antimycin A OCR), ATP-dependent (basal respiration rate—oligomycin OCR), maximal respiration rate (FCCP OCR—rotenone or antimycin A OCR) and oxidative reserve (maximal respiration rate—basal respiration rate). Each plotted value was the mean of a minimum of four replicate wells, and was normalized to total cell numbers plated. Results were presented as means ± s.e.m. One-way ANOVA was used for three group comparisons and Student's *t*-test was used for pairwise comparisons. *P* < 0.05 was considered significant. The study was randomized, and the investigators were blinded to sample allocations among different groups.

mtDNA sequencing. Mutations and general mtDNA haplotypes were determined in oocytes, blood, skin fibroblasts, urine, ES cells and differentiated tissues by whole mtDNA next generation sequencing using the Illumina MiSeq platform^{22,23}. DNA was extracted from cells or tissues using Genra DNA extraction kit (Qiagen), and from oocytes using Pico Pure DNA Extraction Kit (Life Technologies). mtDNA was amplified by a single PCR reaction as previously described²². mtDNA amplification from individual oocytes was performed using 2 primer sets: 7272F 5'-GGCTCATTTCATTTCTCTAACAG-3', 15712R 5'-TTGGCTTAGTGGGCGAAATA-3' and 15635F 5'-TCCATCCTCATCC TAGCAAT-3', 7401R 5'-GGGGGCATCCATATAGTCAC-3.

Detection of mtDNA variants by Sanger sequencing. mtDNA variants present at over 10% heteroplasmy were corroborated independently by Sanger sequencing. Regions of mtDNA containing germline mutations or SNPs were amplified by PCR with previously reported primer sets using PCR Platinum SuperMix High Fidelity kit (Life Technologies)²³. PCR products were purified, sequenced and analysed by Sequencher v5.0 (GeneCodes).

ARMS-qPCR and copy number measurement. The amplification refractory mutation system quantitative PCR assay (ARM-qPCR) was performed to verify

heteroplasmy at mt8993T>G, mt13513G>A and mt3243A>G as previously described²² and to measure carryover in control spindle transfer embryos and ES cells at 7843A>G, 16519T>C and 16278T>C¹⁹. Measurement of mtDNA copy number was performed as previously described³⁴.

Genetic and cytogenetic analyses. Karyotyping by G-banding was performed on 20 metaphase cells from each ES cell line by the OHSU Cytogenetics Laboratory. The comparative genome hybridization array was performed by IVI Gen. CNVs were identified by SNP genotyping as previously described³⁵ and analysed at the clinical genetic diagnosis laboratory in Cincinnati Children's Hospital Medical Center.

Statistical analysis. No statistical methods were used to predetermine sample size. The experiments were not randomized. The investigators were not blinded to allocation during experiments and outcome assessment except for neural progenitor cell and cardiomyocyte differentiations and measurements of mitochondrial oxygen consumption rates. Results are presented as means ± s.d. or s.e.m. *P* < 0.05 was considered significant. Data were analysed using Pearson non-parametric test for correlation, independent group *t*-test or χ^2 test for pairwise comparison and ANOVA with Bonferroni analysis for multiple comparisons (IBM SPSS).

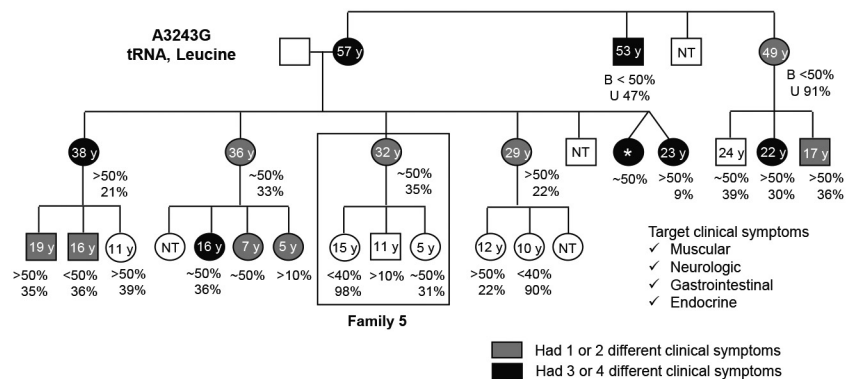
Data availability. All data generated or analysed during this study and included in this letter (and its Supplementary Information) are deposited in the NCBI SRA (SRP091883) and GEO (GSE87897).

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a Information on families with mitochondrial disease

Mitochondrial disease	Family	Affected member	Age (yr)	Clinical manifestation	Mother's age (yr)	mtDNA mutation	Gene
Leigh syndrome	F1	Daughter	2	Seizures, metabolic acidosis, developmental delay, difficulty to thrive, abnormal basal ganglia and enlargement heart ventricle	22	T8993G	<i>ATP6</i> (non-syn)
	F2	Son	2.5	Seizures, chronic acidosis, developmental delay, difficulty to thrive and encephalopathy	23	T8993G	<i>ATP6</i> (non-syn)
	F3	Son	12	Seizures, failure to thrive, difficulty in breathing, enlarged heart ventricle, kidney failure and optic nerve atrophy	36	G13513A	<i>ND5</i> (non-syn)
	F4	Daughter	1	Seizures, difficulty to thrive, lactic acidosis, neuro-motor impairment and history of congenital brain abnormality	28	No	NA
MELAS syndrome	F5	Maternal sister	23	Mitochondrial encephalomyopathy, lactic acidosis and stroke-like episodes	32	A3243G	tRNA (Leucine)

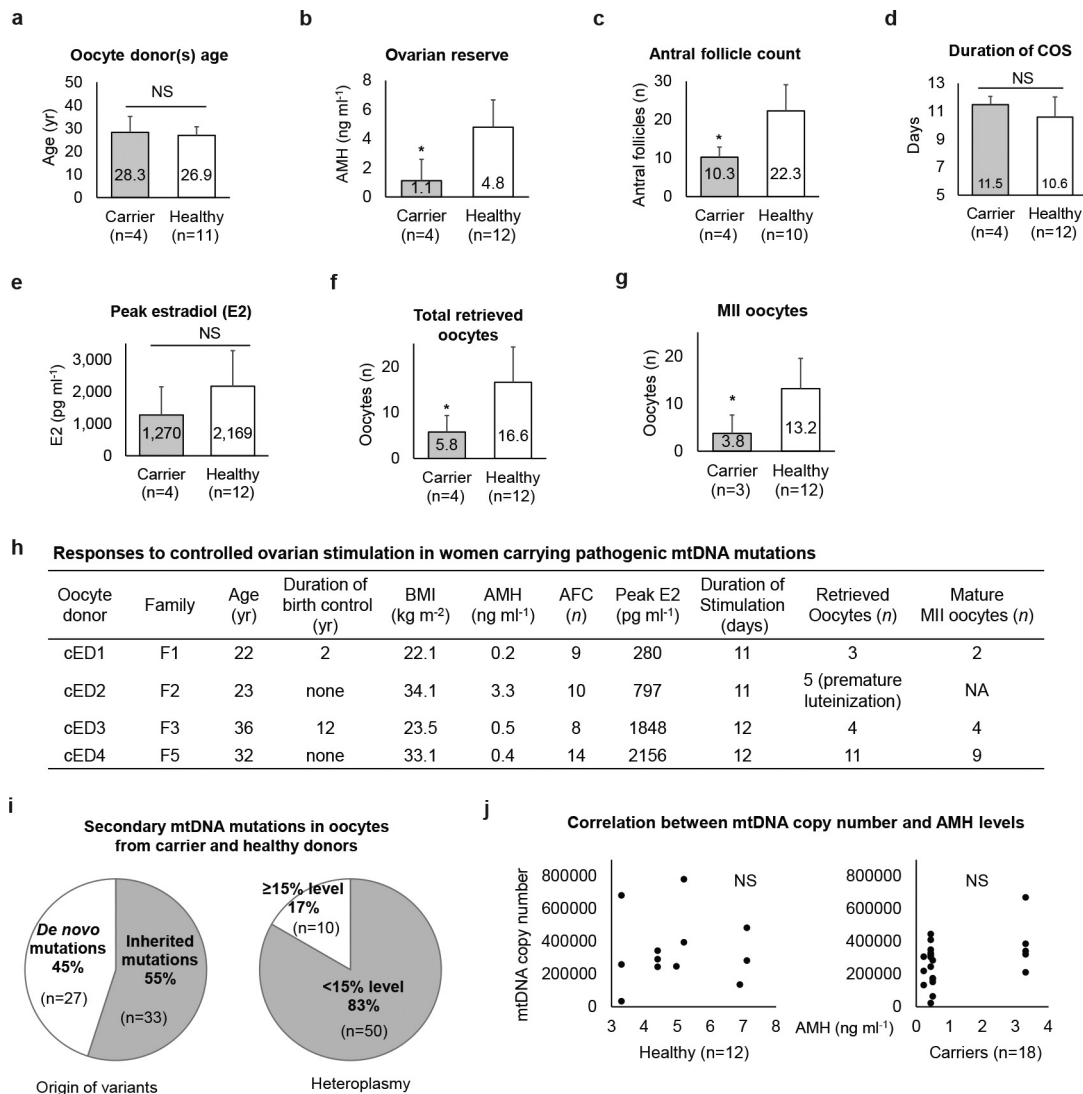
b A3243G mutation distribution in pedigree of family 5 with MELAS syndrome



Extended Data Figure 1 | Families with mitochondrial disease.

a, Families with mitochondrial disease and women of reproductive age were recruited for MRT. All Leigh syndrome families had an existing, severely affected child. F, family; non-syn, non-synonymous; NA, not applicable. **b**, Family 5 was selected from an extensive MELAS syndrome

pedigree. Clinical phenotypes of A3243G MELAS syndrome varied even with similar heteroplasmy levels. Asterisk indicates the first MELAS patient diagnosed clinically. Heteroplasmy in blood (upper panel) and urine (lower panel). Squares, males; circles, females. B, blood; U, urine. NT, not tested. Percentage values refer to heteroplasmy; y, years-old.



Extended Data Figure 2 | Response to controlled ovarian stimulation and oocyte characteristics. **a**, Age of oocyte donor(s) was similar between carrier and healthy groups. **b**, **c**, Ovarian reserve (**b**) and antral follicle count (**c**) were significantly lower in carriers than in healthy oocyte donors. **d**, **e**, Duration of controlled ovarian stimulation (**d**, COS) was longer in carriers and peak oestradiol (**e**, E2) on day of hCG tended to be lower in carriers. **f**, **g**, Total oocyte yield (**f**) and number of mature oocytes (**g**) were significantly lower in carriers than healthy oocyte donors. n = the number of controlled ovarian stimulation cycles. Data are represented as mean \pm s.d. * $P < 0.05$. **h**, Baseline characteristics and cycle outcomes.

Carriers had a lower anti-Müllerian hormone (AMH) levels. Lower peak oestradiol levels were measured in cED1 and cED2. Birth control method: combined oral contraceptive, medroxyprogesterone. **i**, Analysis of heteroplasmic mtDNA variants detected in MII oocytes. *De novo* indicates unique mutations found in individual oocytes; inherited indicates mutations shared with other oocytes, sibling children or egg donors. n = the number of mutations in individual oocytes. **j**, anti-Müllerian hormone levels, a measure of ovarian reserve, were not correlated with mtDNA copy number. n = the number of oocytes. NS, not significant. Source data files are available for **a–g**.

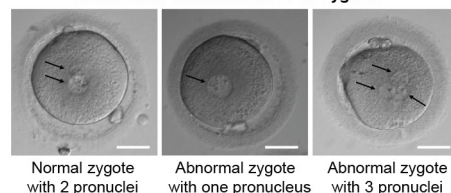
a

Donor oocyte mtDNA haplotypes and SNP differences in ST embryos

Cyto Karyo	Egg donor	hED1 (21-yr)	hED2 (29-yr)	hED3 (23-yr)	hED4 (25-yr)	hED5 (25-yr)	hED6 (28-yr)	hED7 (24-yr)	hED8 (34-yr)	hED9 (32-yr)	hED10 (27-yr)	hED11 (29-yr)
	mtDNA haplotype	L2c	D4a	A2g	B2k	U5a	V3	T2	H1b	H2a	H6a	H56
hED1 (21-yr)	L2c									57		
hED2 (29-yr)	D4a		49									
hED3 (23-yr)	A2g		49									
hED5 (25-yr)	U5a					33			33			
hED6 (28-yr)	V3					33						
hED8 (34-yr)	H1b					33						
hED9 (32-yr)	H2a	57										6
hED11 (29-yr)	H56									6		
cED1 (22-yr)	H1b										21	
cED3 (36-yr)	T2b							22				
cED4 (32-yr)	H49				32							

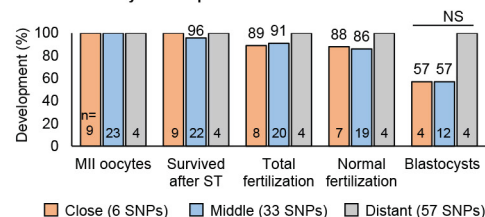
b

Fertilization abnormalities in ST zygotes



e

Embryo development with different donor mtDNA



c Fertilization abnormalities in ST zygotes and embryos

Treatment	Total fertilized zygotes (n)	Total abnormal fertilized zygotes (%)	Abnormal No. of pronuclei			Blastocysts	Karyotypes
			One	Three (%)	Abnormal divided		
Intact control	17	1 (6)	1	0	0	0	
Control ST	32	2 (6)	1	1 (3)	0	1	69,XXY
Carrier ST	12	4 (33)	1	2 (17)	1	0	

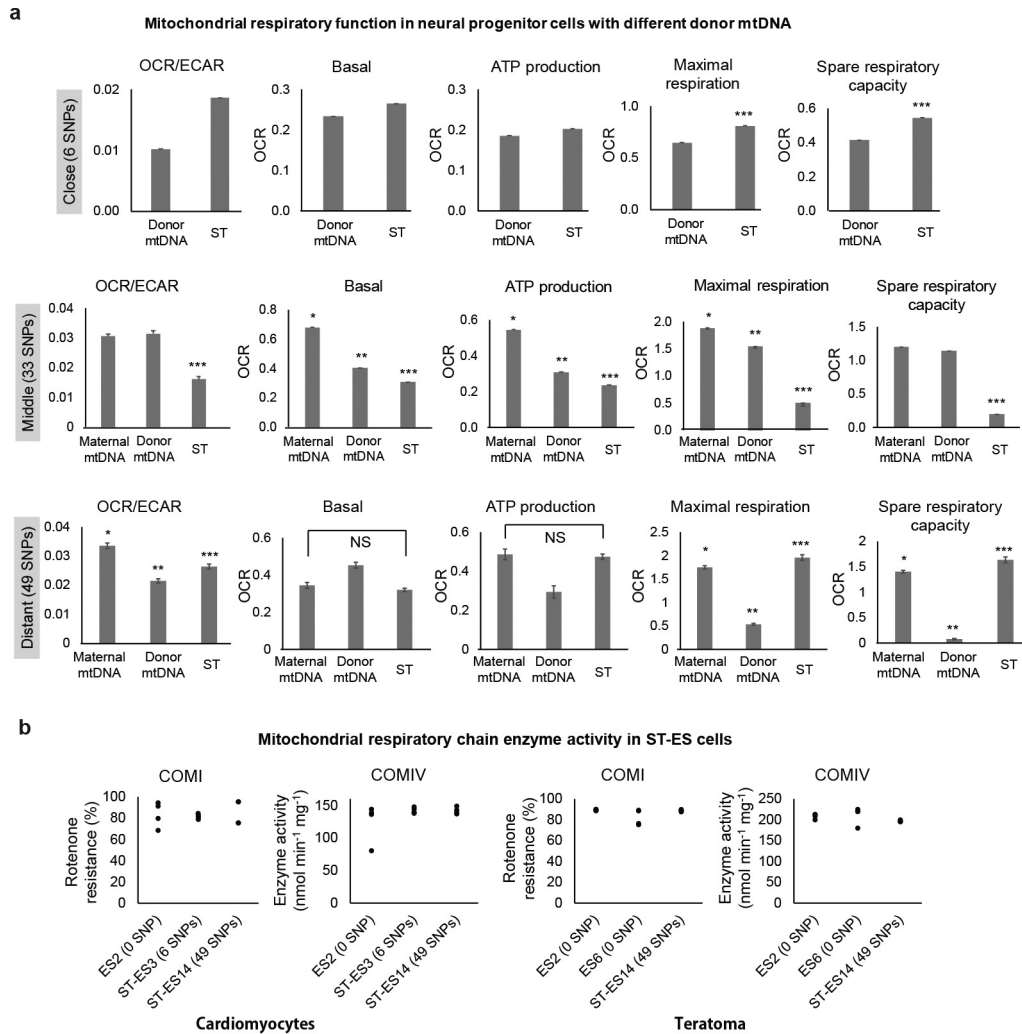
d Fertilization and embryo development of frozen ST oocytes

Treatment	MI I oocytes (n)	Survived after ST (%)	Total fertilization (%)	Normal fertilization (%)	Abnormal No. of pronuclei		Normal morula (%)	Normal blastocysts (%)
					One	Three (%)		
Vitrified cytoplasts/ Fresh spindles	6	6 (100)	6 (100)	4 (67)	1	1 (17)	4 (100)	1 (25)
Vitrified spindles/ Fresh cytoplasts	6	6 (100)	3 (50)	2 (67)	0	1 (33)	2 (100)	1 (50)

Extended Data Figure 3 | Fertilization and development after spindle transfer.

a, Whole mtDNA sequencing was performed to identify haplotypes of all egg donors and spindle transfer was performed to match various haplotypes. Cyto, cytoplasm; Karyo, karyoplast. **b**, Normal fertilization (2 pronuclei) and abnormal fertilization (1 or 3 pronuclei); see arrows. Scale bars, 50 μm. **c**, Carrier spindle transfer showed higher abnormal fertilization than controls. One zygote from the control

spindle transfer group with three pronuclei developed to a blastocyst and displayed a 69,XXY karyotype. **d**, The blastulation rate was similar between vitrified cytoplasm with fresh spindles and vitrified spindles with fresh cytoplasm ($P > 0.05$; t -test). **e**, Control spindle transfer embryo development as a function of donor mtDNA matching distances. The numbers on the top of bars are percentage of embryo development. n = the number of embryos; NS, not significant ($P > 0.05$; t -test).



Extended Data Figure 4 | Mitochondrial function in cells with donor mtDNA. **a**, Energy metabolism ($\frac{OCR}{ECAR}$) was measured and compared between neural progenitor cells (NPCs) derived from MRT and control embryos carrying either donor or maternal mtDNA. OCR, oxygen consumption rate (representing oxidative phosphorylation); ECAR, extracellular acidification rate (representing glycolysis). NPCs from MRT ES cells with 6 and 49 SNP differences displayed comparable oxygen consumption rates to NPCs harbouring maternal or donor mtDNA except maximal respiration in the 49 SNP group. The oxygen consumption rate was reduced in NPCs with 33 SNPs. The oxygen consumption rate data

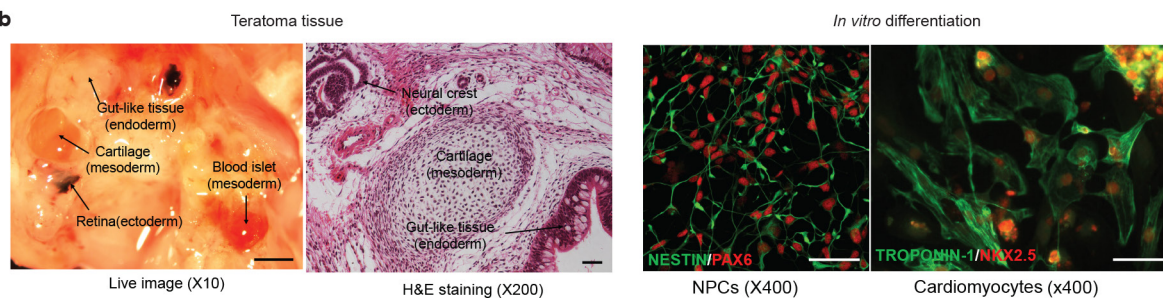
were normalized by live cell DNA content. * $P < 0.05$ versus host mtDNA, ** $P < 0.05$ versus donor mtDNA, *** $P < 0.05$ versus spindle transfer (t -test for 6 SNPs and one-way ANOVA for 33 and 49 SNPs, $n = 16$ technical replicates). Data are shown as mean \pm s.e.m. NS, not significant. **b**, Mitochondrial respiratory chain enzyme activity in differentiated cells from spindle transfer ES cell lines. Mitochondrial complex I and IV activities in differentiated cells from spindle transfer ES cell lines carrying donor mtDNA with 49 SNP differences were comparable to controls with either donor or maternal mtDNA ($P > 0.05$). One-way ANOVA, $n = 4$ technical replicates. Source data file is available for **b**.

a

***In vivo* and *in vitro* differentiation of ST-ES cells**

No.	Cell line	Maternal mtDNA haplotype	Donor mtDNA haplotype	SNP difference	Teratoma	NPCs	Cardiomyocytes
1	ES1	----- H1b control -----	H1b control	0	NT	yes	yes
2	ES2	----- D4a control -----	D4a control	0	yes	yes	yes
3	ES3	----- H56 control -----	H56 control	0	NT	yes	yes
4	ES6	----- A2g control -----	A2g control	0	yes	yes	yes
5	ST-ES3	H2a	H56	6	NT	yes	yes
6	ST-ES6	H1b	U5a	33	NT	yes	yes
7	ST-ES7	U5a	H1b/ U5a (reversed)	33	NT	yes	yes
8	ST-ES14	D4a	A2g	49	yes	yes	yes
9	ST3243-ES1	H49	B2k/ H49 (reversed)	31	yes	yes	yes
10	ST3243-ES2	H49	B2k	31	yes	yes	yes
11	NT-ES8	X2c	D4a/ X2c (reversed)	39	yes	yes	yes

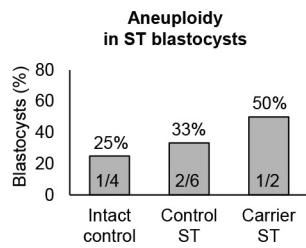
b



Extended Data Figure 5 | Differentiation of ES cells derived from MRT embryos. **a**, Summary of *in vitro* and *in vivo* differentiation of ES cells derived from spindle transfer or SCNT embryos carrying donor mtDNA. Similar to control IVF ES cells, all tested MRT ES cells produced teratoma tumours *in vivo* and formed neural progenitor cells (NPCs)

and cardiomyocytes *in vitro*. NT, not tested. **b**, Histological analysis of differentiated tissues from MRT ES cells. The representative tissues were collected and used for mtDNA carry-over analysis and measurements of mitochondrial function. Scale bars, 1 mm (live image) and 50 μ m (haematoxylin and eosin (H&E) and immunofluorescence staining).

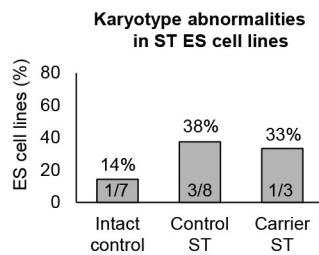
a



Summary of array CGH in intact and ST blastocysts

Treatment	#	Age of nuclear donor (yr)	Egg donor ID	aCGH Results	Sex
Intact control	1	34	hED8	Abnormal: -21	Male
	2	29	hED11	Normal	Male
	3	25	hED4	Normal	Female
	4	25	hED4	Normal	Male
Control ST	1	34	hED8	Abnormal: +7, +15	Male
	2	32	hED9	Normal	Male
	3	32	hED9	Normal	Female
	4	32	hED9	Normal	Male
	5	29	hED11	Abnormal: +12	Male
	6	21	hED1	Normal	Male
Carrier ST	1	36	cED3	Abnormal: +9	Male
	2	32	cED4	Normal	Female

b

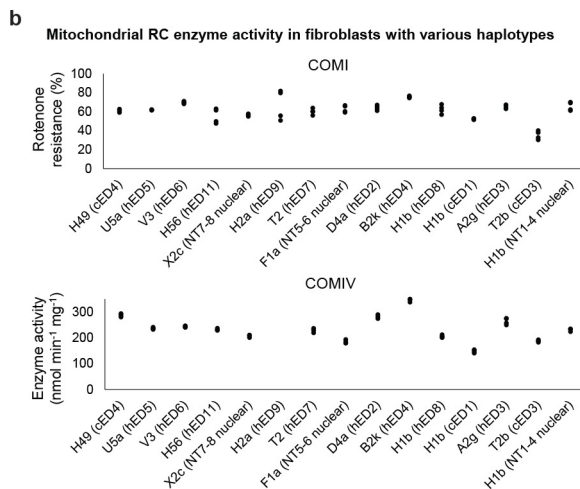
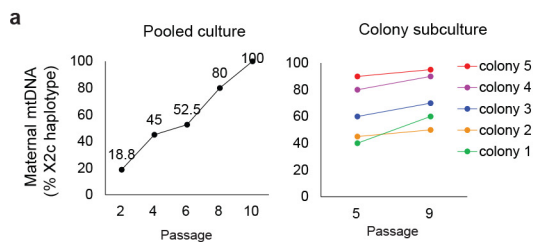


G-banding analysis of ES cell lines derived from intact and ST blastocysts

Treatment	#	Age of nuclear donor (yr)	Egg donor ID	ES cell line	Karyotype
Intact control	1	34	hED8	ES1	46,XX
	2	29	hED2	ES2	46,XX
	3	29	hED11	ES3	46,XY
	4	25	hED4	ES4	46,XY
	5	25	hED5	ES5	46,XX
	6	23	hED3	ES6	46,XX
	7	23	hED3	ES7	45,XY,-21
Control ST	1	34	hED8	ST-ES5	Mosaicism 47,XX +10 [12]/46,XX [8]
	2	34	hED8	ST-ES6	46,XX
	3	32	hED9	ST-ES3	46,XX
	4	29	hED2	ST-ES14	46,XY
	5	28	hED6	ST-ES10	45,X
	6	25	hED5	ST-ES7	46,XY
	7	25	hED5	ST-ES8	92,XXXX
	8	25	hED5	ST-ES9	46,XY
Carrier ST	1	36	cED3	13513 ST-ES	47,XY,+9
	2	32	cED4	3243 ST-ES1	46,XX
	3	32	cED4	3243 ST-ES2	46,XY

Extended Data Figure 6 | Nuclear genome abnormalities associated with spindle transfer. **a**, Aneuploidy rate in blastocysts determined by a comparative genome hybridization array was not significantly different in spindle transfer groups compared to controls. **b**, Karyotype abnormality

rate in ES cells determined by G-banding analysis was also comparable among spindle transfer groups ($P > 0.05$). Normal, 46 XX or 46 XY. Number inside bars, the number of blastocysts or ES cell lines (abnormal/total). Simple χ^2 tests.



Extended Data Figure 7 | Maternal mtDNA heteroplasmy changes and mitochondrial function. **a**, Maternal mtDNA (X2c haplotype) heteroplasmy increased during extended *in vitro* culture and reached homoplasmy in pooled cultures or in individual colony subcultures ($n = 1$). See also Supplementary Table 4. **b**, Respiratory chain complex 1

(COM I) and complex 4 (COM IV) enzyme activities were measured in fibroblasts carrying various human mtDNA haplotypes used in this study. No significant differences were noted ($P > 0.05$). One-way ANOVA, $n = 4$ technical replicates. Source data file is available for **b**.

Extended Data Table 1 | *De novo* copy number variations (CNVs) in ES cells derived from control and spindle transfer blastocysts

Treatment	#	ES cell line ID	CNVs	Chromosome Location	CNV chromosome	Interpretation
Intact control	1	ES1	Deletion (neutral)	15q11.2	male	699 kb region of chromosome 15 (15q11.2) shows low level mosaicism (approximately 10%-20%) for homozygosity of uncertain clinical significance.
	2	ES5	Deletion (neutral)	15q11.2	male	1.1 Mb region of chromosome 15 (15q11.2) shows mosaicism for homozygosity of uncertain clinical significance.
			Deletion (neutral)	Xp22.33-q28	male	Less than 10% of cells of the entire chromosome X (Xp22.33-Xq28) of uncertain clinical significance.
Control ST	1	ST-ES5	Duplication	10p15.3-q26.3	male	Mosaic trisomy 10.
			Deletion (neutral)	15q11.2	male	549 kb region of chromosome 15 (15q11.2) shows mosaicism for homozygosity of uncertain clinical significance.
			Deletion	Xp22.33-q28	male	Monosomy X (Xp22.33-Xq28) of uncertain clinical significance.
	2	ST-ES6	None	NA	NA	NA
	3	ST-ES7	None	NA	NA	NA
4	ST-ES9	None	NA	NA	NA	
Carrier ST	1	3243ST-ES1	None	NA	NA	NA
	2	3243ST-ES2	Deletion (neutral)	15q11.2	male	791 kb region of chromosome 15 (15q11.2) shows mosaicism for homozygosity of uncertain clinical significance.

Normal, diploid chromosomal arrangements are identical to original male and female parents. Grey box, shared CNVs among independent cell lines; NA, not applicable.

Extended Data Table 2 | MRT outcomes for families with pathogenic mtDNA mutations

	Family 1	Family 2	Family 3		Family 4		Family 5		
Mitochondrial disease	Leigh syndrome	Leigh syndrome	Leigh syndrome	Leigh syndrome	Leigh syndrome	Leigh syndrome	MELAS		
Pathogenic mtDNA mutation	T8993G	T8993G	G13513A		No		A3243G		
Age of carrier (yr)	22	23	36		28		32		
COS	Yes	Yes	Yes		Excluded		Yes		
No. retrieved oocytes	3	5	4		NA		11		
No. ST oocytes	2	Canceled	4		NA		7		
ST blastocysts	0	NA	2		NA		4		
Grade at D6	NA	NA	5AA	5BB	NA	5AA	5AA	5BB	5CC
Aneuploidy	NA	NA	Yes:+9	NT	NA	NT	No	No	No
Sex	NA	NA	M	NA	NA	NA	M	F	F
mtDNA carryover	NA	NA	< 1%	NA	NA	NA	< 1%	< 1%	< 1%*

Six spindle transfer embryos derived from carrier oocytes reached the blastocyst stage. At least two blastocysts were eligible for transfer. M, male; F, female. NA, not applicable; NT, not tested; D6, blastocysts at day 6 of development. The first number in grade indicates expansion, grade 5 means blastocyst is hatched. The second letter indicates quality of the inner cell mass (ICM), A refers to large ICM of tightly packed cells; B, smaller ICM, loose cells; C, very small ICM. The third letter grades the trophectoderm quality.

*mtDNA in ES cells from this blastocyst reversed to the original maternal haplotype but did not carry the A3243G mutation.