Resource

**Cell Reports**

**Pressure-Driven Mitochondrial Transfer Pipeline Generates Mammalian Cells of Desired Genetic Combinations and Fates**

**Graphical Abstract**

**Highlights**

- We report a “proof-of-principle” mitochondrial transfer pipeline by MitoPunch

- MitoPunch generates cells with unique mtDNA-nDNA pairs, regardless of cell source

- Replacement mtDNA in non-immortal cells remains stable with cell fate conversions

- Enables studies of mtDNA-nDNA interactions with reprogramming and differentiation

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**In Brief**

Patananan and colleagues demonstrate a pipeline for transferring isolated mitochondria into mtDNA-deficient recipient cells. mtDNA-depleted fibroblasts permanently retain acquired non-native mtDNA through cell fate transitions. Initially, mitochondrial recipients show mtDNA-deficient cell transcriptome and metabolome profiles, with improvement to control profiles by reprogramming to pluripotency and subsequent differentiation.

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Generating mammalian cells with desired mitochondrial DNA (mtDNA) sequences is enabling for studies of mitochondria, disease modeling, and potential regenerative therapies. MitoPunch, a high-throughput mitochondrial transfer device, produces cells with specific mtDNA-nuclear DNA (nDNA) combinations by transferring isolated mitochondria from mouse or human cells into primary or immortal mtDNA-deficient (ρ0) cells. Stable isolated mitochondrial recipient (SIMR) cells isolated in restrictive media permanently retain donor mtDNA and reacquire respiration. However, SIMR fibroblasts maintain a ρ0-like cell metabolome and transcriptome despite growth in restrictive media. We reprogrammed non-immortal SIMR fibroblasts into induced pluripotent stem cells (iPSCs) with subsequent differentiation into diverse functional cell types, including mesenchymal stem cells (MSCs), adipocytes, osteoblasts, and chondrocytes. Remarkably, after reprogramming and differentiation, SIMR fibroblasts molecularly and phenotypically resemble unmanipulated control fibroblasts carried through the same protocol. Thus, our MitoPunch “pipeline” enables the production of SIMR cells with unique mtDNA-nDNA combinations for additional studies and applications in multiple cell types.
respiration. As many as 1:5,000 people have mtDNA mutations that impair high-energy-demand tissues and contribute to debilitating diseases, including cancer, diabetes, and metabolic syndromes (Schafer et al., 2004). In addition, cells may contain a mixture of different mtDNA sequences, a situation termed heteroplasm, with up to 1 in 8 asymptomatic individuals carrying an unsuspected pathogenic mtDNA mutation (Elliott et al., 2008; Rebolledo-Jaramillo et al., 2014). Thus, an ability to controllably manipulate mtDNA sequences could enable studies of mitochondria and potentially develop disease models or therapies for mtDNA disorders.

In human reproduction, several types of mitochondrial replacement strategies were developed to exchange pathogenic mtDNA in a zygote with non-detrital mtDNA from a healthy donor oocyte. These approaches have potential for preventing transmission of mtDNA disorders from mother to their children (Wolf et al., 2015; Wolf et al., 2019). However, in vitro methods to change mtDNA sequences within somatic cells and tissues remain limited (Patananan et al., 2016). Cell fusions that produce “cybrids” permanently retain donor mitochondria (Wong et al., 2017), although fusion partners are typically transformed cells that cannot be reprogrammed. Also, endonucleases imported into mitochondria can shift heteroplasm ratios to alter mitochondria and cell functions by targeting specific sequences for destruction. However, these endonucleases are laborious to produce, are limited to certain mtDNA sequences, and do not yield homoplasmy (Campbell et al., 2016; Kitani et al., 2015). Thus, an ability to controllably manipulate mtDNA sequences could enable studies of mitochondria and potentially develop disease models or therapies for mtDNA disorders.

Several methods transfer isolated mitochondria into mtDNA-deficient cells, known as ρ0 (rho null) cells, to restore respiration (Kim et al., 2018; Nigou Mombo et al., 2017). In addition, some studies reported endocytosis of mitochondria by mammalian cells (Clark and Shay, 1982; Kesner et al., 2016). However, these studies were not concerned with theft mtDNA from cells and generating stable isolated mitochondrial recipient (SIMR) clones that permanently retain donor mtDNA (Kesner et al., 2016; Kitani et al., 2014; Sun et al., 2019). A recent study did produce a limited number of SIMR clones by coincubating high concentrations of isolated HEK293T donor mitochondria with ρ0 osteosarcoma cells (Patel et al., 2017). We (Dawson et al., 2020) and others (Ali Pour et al., 2020) have recently reported similar findings in which cells are capable of endocytosing exogenous mitochondria and even altering metabolic functions for a limited period of time (~1 week), but these exogenous mtDNAs are lost over time. To address this problem, we previously developed a photothermal nanoblade to stably transfer small quantities of isolated mitochondria into ρ0 osteosarcoma cells (Wu et al., 2016). Unfortunately, the nanoblade is laborious and low throughput, and two of three SIMR clones reported did not reset the ρ0 cell metabolome. A technique that generates many non-transformed stable clones is desirable to examine novel mtDNA-nuclear DNA (nDNA) combinations through reprogramming to pluripotency and differentiation into multiple cell types.

Here, we describe a simple-to-use mitochondrial transfer technique called “MitoPunch” to rapidly generate numerous non-transformed SIMR clones. We apply MitoPunch to implement a pipeline that demonstrates donor mtDNA functions in recipient host primary cells at different cell fates. Our study establishes this resource pipeline to generate primary SIMR cells using non-immortalized materials. We also measure the status of the metabolome, transcriptome, and biophysical properties of SIMR cells with defined mtDNA-nDNA combinations to guide future studies generating somatic cells with desired mtDNA-nDNA combinations.

RESULTS

MitoPunch Generates SIMR Cells with a Range of Cell Types and mtDNAs

MitoPunch is a massively parallel, pressure-driven, large cargo transfer platform based on prior photothermal nanoblade and biophotonic laser-assisted cell surgery tool (BLAST) technologies (Sercel et al., 2020; Wu et al., 2016; Wu et al., 2015). Mito-Punch uses a mechanical plunger to physically deform a pliable polydimethylsiloxane (PDMS) reservoir containing isolated mitochondria suspended in phosphate-buffered saline (1× PBS [pH 7.4]) (Figure 1A). Plunger activation propels the suspended cargo within the PDMS delivery chamber through a porous membrane containing numerous 3-μm-diameter holes on which a confluent layer of adherent cells is grown. This action directly forces isolated mitochondria into the cytosol of recipient cells.

To demonstrate MitoPunch generation of SIMR cells, we transferred isolated mitochondria from ~1.5 × 10^5 HEK293T cells into ~2 × 10^5 143BTK–ρ0 osteosarcoma cells. Post-transfer, we select for and isolate SIMR colony clones with permanently retained donor mtDNA using uridine-deficient media. This selection is enabling because respiration-defective ρ0 cells have inactive dihydroorotate dehydrogenase and depend on exogenous uridine or restored respiration for pyrimidine biosynthesis (Grégoire et al., 1984). Compared to the coinubation of the same amount of isolated mitochondria with cells (Clark and Shay, 1982; Kesner et al., 2016), only 143BTK–ρ0 cells with HEK293T mitochondria from MitoPunch transfer (143BTK–ρ0+HEK293T) permanently retained donor mtDNA and survived uridine-deficient media selection (Figure 1B). In a representative set of mitochondrial transfer experiments, MitoPunch generated ~75 independent crystal-violet-stained SIMR clones in comparison to no clones obtained by coinubation (Figure 1B).

We next examined whether MitoPunch could generate SIMR clones with defined mtDNA-nDNA pairs that transfer features of mitochondrial disease. We isolated mitochondria from cybrid cells containing either an A3243G mtDNA substitution commonly associated with mitochondrial encephalopathy, lactic acidosis, and stroke-like episodes (MELAS) or wild-type (WT), non-mutant mtDNA from the same individual (Picard et al., 2014). The A3243G point mutation is in the tRNA^Leu^ gene and results in altered production and assembly of ETC complexes with impaired oxidative phosphorylation (Chomyn et al., 1992; Sasarman et al., 2008). Following MitoPunch into 143BTK–ρ0 recipients and 2 weeks of selection, two of several dozen independent SIMR clones that permanently retained MELAS (143BTK–ρ0+MELAS) or WT (143BTK–ρ0+WT) mtDNA were tested for oxygen consumption rate (OCR) using the Seahorse.
Figure 1. MitoPunch Is a Versatile Mitochondrial Transfer Technology
(A) Schematic representation of the MitoPunch mitochondrial transfer platform.
(B) Images of crystal-violet-stained SIMR colonies from coincubation or MitoPunch delivery of either 1× PBS (pH 7.4) (sham control) or isolated HEK293T cell mitochondria into 143BTK−p0 osteosarcoma cells after selection in uridine-depleted medium. Data are the means ± SD of three technical replicates.
(C) OCR measurements for ~1.5 × 10^6 143BTK−p0 SIMR, 143BTK−p0+MELAS, WT cybrid, MELAS cybrid, 143BTK−p0+WT SIMR, and 143BTK−p0+WT SIMR cells by Seahorse XF96 Extracellular Flux Analyzer. Values were calculated by standard procedures (see STAR Methods). Data are the means ± SD of four technical replicates.

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Extracellular Flux Analyzer. Results showed 143BTK–p0+MELAS clones had significantly impaired basal respiration, maximal respiration, and spare respiratory capacity compared to patient-matched 143BTK–p0+WT and native 143BTK–control cells (Figure 1C), indicating stable mtDNA transfer of the primary metabolic deficit of the MELAS phenotype.

We then expanded mitochondrial donor and recipient cell pairings beyond these initial studies to demonstrate the versatility of MitoPunch. As examples, mitochondria were isolated from harvested C57BL/6 mouse tissues and MitoPunch transferred into C3H/An-derived L929 p0 immortalized fibroblasts. Two weeks of selection yielded dozens of SIMR clones from each mitochondrial source. SIMR clones generated with high-energy-demand heart, lung, or muscle-derived mitochondria showed the most robust respiratory profiles, in contrast to SIMR clones that received low-energy-demand spleen- or kidney-derived mitochondria (Figure 1D). We also evaluated MitoPunch delivery of heteroplasmic mtDNA mixtures into cells. Mitochondria isolated from mouse cybrid lines containing mtDNA mutations in the cytochrome B (mt-Cytb), NADH dehydrogenase subunit 4 (mt-nd4), and NADH dehydrogenase subunit 6 (mt-nad6) genes were MitoPunch transferred individually or in 1:1 mixtures by protein content into L929 p0 fibroblasts. SIMR cells with a single source of mutant mtDNA continued to show severe respiratory impairments (Figure 1E). In contrast, SIMR cells in a mixture of non-overlapping mutant mtDNAs showed markedly improved respiratory profiles, strongly suggesting that both mtDNAs were stably maintained (Figure 1E). Thus, MitoPunch and selection is a versatile approach for generating human or mouse SIMR cells with desired mtDNA-nDNA pairs. Co-transfer of multiple mtDNA types into the same recipient cell also provides a simple method to examine complementation for mutant mtDNA mixtures.

**MitoPunch Generates Non-Transformed, Non-Malignant SIMR Cells**

To obtain SIMR cells with mtDNA-nDNA combinations using non-immortalized recipient cells, we established a human fibroblast mitochondrial recipient pipeline. Hayflick-limited BJ fibroblasts, neonatal dermal fibroblasts (NDFs), and adult dermal fibroblasts (ADFs) were treated for 3 weeks with FDA-approved 2′,3′-dideoxycytidine (ddC) (Nelson et al., 1997) to deplete endogenous mtDNA. Primary p0 human fibroblasts had undetectable mtDNA (Figures S1A and S1B) and cellular respiration (Figures S1C and S1D) by qPCR and Seahorse assay, respectively. Because ddC could cause nDNA alterations, we examined p0 fibroblasts by whole-genome sequencing and identified only a few non-synonymous mutations at 0.6 mutations per megabase, on average, with no chromosomal breaks and no changes in DNA copy number (Table S1).

Subsequently, mitochondria isolated from a human peripheral blood mononuclear cell lot (PBMC1) were transferred into fresh p0 fibroblasts, followed by an empirical and reproducible selection protocol with uridine-deficient galactose medium. From 5–10 BJ fibroblasts, NDFs, or ADFs, p0+PBMC1 SIMR clones were isolated that showed the correct mtDNA-nDNA sequence pairs and human leukocyte antigen (HLA) recipient cell haplotypes (Figures 2A–2C). Primary, non-immortal SIMR clones were also obtained from independent PBMC2 and HEK293T cell mitochondrial transfers. We observed variable efficiencies for HEK293T cell and the PBMC2 mitochondrial transfers, whereas ADFs p0+PBMC2 did not yield clones (Figure S1E).

Analysis of the bulk culture representing 23 BJ p0+HEK293T SIMR clones confirmed the correct mtDNA-nDNA pairing and HLA haplotype (Figures S1F and S1G).

We examined the respiratory function of BJ p0+PBMC1 and BJ p0+HEK293T SIMR fibroblasts by Seahorse assay, which showed statistically improved basal and maximal respiration and spare respiratory capacity for both SIMR cell types compared to BJ p0 fibroblasts, albeit remaining lower than levels for control BJ fibroblasts (Figures 2D and S1H). Immunofluorescence (IF) microscopy showed BJ p0 fibroblasts with a fragmented mitochondrial network morphology lacking mtDNA-containing nucleoids, as observed previously for p0 cells (Kukat et al., 2008) (Figure 2E). In contrast, native BJ fibroblasts showed a reticular mitochondrial network with dozens of nucleoids per cell (Figure 2E). By IF nucleoid speckle numbers, both BJ p0+PBMC1 and BJ p0+HEK293T SIMR cells appeared to restore mtDNA content to levels equivalent to or exceeding that of native BJ fibroblasts (Figures 2E and S1I). SIMR cell mitochondria showed a reticular mitochondrial network morphology similar to that of native BJ fibroblasts, although with denser and more swollen mitochondria (Figures 2E and S1I). Despite SIMR fibroblasts permanently retaining donor mtDNA, OCR and IF suggest that assimilation of transferred mtDNA results in cells with features in between those of BJ p0 and native BJ fibroblasts.

**SIMR Fibroblasts Are Reprogrammable**

We reprogrammed BJ p0+PBMC1 and BJ p0+HEK293T SIMR fibroblasts along with native BJ fibroblasts using OCT4, SOX2, KLF4, cMYC, NANOG, and LIN28 RNAs and quantified for TRA-1-60″ staining clones. In two independent experiments, native BJ fibroblasts yielded an average of 136 reprogrammed TRA-1-60″ clones (0.068% efficiency), compared to 21 (0.011%) and three (0.0015%) clones for BJ p0+PBMC1 and BJ p0+HEK293T cells, respectively (Figures 3A and S1J). Three unique reprogrammed clones of BJ p0+PBMC1-iPSCs (1, 2, and 11) and BJ p0+HEK293T-iPSCs (1, 2, and 4) were tested for pluripotency biomarkers and stained positive for OCT3/4 and SOX2 transcription factors by flow cytometry, as did BJ-induced pluripotency cell iPSC control, but not native BJ fibroblasts, as expected (Figures 3B and S1K). Conversely, the differentiated cell biomarker CD44 (Quintanilla et al., 2014) was negative in all reprogrammed BJ p0+PBMC1-iPSC, BJ p0+HEK293T-iPSC, and control BJ-iPSC clones and immunostained only the native BJ fibroblasts.
BJ-iPSC and all SIMR-iPSC reprogrammed clones were also SSEA-4+ (Abujarour et al., 2013) and OCT4+ by IF (Figures 3C and S1L). Seahorse assays of BJ p0+PBMC1-iPSC and BJ p0+HEK293T-iPSC clones showed minimal or no statistical differences in basal respiration, maximal respiration, and spare respiratory capacity compared to the native BJ-iPSC control (Figures 3D and S1M). Thus, SIMR fibroblast reprogramming generated iPSCs with donor mtDNA.

Other studies have shown iPSC reprogramming of fibroblasts from individuals with pathogenic mtDNA mutations (Cherry et al., 2013; Folmes et al., 2013; Hämäläinen et al., 2013; Kang et al., 2016; Kodaira et al., 2015; Ma et al., 2015; Matsubara et al., 2018; Pek et al., 2019; Perales-Clemente et al., 2016; Russell et al., 2018; Yang et al., 2018), but this has not been attempted for SIMR fibroblasts with donated, non-native mutant mtDNA. Therefore, mitochondria containing an A3243G MELAS mtDNA mutation or WT mtDNA were isolated, followed by MitoPunch transfer into BJ p0 fibroblasts and selection. BJ p0+MELAS SIMR fibroblasts showed impaired proliferation during reprogramming and did not yield iPSCs (data not shown). Therefore, we switched to NDF p0 recipient fibroblasts and generated SIMR fibroblasts using isolated MELAS (NDF p0+MELAS), WT (NDF p0+WT), or NDF (NDF p0+NDF) mitochondria. Seahorse assays showed that NDF p0+MELAS fibroblasts had a significant reduction in basal respiration, maximal respiration, and spare respiratory capacity compared to native NDF, NDF p0+NDF, and NDF p0+WT.
fibroblasts (Figure S1N). Restriction fragment length polymorphism (RFLP) PCR analyses confirmed the generation of homoplasmic NDF\(r\)0+MELAS fibroblasts (Figure S1O). We also generated \(\sim\)25–50% heteroplasmic NDF\(r\)0+MELAS/WT fibroblasts, which was verified by RFLP analyses (Figure S1O). Homoplasmic NDF\(r\)0+MELAS fibroblasts underwent RNA-based reprogramming as described earlier, but all developing iPSC clones spontaneously differentiated (Figure S1P). Reprogramming of NDF\(r\)0+MELAS/WT heteroplasmic fibroblasts (Figures S1O) yielded 20 iPSC clones, but all clones retained only WT mtDNA by RFLP analysis (Figures S1P and S1Q). To examine whether the reprogramming method influenced mutant mtDNA SIMR-iPSC generation, NDF\(r\)0+MELAS fibroblasts underwent integrating DNA, lentiviral, and Sendai virus reprogramming strategies. In all cases, NDF\(r\)0+MELAS cells spontaneously differentiated despite early signs of reprogramming (Table S2). In addition, no NDF\(r\)0+MELAS-iPSCs were obtained when reprogramming was performed with additional uridine supplementation, antioxidant N-acetylcysteine, a Rho-associated protein kinase (ROCK) inhibitor, or low oxygen tension (data not shown). Similar results were also obtained with all four reprogramming strategies for NDF\(r\)0 SIMR fibroblasts containing additional mtDNA mutations including a cytochrome B deletion, a Kearns-Sayre common deletion, and A8344G or T8993G mtDNA substitutions (Table S2). Thus, SIMR fibroblasts readily maintain a large variety of mtDNA sequences, in contrast to SIMR-iPSCs, which can be generated only with non-detrimental mtDNA sequences. Further biochemical investigations are needed to determine how mtDNA sequences dictate SIMR reprogramming, whereas native mutant mtDNA fibroblasts can be reprogrammed (Hämäläinen et al., 2013; Ma et al., 2015; Pek et al., 2019).

SIMR-iPSCs Produce Functional, Differentiated Cell Types

We next determined whether SIMR-iPSCs with isogenic nuclei and non-native donor mtDNAs could differentiate. We chose to examine defined medium differentiation of mesenchymal stem cells (MSCs) because of their relevance to potential therapies and current use in over 850 clinical trials (Hsu et al., 2016). A BJ-iPSC control, BJ\(r\)0+PBMC1-iPSCs, and BJ\(r\)0+HEK293T-iPSCs were differentiated into MSCs and validated with an antibody panel against surface biomarkers established by the International Society for Cellular Therapy (ISCT) (Dominici et al., 2006). Flow cytometry verified that the BJ-MSC control, BJ
SIMR Cell Metabolism and RNA Transcript Changes with Fate Transitions

We used ultra-high-performance liquid chromatography-mass spectrometry (UPLC-MS) to quantify 154 steady-state metabolites in native BJ, BJ p0, BJ p0+PBMC1, and BJ p0+HEK293T clones, and BJ p0+HEK293T clones at fibroblast, iPSC, and MSC fates. Hierarchical clustering showed distinct, grouped profiles for fibroblasts, iPSCs, and MSCs independent of mitochondrial transfer status (Figures S2A and S2B; Table S3). Principal component analysis (PCA) of metabolite data also showed three main clusters representing fibroblast, iPSC, and MSC fates but no clear differences between SIMR and native control cells within each fate (Figures S2C and S2D). Metabolite set variation analysis (MSVA) and Euclidean distance analysis of the BJ p0+PBMC1-iPSC and BJ p0+PBMC1-MSC clones showed similar metabolite pathway profiles to themselves and to the respective BJ-iPSC and BJ-MSC controls (Figures S2A, S2E, and S2F). In contrast, BJ p0+HEK293T-iPSC clones 1 and 2 clustered separately from clone 4 and the BJ-iPSC control for several metabolic pathways, particularly purine, pyrimidine, glutathione, and ethanol metabolism (Figures S2B, S2E, and S2F). This separation in BJ p0+HEK293T-iPSC clones was no longer present upon further differentiation to BJ p0+HEK293T-MSC clones (Figures S2B, S2E, and S2F). In summary, steady-state metabolite analyses showed that SIMR cells are comparable to native control cells, with only a few differences that are resolved upon iPSC reprogramming and differentiation to MSCs.

We utilized RNA sequencing (RNA-seq) to evaluate whole-transcriptome profiles for SIMR cells at fibroblast, iPSC, and MSC fates. DESeq2 was used to identify significantly differentially expressed genes (DEGs), defined as genes showing an absolute log 2-fold change > 0.5 and adjusted p < 0.05. For both BJ p0+PBMC1 and BJ p0+HEK293T SIMR cells, the greatest number of DEGs compared to native BJ control cells with an adjusted p < 0.05 occurred at the fibroblast fate (Figures 5A and S3A). RNA-seq identified 1741, 194, and 224 elevated and 1827, 68, and 115 repressed DEGs by comparing BJ p0+PBMC1 cells to native BJ parent cells at the fibroblast, iPSC, and MSC fates, respectively (Figure 5A; Table S4). Transcriptomic analysis of the independently generated BJ p0+HEK293T cells similarly identified 1,377, 537, and 239 elevated and 1,564, 648, and 210 repressed DEGs compared to native BJ parent cells at fibroblast, iPSC, and MSC fates, respectively (Figure S3A; Table S4).

Reactome pathway enrichment analysis of DEGs showed diverse pathways altered in SIMR fibroblast transcript profiles compared to those in native BJ fibroblasts, including those associated with extracellular matrix organization and the complement cascade (Figure S3B; Table S5). Differential expression and pathway enrichment analyses comparing all SIMR-iPSCs and SIMR-MSCs to native BJ-iPSC and BJ-MSC controls, respectively, identified a dramatically smaller number of DEGs, with overrepresented pathways driven primarily by a cluster of histone transcripts (Figures S3C and S3D; Table S5).

Further detailed transcriptome analyses uncovered metabolic pathway differences based on cell condition and fate. Somatic cell reprogramming to iPSCs requires a metabolic shift from predominantly oxidative phosphorylation to mainly glycolysis, which correlates with all BJ p0+PBMC1-iPSC and BJ p0+HEK293T-iPSC clones showing elevated expression of glycolysis-associated transcripts by gene set variation analysis (GSVA) (Figures S4A and S4B). Additionally, GSVA showed increased expression of ETC transcripts in BJ p0, BJ p0+PBMC1, and BJ p0+HEK293T SIMR fibroblasts compared to that in native BJ fibroblasts (Figures S4A and S4B). However, immunoblotts for succinate dehydrogenase (SDHb; complex II), ubiquinol-cytochrome c reductase core protein 2 (UUQCRC2; complex III), cytochrome C oxidase II (MT-COXII; complex IV), and ATP synthase F1 subunit alpha (ATPSA; complex V) demonstrated the opposite result, with ETC proteins in SIMR fibroblasts reduced compared to those in the native BJ fibroblast control (Figure S4C). Overall, whole-transcriptome data analysis showed that initial large differences between SIMR clones and native control cells at the fibroblast fate progressively dissipated during reprogramming and differentiation.

We examined the RNA transcript levels of 1,158 nuclear genes listed in the MitoCarta2.0 database that encode proteins that...
localize to the mitochondria. Hierarchical clustering analysis of transcripts from these genes identified a separation of BJ p0+PBMCM1, BJ p0+HEK293T, and BJ p0 transcripts away from the native BJ transcripts at the fibroblast fate (Figures 5B and S3B). Pathway analysis of DEGs between these two groups showed an enrichment for genes encoding ETC proteins in the native BJ fibroblasts (Table S4). Of note, this differential clustering was not observed at the iPSC and MSC fates (Figures 5B and S5A).

Closer examination of mtDNA-encoded transcripts only demonstrated, as anticipated, that BJ p0 cells have dramatically lowered expression of all 13 coding gene transcripts compared to transcripts from SIMR and native fibroblast cells (Figures 5C and S5B). Additionally, both BJ p0+PBMCM1 and BJ p0+HEK293T fibroblasts showed significantly reduced expression of the 13 mtDNA-encoded genes compared to native BJ fibroblasts (Figures 5C and S5B). By contrast, no significant difference was observed in mtDNA transcript levels after reprogramming SIMR and control fibroblasts to iPSCs, followed by differentiation to MSCs (Figures 5C and S5B).

We then used the MitoXplorer pipeline (Yim et al., 2020) to quantify the representation of 38 distinct mitochondrial processes within the identified DEGs. As anticipated from abolished respiration, BJ p0 fibroblasts showed DEGs for 29 mitochondrial processes compared to native BJ fibroblasts, especially within oxidative phosphorylation, mitochondrial genome translation, and amino acid metabolism processes (Figure 5D). Similarly, BJ p0+PBMCM1 and BJ p0+HEK293T fibroblasts had altered gene expression profiles in 30 and 29 mitochondrial processes, respectively, in comparison to native BJ fibroblasts (Figures 5D and S5C). Further analysis at this fate using MitoXplorer highlighted differences between the two SIMR lines, as the BJ p0+PBMCM1 had fewer DEGs for mtDNA-associated oxidative phosphorylation and mitochondrial genome translation compared to BJ p0+HEK293T fibroblasts. Furthermore, both SIMR fibroblast lines had more DEGs associated with nuclear-encoded mitochondrial translation and calcium signaling and transport than the BJ p0 line, when compared to native BJ fibroblasts. Of note, the number of affected mitochondrial processes was dramatically reduced by reprogramming, with only 7 and 16 processes exhibiting DEGs in BJ p0+PBMCM1-iPSCs and BJ p0+HEK293T-iPSCs, respectively (Figures 5D and S5C). Finally, BJ p0+PBMCM1-MSCs and BJ p0+HEK293T-MSCs exhibited only 3 and 8 mitochondrial processes with DEGs, respectively, with only ROS defense similarly altered between the two SIMR-MSC lines and MSC control (Figures 5D and S5C). These data uncover transcriptomic alterations to translation, among additional mitochondrial processes, at the fibroblast fate as a potential difference maker for exogenous mtDNA function in SIMR cells. Combined, the results show that, although SIMR cell mitochondrial function becomes more similar to that of the BJ control with reprogramming and differentiation, differences still exist that are based on the transferred mtDNA sequence.

**DISCUSSION**

Here, we use MitoPunch to report on a rapid, versatile pipeline to generate transformed or non-immortal cells with specific mtDNA-nDNA combinations, an advance with certain advantages over cybrid technology (Patananan et al., 2016; Wong et al., 2017), uncontrolled selection in physiologic mitochondrial “bottlenecks” (Latorre-Pellicer et al., 2019), or time-consuming screens for cells with desired mtDNA mutations (Fayzulin et al., 2015; Lorenz et al., 2017). We show that the transcriptome and metabolome of SIMR fibroblasts resemble those of p0 matched recipient cells and that reprogramming to pluripotency followed by differentiation resets these profiles to closely resemble those of unmanipulated control cells. Our studies would be difficult or impossible using other mitochondrial transfer approaches, such as those that use immortal cell lines incapable of reprogramming. Although it is also possible to generate SIMR cells with the nanoblade and microinjection, these low-throughput methods suffer practical and experimental limitations. In contrast, MitoPunch is an accessible approach to generate numerous clones with desired, stable mtDNA-nDNA combinations within 2 weeks.

SIMR clone formation was achieved for all p0 recipient fibroblasts studied. Of note, some mtDNA-nDNA combinations produced SIMR clones at lower efficiencies than other combinations, which could only be detected using a high-throughput platform like MitoPunch. In contrast to 143BTK–p0+HEK293T...
SIMR cells that produced ~75 clones, MitoPunch transfers into primary fibroblasts yielded fewer SIMR clones, possibly related to the Hayflick limit or a sub-optimal response to the MitoPunch procedure. For example, ADF p0 recipients grew the slowest, reached senescence shortly after mtDNA depletion (data not shown), and resulted in the fewest SIMR clones. On average, twice as many BJ p0+HEK293T clones were obtained compared to BJ p0+PBMC1 clones, which may be from a more compatible mtDNA-nDNA pairing and more favorable metabolic profile (Latorre-Pellicer et al., 2016). SIMR fibroblasts also showed a 10-fold reduction in reprogramming efficiency compared to native fibroblast controls. A similar reduction was observed in mouse embryo fibroblast reprogramming with non-native mtDNA, perhaps from a lower mtDNA-nDNA compatibility (Latorre-Pellicer et al., 2019).

Evidence for incomplete mtDNA-nDNA assimilation in SIMR fibroblasts was observed in transcriptome data that showed hundreds of DEGs between SIMR and unmanipulated fibroblasts. Also, MitoCarta2.0- and mtDNA-encoded transcripts were most similar for SIMR and p0 fibroblasts, despite SIMR cell culture in restrictive medium requiring ETC activity for growth and survival. These data suggest that exogenous mtDNA in SIMR fibroblasts do not fully communicate and influence the nDNA, despite being able to support a selectable level of ETC activity and adequate synthesis of metabolites for cell proliferation. Supporting evidence for this suggestion includes that ddC exposure yielded minimal nDNA damage and that the metabolome and transcriptome profiles of SIMR fibroblasts are mostly reset to unmanipulated BJ-iPSC profiles in SIMR-iPSCs. Our results agree with a report showing that p0 cells have altered metabolism and an epigenome that can only be partially reset by cybrid formation (Smiraglia et al., 2008). Our study provides a platform for investigating the resetting of p0 cell transcription and metabolism after stable mtDNA transplantation and subsequent cell fate changes. Further work is needed to determine whether cells that receive exogenous mtDNA by other forms of mitochondrial transfer also have disrupted mtDNA transcription profiles.

In summary, we provide a proof-of-principle mitochondrial transfer pipeline to generate cells of different fates with specific mtDNA-nDNA combinations, including clonal lines with genome pairings not found in nature. Future studies will generate SIMR-derived cells representing high-energy-demand tissues, such as cardiomyocytes or neurons, and will investigate the current inability to generate SIMR-iPSCs containing mutant mtDNAs to enable patient-specific disease and drug screening models with isogenic nuclei for mtDNA diseases. Furthermore, our results show that the interpretation of mitochondrial transfer experiments must consider that cells initially generated may not show complete mtDNA-nDNA integration and subsequent restoration of cellular pathways. This is particularly salient for cells that lack subsequent reprogramming potential, such as transient or transformed mitochondrial transfer cell lines, since our results show that reprogramming and differentiation are required for resetting the nDNA expression profile. Finally, this mitochondrial transfer pipeline bypasses the evolutionary pairwise selection of mtDNAs and nDNAs in cells to expand upon the repertoire of genomic combinations present in the human population and generates a library of cells at various fates with defined mtDNA-nDNA combinations and unique functional properties for research and potential therapeutic applications.

**STAR METHODS**

Detailed methods are provided in the online version of this paper and include the following:

- **KEY RESOURCES TABLE**
- **RESOURCE AVAILABILITY**
  - Lead Contact
  - Materials Availability
  - Data and Code Availability
- **EXPERIMENT MODEL AND SUBJECT DETAILS**
  - Cell Lines
  - Human Tissues
- **METHOD DETAILS**
  - mtDNA Depletion and qPCR Verification
  - Mitochondrial Transfer into p0 Recipients
  - Confocal microscopy
  - Mitochondrial Oxygen Consumption Measurements
  - Mitochondrial Isolation from Mouse Tissues and Delivery
  - iPSC Reprogramming
  - MSC Differentiation
  - Human mtDNA D-Loop Sequencing
  - ROS Quantification
  - iPSC Flow Cytometry
  - MSC Flow Cytometry
  - Fluorescence Microscopy
  - MSC Immunosuppression Assay
  - Tri-lineage Differentiation
  - Tri-lineage Differentiation Analyses
  - UPHLC-MS Metabolomics Processing
  - RNA Extraction
  - RNA-Seq Library Preparation
  - Restriction-Fragment Length Polymorphism Heteroplasmy Assay

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**Figure 5. Transcriptome Features of SIMR and Native mtDNA Cells**

(A) Number of DEGs for BJ p0+PBMC1 cells compared to native BJ cells at fibroblast, iPSC, and MSC fates.
(B) Hierarchical clustering of nuclear-encoded MitoCarta2.0 database genes from native BJ, BJ p0, and BJ p0+PBMC1 cells at fibroblast, iPSC, and MSC fates.
(C) Normalized, batch-adjusted read counts shown as box-and-whisker Tukey plots for 13 MitoCarta2.0-annotated mtDNA-encoded genes for native BJ, BJ p0, and BJ p0+PBMC1 cells at the fibroblast, iPSC, and MSC fates. Statistical significance was by Welch’s t test.
(D) Mitoxplorer-categorized DEGs for BJ p0+PBMC1 compared to native BJ cells at the fibroblast, iPSC, and MSC fates divided into the 38 mitochondrial processes.

See also Figures S2, S3, S4, and S5 and Tables S3, S4, and S5.
QUANTIFICATION AND STATISTICAL ANALYSIS
- mtDNA Depletion and qPCR Verification
- Mitochondrial Oxygen Consumption Rate Measurements
- Metabolomics Data Analysis
- RNA-Seq Pre-Processing
- Differential Gene Expression Analysis
- MitoXplorer Analysis
- Gene Expression PCA and hierarchical clustering
- Metabolic Transcript Gene Set Variation Analysis (GSVA)
- Gene Set Overrepresentation Analysis (ORA)
- HLA Class I Genotyping

SUPPLEMENTAL INFORMATION
Supplemental Information can be found online at https://doi.org/10.1016/j.celrep.2020.108562.

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AUTHOR CONTRIBUTIONS
Conceptualization: A.N.P., A.J.S., T.-H.W., S.R., K.R.N., and M.A.T.; Methodology: A.N.P., A.J.S., T.-H.W., F.M.A., A.V., M.A.T.; Software: T.L.N., F.M.A., A.V., and D.B.; Formal Analysis: A.N.P., A.J.S., T.-H.W., F.M.A.; Data Curation: A.N.P., F.M.A., A.V., D.B., and C.V.; Writing – Original Draft: A.N.P., A.J.S., F.M.A., and D.B.; Writing – Review & Editing: A.N.P., A.J.S., F.M.A., T.-H.W., K.R.N., and M.A.T.; Visualization: A.N.P. and A.J.S.; Supervision: A.N.P. and T.-H.W.; Project Administration: A.N.P. and T.-H.W.; Funding Acquisition: A.N.P., A.J.S., S.R., K.R.N., P.-Y.C., K.P., and M.A.T.

DECLARATION OF INTERESTS
M.A.T. and P.-Y.C. are co-founders, board members, shareholders, and consultants for NanoCav, a private start-up company working on mitochondrial transfer and quantitative phase microscopy techniques and applications. T.-H.W. was an employee of NanoCav and is employed by ImmunityBio, and S.R. and K.R.N. are board members of NanoCav and employed by ImmunityBio. The other authors report no competing interests.

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## STAR+METHODS

### KEY RESOURCES TABLE

| REAGENT or RESOURCE                           | SOURCE                        | IDENTIFIER          |
|------------------------------------------------|-------------------------------|---------------------|
| **Antibodies**                                |                               |                     |
| OCT3/4                                        | BD Bioscience                 | Cat#561628, RRID: AB_10895977 |
| SOX2                                          | BD Bioscience                 | Cat#561610, RRID: AB_10712763 |
| Mouse IgG1 κ Isotype Control                  | BD Bioscience                 | Cat#557782, RRID: AB_396870 |
| Mouse IgG1, κ Isotype Control                 | BD Bioscience                 | Cat#565373, RRID: AB_1645606 |
| SSEA4                                         | eBioscience                   | Cat#12-8843-42, RRID: AB_11151520 |
| OCT4                                          | eBioscience                   | Cat#53-5841-82, RRID: AB_1210530 |
| TRA-1-60                                      | Stemgent                      | Cat#09-0068, RRID: AB_2233143 |
| TOMM20                                        | Abcam                         | Cat#ab78547, RRID: AB_2043078 |
| dsDNA                                         | Abcam                         | Cat#ab27156, RRID: AB_470907 |
| **Biological Samples**                        |                               |                     |
| LP351 Human PBMCs (PBMC1)                     | HemaCare                      | Donor ID: D326351 |
| LP298 Human PBMCs (PBMC2)                     | HemaCare                      | Donor ID: D316153 |
| **Chemicals, Peptides, and Recombinant Proteins** |                           |                     |
| 2',3'-dideoxycytidine                         | Sigma                         | D5782, CAS 7481-89-2 |
| Dialyzed FBS                                  | Life Technologies             | Cat#26400-044       |
| DMEM without glucose                          | Gibco                         | Cat#11966025        |
| Matrigel                                      | Corning                       | Cat#356234          |
| Lipofectamine RNAiMAX                         | ThermoFisher                  | Cat#13778100        |
| mTeSR 1                                       | StemCell Technologies         | Cat#65850           |
| Phusion high-fidelity PCR master mix with HF buffer | NEB                           | Cat#M05315          |
| Gentle Cell Dissociation Reagent              | StemCell Technologies         | Cat#07174           |
| Accutase                                      | BD Biosciences                | Cat#561527          |
| Hoechst 33342                                 | ThermoFisher                  | Cat#R37605          |
| Apa1                                          | NEB BioLabs                   | Cat#R0114S          |
| **Critical Commercial Assays**                |                               |                     |
| DNeasy Blood & Tissue kit                     | QIAGEN                        | Cat#69504           |
| SYBR Select Master Mix for CFX                | Life Technologies             | Cat#4472942         |
| Qproteome Mitochondria Isolation kit          | QIAGEN                        | Cat#37612           |
| **StemRNA-NM Reprogramming kit**             |                               |                     |
| ReproRNA-OKSGM                                | Stem Cell Technologies        | Cat#09530           |
| STEMCCA Constitutive Polycistronic (OKSM) Lentivirus Reprogramming kit | MilliporeSigma            | Cat#SCR510          |
| CytoTune-iPS 2.0 Sendai Reprogramming kit     | Fisher Scientific             | Cat#A16517          |
| STEMdiff Mesenchymal Progenitor Kit           | StemCell Technologies         | Cat#05240           |
| QIAGEN QIAQuick Gel Extraction kit            | QIAGEN                        | Cat#28704           |
| Fixation/Permeabilization Solution Kit with BD GolgiPlug | BD Biosciences            | Cat#555028          |
| Human MSC Analysis Kit                        | BD Biosciences                | Cat#562245          |
| V3 96-well plate                              | Agilent                       | Cat#101085-004      |
| CD4+ T cell isolation kit                     | Miltenyi Biotec               | Cat#130-096-533     |
| Vybrant CFDA SE. Cell Tracer Kit              | ThermoFisher                  | Cat#V12883          |

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### REAGENT or RESOURCE SOURCE IDENTIFIER

| REAGENT or RESOURCE | SOURCE | IDENTIFIER |
|---------------------|--------|------------|
| Dynabeads Human T-activator CD3/CD28 | ThermoFisher | Cat#11131D |
| Mesencult-ACF Basal Medium | StemCell Technologies | Cat#05449 |
| Mesencult Adipogenic Differentiation Kit | StemCell Technologies | Cat#05412 |
| Mesencult Osteogenic Differentiation medium | StemCell Technologies | Cat#05465 |
| ACF Enzymatic Dissociation/Inhibition Solutions | StemCell Technologies | Cat#05426 |
| Mesencult-ACF Chondrogenic Differentiation Medium | StemCell Technologies | Cat#05455 |
| Y-27632 | Stem Cell Technologies | Cat#72304 |
| BCA protein assay | ThermoFisher | Cat#23225 |
| CellROX Green Flow Cytometry Assay Kit | ThermoFisher | Cat#C10492 |
| KAPA Stranded RNA-Seq Kit with Ribo-Erase | Kapa Biosystems, Roche | Cat#07962304001 |

### Deposited Data

| Data Type | Source | GEO | Metabolite relative amounts | N/A |
|-----------|--------|-----|-----------------------------|-----|
| RNaseq count matrices and raw reads | This paper | GEO: GSE115871 | N/A |
| Metabolite relative amounts | This paper | N/A | N/A |

### Experimental Models: Cell Lines

| Model Name | Source | Catalog Number |
|------------|--------|----------------|
| HEK293T | ATCC | CRL-3216 |
| BJ Foreskin Fibroblast | ATCC | CRL-2522 |
| Primary Dermal Fibroblast; Normal, Human, Adult | ATCC | PCS-201-012 |
| Primary Dermal Fibroblast Normal; Human, Neonatal | ATCC | PCS-201-010 |
| Leigh Syndrome ATP Synthase 6 (T8993G) Fibroblast | Coriell Institute | GM13411 |
| Kearns-Sayre Syndrome (common deletion) Fibroblast | Coriell Institute | GM06225 |
| MELAS (A3243G) Cybrid (CL3) | Gift from Douglas Wallace (Children’s Hospital of Philadelphia Research Institute) | N/A |
| Wildtype Cybrid (CL9) | Gift from Douglas Wallace (Children’s Hospital of Philadelphia Research Institute) | N/A |
| MELAS (A3243G) Cybrid | Gift from Carlos Moraes (University of Miami) | N/A |
| MERRF (A8344G) Cybrid | Gift from Carlos Moraes (University of Miami) | N/A |
| Δ Cytochrome B 3.0 Cybrid | Gift from Carlos Moraes (University of Miami) | N/A |
| L929 r0 Mouse Fibroblast | Gift from Jose Antonio Enriquez Dominguez (Centro Nacional de Investigaciones Cardiovasculares Carlos III (CNIC)) | N/A |

### Oligonucleotides

| Name | Source | Catalog Number |
|------|--------|----------------|
| ND1 forward -CCCTA AAAACCGGCAACATCT | IDT | N/A |
| ND1 reverse - CGAT GGTGAGAGCTAAGGGTC | IDT | N/A |
| GAPDH Forward - TGCAC CACCAACTGCTTAGTC | IDT | N/A |
| GAPDH Reverse - GGCA TGGACTGTCATGAG | IDT | N/A |
| RPLP0 Forward -CGA CCTGGAAGTCCAACTAC | IDT | N/A |

(Continued on next page)
Continued

| REAGENT or RESOURCE | SOURCE | IDENTIFIER |
|---------------------|--------|------------|
| RPLP0 Reverse -ATCT | IDT    | N/A        |
| GCTGCATCTGCTTG      |        |            |
| D Loop Forward - TTCCAA | IDT | N/A        |
| GGACAAATCAGAGAAAAAGT |        |            |
| D Loop Reverse - AGCC | IDT | N/A        |
| CGTCTAAACATTTCAGTGTA |        |            |
| RFLP Forward - CTC | IDT    | N/A        |
| GGAGCAGAACCCAACCT   |        |            |
| RFLP Reverse - CGAA | IDT    | N/A        |
| GGTTTTAGTAGGCCCGT    |        |            |

Software and Algorithms

FlowJo Software Version 10.4.2 (FlowJo, LLC) N/A

Salmon v0.9.1 (Harrow et al., 2012; Mudge and Harrow, 2015; Patro et al., 2017) https://github.com/COMBINE-lab/salmon/releases

Statistical Language R v3.6.0 (R Core Team, 2017) https://www.r-project.org/

Bioconductor v3.9.0 (Huber et al., 2015) https://bioconductor.org/news/bioc_3_3_release/

R Bioconductor package tximport v1.12.3 (Soneson et al., 2015) https://support.bioconductor.org/p/106345/

R Bioconductor package DESeq2 v1.24.0 (Huber et al., 2015; Love et al., 2014) https://bioconductor.org/packages/release/bioc/html/DESeq2.html

R package ggpublisher v0.1.6 (Kassambara, 2017) https://www.rdocumentation.org/packages/ggpublisher/releases

R package pheatmap v1.0.12 (Waines et al., 2016) https://www.rdocumentation.org/packages/pheatmap/releases

R package ggpubr v0.1.6 (Kassambara, 2017) https://www.rdocumentation.org/packages/ggpubr/versions/0.1.6

R package gplots v3.0.1 (Kolde, 2015) https://cran.r-project.org/web/packages/gplots/index.html

R package FactoMineR v2.2 (Husson, 2020) https://www.rdocumentation.org/packages/FactoMineR/releases

R package factoextra v1.0.6.3 (Kassambara, 2019) https://www.rdocumentation.org/packages/factoextra/releases

R Bioconductor package Mfuzz v2.38.0 (Kumar and Futschik, 2007) https://bioc.ism.ac.jp/packages/3.6/bioc/html/Mfuzz.html

R package ggplot2 v3.2.0 (Wickham, 2019) https://cran.r-project.org/web/packages/ggplot2/index.html

R Bioconductor package GSVA v1.32.0 (Hänzelmann et al., 2013) https://anaconda.org/bioconda/bioconductor-gsva

R Bioconductor package limma v3.40.6 (Benjamini and Hochberg, 1995; Ritchie et al., 2015) https://www.bioconductor.org/install/

R Bioconductor package clusterProfiler v3.12.0 (Yu and He, 2016) https://bioconductor.org/packages/3.7/bioc/vignettes/clusterProfiler/inst/doc/clusterProfiler.html

R Bioconductor package ReactomePA v1.28.0 (Yu et al., 2012) https://anaconda.org/bioconda/bioconductor-reactomepa

TraceFinder v3.3 (ThermoFisher) Cat#OPTION-30493

MitoXplorer 1.0 (Yim et al., 2020) http://mitoxplorer.ibdm.univ-mrs.fr

RESOURCE AVAILABILITY

Lead Contact
Further information and requests for resources and reagents should be directed to and will be fulfilled by the Lead Contact, Dr. Michael A. Teitell (mteitell@mednet.ucla.edu).
Materials Availability
All materials generated in this study are available upon reasonable request to the Lead Contact, Dr. Michael A. Teitell.

Data and Code Availability
All raw RNA-Seq reads, transcript abundance values, and processed gene count matrices are submitted to the NCBI Gene Expression Omnibus (GEO). The accession number for the RNA-seq reads reported in this paper is GEO: GSE115871. All other data are available upon request. All software used is available either commercially or as freeware. All custom code is available on GitHub at https://bitbucket.org/ahsanfasih/mitoDesigner/src/master/.

EXPERIMENT MODEL AND SUBJECT DETAILS

Cell Lines
HEK293T cells expressing mitochondria-targeted DsRed protein (pMitoDsRed, Clontech Laboratories) were made as previously described (Miyata et al., 2014). Primary, non-transformed human fibroblast sources include BJ (ATCC, Cat. # CRL-2522), ADF (ATCC, Cat. # PCS-201-012), and NDF (ATCC, Cat. # PCS-201-010). Isogenic cybrid cell lines derived from the same patient containing either the homoplasmic A3243G MELAS substitution or homoplasmic WT sequence were obtained from Douglas Wallace (Children’s Hospital of Philadelphia Research Institute). An alternative A3243G MELAS cybrid cell line, in addition to A8344G MERRF and Δcytochrome B 3.0 cybrid cell lines, were from Carlos Moraes (University of Miami). Two primary A3243G MELAS fibroblast lines were from Anu Suomalainen Wartiovaara (University of Helsinki). Primary fibroblasts associated with Leigh Syndrome (T8993G, Cat. # 30-002-CI), GlutaMax (ThermoFisher, Cat. # 35050-061), and non-essential amino acids (MEM NEAA, ThermoFisher, Cat. # 11-0013CV) supplemented with 10% Fetal Bovine Serum (FBS, Hyclone, Cat. # SH30088.03HI0), penicillin-streptomycin (Corning, Cat. # 30-002-CI), GlutaMax (ThermoFisher, Cat. # 35050-061), and non-essential amino acids (MEM NEAA, ThermoFisher, Cat. # 11-140-050). BJ, ADF, and HEK293T-DsRed cells were grown at 37°C and 5% CO2 in complete media containing DMEM (Corning, Cat. # 10013CV) supplemented with 10% Fetal Bovine Serum (FBS, Hyclone, Cat. # SH30088.03HI0), penicillin-streptomycin (Corning, Cat. # 30-002-CI), GlutaMax (ThermoFisher, Cat. # 35050-061), and non-essential amino acids (MEM NEAA, ThermoFisher, Cat. # 11-140-050). BJ, ADF, and HEK293T-DsRed cells grown in complete media containing 50 μg/ml uridine (Sigma, Cat. # U3003). iPSCs were grown on matrigel (Corning, Cat. # 356234) coated plates in mTeSR1 media (StemCell Technologies, Cat. # 85850) according to the manufacturer’s protocol. MSCs were grown in defined, MesenCult-ACF media (StemCell Technologies, Cat. # 05449) following the manufacturer’s protocol. Cells tested negative repeatedly for mycoplasma using a universal mycoplasma detection kit (ATCC, Cat. # 30-1012K).

Human Tissues
The following human tissues were used: PBMC1 (PBMCs from leukopak donor 351, Caucasian female, 42 year old, Donor ID: D326351, HemaCare Corp) and PBMC2 (PBMCs from leukopak donor 298, Hispanic/Latino male, 25 year old, Donor ID: D316153, HemaCare Corp).

METHOD DETAILS

mtDNA Depletion and qPCR Verification
A 1000x stock of ddC (Sigma, Cat. # D5782) was prepared in water and added to BJ, ADF, and NDF cells grown in complete media with 50 μg/ml uridine to an appropriate final concentration. Cells were passaged every 3-4 d with fresh ddC added over 3 weeks. Following ddC treatment, total DNA was extracted (QIAGEN, Cat. # 69504) and mtDNA quantified using SYBR Select Master Mix for CFX (Life Technologies, Cat. # 4472942). mtDNA-encoded MT-ND1 was amplified with the following primers: forward: CCCTAAAACCCGCCACATCT; reverse: CGATGGTGAGAGCTAAGGTC. mtDNA levels were normalized to nucleus-encoded GAPDH using the following primers: forward: TGCACCCACCACTGCTTAGC; reverse: GGATGGACTGTGGTCATGAG. RPLP0 served as an alternative nucleus-encoded gene for normalization using the following primers: forward: CGACCTGGAAGTCCAACCT; reverse: ATCTGCTGCATCTGCTTG. qPCR was run on a BioRad CFX Thermal Cycler using the following protocol: 1) 50°C for 2 min, 2) 95°C for 2 min, and 3) 40 cycles at 95°C for 10 s and 60°C for 45 s. Samples were compared by calculating ΔΔCT and fold differences.

Mitochondrial Transfer into ρ0 Recipients
Mitochondria were harvested from HEK293T-DsRed cells, PBMCs (PBMC1 or PBMC2), or other cell types using a Qproteome Mitochondria Isolation Kit (QIAGEN, Cat. # 37612) following the manufacturer’s protocol. Mitochondrial pellets were re-suspended in 1x PBS, pH 7.4, at 1 mg total protein/ml. Mitochondrial suspensions were delivered into ρ0 cells using MitoPunch. The MitoPunch platform is a force-based mitochondrial transfer device. Briefly, a 5V solenoid (Sparkfun, Cat. # ROB-11015) is mounted on a threaded plug (Thor Labs, Cat. # SM1PL) and inserted into a threaded cage plate (Thor Labs, Cat. # CP02T). Above the solenoid, assembly rods (Thor Labs, Cat. # ER3) support an upper plate (Thor Labs, Cat. # CP02). The upper plate holds a custom machined aluminum washer (outer diameter, 25 mm; inner diameter, 10 mm) that supports a deformable PDMS (10:1 ratio of Part A base: Part B curing agent) fluid reservoir above the solenoid. The PDMS reservoir is composed of a bottom circular layer (25 mm diameter, 0.67 mm height) chemically bonded to an upper circular ring layer (outer diameter, 25 mm; inner diameter, 10 mm; height,
1.30 mm) and can hold approximately 120 μl of isolated mitochondrial suspension. Cells are seeded onto a porous membrane with 3 μm pores (Corning, Cat. # 353181) 24 h prior to mitochondrial transfer.

To perform mitochondrial transfer, this membrane with adherent cells is secured on top of the PDMS reservoir using an upper plate (Thor Labs, Cat. # CP02). The solenoid is controlled by a 5 V power supply mini board (Futurlec, Cat. # MINIPOWER) and powered by a 12V, 3 Amp DC power supply (MEAN WELL, Cat. # RS-35-12). The activated solenoid strikes the center of the PDMS chamber, deforming the bottom circular layer by approximately 1.3 mm. This deformation rapidly injects the mitochondrial suspension through the membrane and into the monolayer cell culture on the opposite side. A tunable MitoPunch prototype was developed by NanoCav LLC with variable plunger force which improves SIMR generation efficiency especially in replication-limited fibroblasts.

As a comparison to MitoPunch, we performed isolated mitochondria coincubation control experiments. An equal number of coincubation recipient cells were seeded alongside MitoPunch recipients in 12 well dishes instead of the porous membrane. After ~24 h, an equal volume of mitochondrial isolate as loaded into the PDMS reservoir for MitoPunch was pipetted into the cell medium of each coincubation recipient well and incubated at 37°C for 2 h before being released, collected, and plated on 10 cm dishes for selection as described below.

For human fibroblasts and mouse recipients, cells were grown in complete media with 50 μg/mL uridine for 4 d following mitochondria delivery and on day 5 post-delivery, cells were shifted to uridine-free complete media prepared with 10% dialyzed FBS (Life Technologies, Cat. # 26400-044). On day 8 post-delivery, cells were shifted to glucose-free, galactose-containing medium (DMEM without glucose, GIBCO, Cat. # 11966025) supplemented with 10% dialyzed FBS and 4.5 g/l galactose. Colonies emerged at approximately 10 d post-delivery and were counted by microscopy or isolated using cloning rings. For human 143BTK−/− recipients, cells were grown in complete media with 50 μg/mL uridine following mitochondria delivery and shifted to uridine-free complete media prepared with 10% dialyzed FBS on day 3 post-delivery, and clones emerged approximately 10 d post-delivery and were quantified.

**Confocal microscopy**

Cells, ~1x10⁵, were plated on glass coverslips (Zeiss, Cat. # 474030-9000) in 6 well dishes in 2 mL of media and cultured for approximately 24 h. The media was aspirated and cells then fixed by incubation of 1 mL freshly diluted 4% paraformaldehyde (Thermo Fisher Scientific, Cat. # 28906) in 1x PBS, pH 7.4, for 15 min at RT. Paraformaldehyde was removed and cells were washed 3x with PBS, and then washed 3x with PBS with 5 min RT incubation during each wash. Cells were permeabilized by a 10 min RT incubation with 0.1% Triton X-100 (Sigma, Cat. # X100). Permeabilized cells were washed 3x with PBS and then blocked by incubation for 1 h at RT with PBS 2% bovine serum albumin (BSA) dissolved in PBS. After blocking, cells were incubated with primary antibodies at 1:1000 dilution in PBS with secondary antibodies (Invitrogen, Cat. #’s A31573 and A21202) diluted 1:100 in 2% BSA blocking buffer against dsDNA (Abcam, Cat. # ab27156) and TOM20 (Abcam, Cat. # ab78547), and then washed 3x with 5 min RT incubation with PBS. After washing, cells were incubated for 1 h with secondary antibodies (Invitrogen, Cat. #’s A31573 and A21202) diluted 1:100 in 2% BSA blocking buffer protected from light at RT, and washed 3x with 5 min incubations with PBS. To mount coverslips on slides, samples were removed from the 6 well dish, dipped in deionized water, dried with a Kimwipe, and mounted using ProLong Gold Antifade Mountant with DAPI (Invitrogen, Cat. # P3691) on microscope slides (VWR, Cat. # 48311-601). Mounted samples were dried protected from light at RT for 48 h prior to imaging with a Leica SP8 confocal microscope.

**Mitochondrial Oxygen Consumption Measurements**

OCR was measured using a Seahorse XF96 Extracellular Flux Analyzer (Agilent). For fibroblasts or MSCs, 1 – 2 x10⁵ cells per well were seeded onto a V3 96-well plate (Agilent, Cat. # 101085-004) and grown overnight before analysis. iPSCs were treated similarly but plated on matrigel-coated V3 plates. A mitochondrial stress test quantified OCR at basal respiration and after the sequential addition of mitochondrial inhibitors oligomycin, carbonyl cyanide-p-trifluoromethoxyphenylhydrazone (FCCP), and rotenone.

**Mitochondrial Isolation from Mouse Tissues and Delivery**

Spleen, liver, lung, bone marrow, heart, skeletal muscle, and kidney were harvested from an ~8 month-old female C57BL/6 mouse. Briefly, tissue was dissociated by passage through a cell strainer using the plunger of a syringe, and mitochondria were isolated from dissociated tissue using the Qproteome Mitochondria Isolation Kit (QIAGEN, Cat. # 37612) following the manufacturer’s protocol. Mitochondrial suspensions were delivered into L929 fibroblasts using MitoPunch. L929 fibroblast recipients were grown in complete media with 50 μg/mL uridine following mitochondria delivery and shifted to uridine-free complete media prepared with 10% dialyzed FBS 2 h before being released, collected, and plated on 10 cm dishes for selection as described below.

**iPSC Reprogramming**

Reprogramming of fibroblast lines to iPSCs was done using the StemRNA-NM Reprogramming kit (Stemgent, Cat. # 00-0076) following the manufacturer’s protocol. Briefly, fibroblasts were plated on a matrigel (Corning, Cat. # 356234) coated 6-well plate at 2x10⁵ cells/well on 0 d. Daily transfections of non-modified (NM)-RNA reprogramming cocktail were performed for 4 d using Lipofectamine RNAiMAX (ThermoFisher, Cat. # 13778100). On 10-12 d, iPSC colonies were identified by staining with TRA-1-60.
with MesenCult -ACF Plus Medium with 10^6 cells/cm^2, and cultured for 2 d on Matrigel with mTeSR1 medium before the medium was changed to STEMdiff -ACF Mesenchymal Induction Medium. STEMdiff -ACF Mesenchymal Induction Medium was changed daily for 3 d, and on day 4, the medium was changed to MesenCult -ACF Plus Medium. Cells were fed again with MesenCult -ACF Plus Medium on day 5. On day 6, cells were collected with Gentle Cell Dissociation Reagent (StemCell Technologies, Cat. # 07174) and passaged onto plastic plates with MesenCult -ACF Plus Medium with 10 μM ROCK inhibitor (Y-27632; Stem Cell Technologies, Cat. # 72304). Daily half-medium changes were made for ~1 week when cells were ~80% confluent. Cells were further passaged by dissociation with ACF Enzymatic Dissociation Solution and resuspended in MesenCult -ACF Plus Medium before further analysis.

**Human mtDNA D-Loop Sequencing**

Total DNA was extracted from 1x10^6 cells using the QIAGEN DNasy Blood and Tissue kit. PCR was performed using Phusion high-fidelity PCR master mix with HF buffer (NEB, Cat. # M0531S) and the following primers: forward – TTCAAGGACAAATCAGATG, reverse – GAAAAAGT, reverse – AGCCCGTCTAAACATTTTCAGTGTA. PCR was run on an Eppendorf vapo.protect thermal cycler at 1) 98°C for 2 min, 2) 30 cycles at 98°C for 15 s, 58°C for 30 s, 72°C for 30 s, and 3) 72°C for 5 min. PCR products were run on a 0.8%–1% agarose TAE gel, extracted with the QIAGEN QIAQuick Gel Extraction kit (QIAGEN, Cat. # 28704), and Sanger sequenced using the same PCR primers.

**ROS Quantification**

CellROX Green Flow Cytometry Assay Kit (ThermoFisher, Cat. # C10492) was used according to the manufacturer’s protocol. Briefly, 7.5x10^4 cells were plated in a 6-well plate ~24 h prior to measurements. 250 μM tert butyl hydroperoxide (TBHP) and 750 μM CellROX reagent were added to the cells ~2 h and 1 h prior to quantification, respectively. Cells were released using Accutase, washed once with FACS buffer (5% FBS in 1x DPBS, pH 7.4), and quantified using a LSRFortessa flow cytometer (BD Bioscience).

**iPSCs Flow Cytometry**

iPSCs were harvested by 15 min RT incubation with Gentle Cell Dissociation Reagent. Cells were centrifuged at 300 x g for 5 min, washed in 1ml DPBS + 10% FBS, and re-suspended in 100 μl BD Perm/Fix Buffer (BD Bioscience). Cells were incubated at 4°C for 15 min and washed twice in DPBS + 10% FBS. Following the second wash, cells were incubated in 50μl DPBS + 10% FBS containing conjugated antibodies. Antibodies used were OCT3/4 AlexaFluor488 (BD Bioscience 561628 1:10), SOX2 V450 (BD Biosciences 557782 1:10), and CD44 PE (BD Bioscience 562245 1:21). Cells were incubated with conjugated antibodies for 30 min for 30 min and then washed twice in DPBS + 10% FBS. Data was acquired on a LSRFortessa flow cytometer (BD Bioscience) and analyzed using FlowJo software (FlowJo, LLC).

**MSC Flow Cytometry**

MSCs were harvested by 5 min, 37°C incubation with Accutase (BD Biosciences, Cat. # 562245) for 30 min and then washed twice in DPBS + 10% FBS. Data was acquired on a LSRFortessa flow cytometer (BD Bioscience) and analyzed using FlowJo software (FlowJo, LLC).

**Fluorescence Microscopy**

iPSCs were cultured on matrigel-coated 6-well plates and fixed with 4% paraformaldehyde for 10 min. Blocking was done for 1 h in 1x PBS, pH 7.4, with 5% FBS and 0.3% Triton X-100. Cells were stained with SSEA4 (eBioscience, Cat. # 12-8843-42) and OCT4 (eBioscience, Cat. # 53-5841-82) antibodies, and Hoechst 33342 dye (ThermoFisher, Cat. # R37605) overnight at 4°C in blocking buffer. Phase contrast and fluorescence images were obtained with a Zeiss Axio Observer Z1 microscope and Hamamatsu EM CCD camera (Cat. # C9100-02).

**MSC Immunosuppression Assay**

MSC inhibition of T cell proliferation was performed as described previously (Hsu et al., 2015). Briefly, MSCs were plated in a 12-well plate the day before assay. PBMCs were isolated by Ficoll gradient from a healthy de-identified leukopak donor. CD4^+ T cells were isolated from PBMCs using the CD4^+ T cell Isolation Kit (Miltenyi Biotec, Cat. # 130-096-533) and labeled with CFDA SE
(ThermoFisher, Cat. # V12883). Labeled CD4⁺ T cells were stimulated with Dynabeads Human T-activator CD3/CD28 (ThermoFisher, Cat. # 11131D) at a ratio of one bead per T cell. T cells were added into MSC cultures at the following T cell:MSC ratios: 1:2, 1:1, 5:1, and 10:1. After 5 d of co-culture, T cell proliferation was measured using CFSE signature dye dilution by flow cytometry.

**Tri-lineage Differentiation**

Adipocytes, osteoblasts, and chondrocytes were generated from MSCs. For adipocyte differentiation, MSCs between passages 3 – 4 were plated on 6-well plates with MesenCult-ACF Basal Medium (StemCell Technologies, Cat. # 05449) at 4 – 5 × 10⁵ cells per well. Differentiation was performed using the MesenCult Adipogenic Differentiation Kit (StemCell Technologies, Cat. # 05412) according to the manufacturer’s protocol. Media changes were done every 3 – 4 d until 13 d. For osteogenic lineage differentiation, MSCs between passages 3 – 4 were plated on a 6-well plate with MesenCult-ACF Basal Medium (StemCell Technologies) at 3 – 4 × 10⁶ cells per well. Differentiation was performed using MesenCult Osteogenic Differentiation medium (StemCell Technologies, Cat. # 05465) according to the manufacturer’s protocol. Medium changes were done every 3 – 4 d until 13 d. For 3-D pellet chondrogenic differentiation, MSCs were first released from T25 flasks using ACF Enzymatic Dissociation/Inhibition Solution (StemCell Technologies, Cat. # 05426) and collected in polypropylene tubes at 2.5 × 10⁶ cells per tube with MesenCult-ACF Chondrogenic Differentiation Medium (StemCell Technologies, Cat. # 05455) according to the manufacturer’s protocol. Medium changes were done every 3 – 4 days until 13 d.

**Tri-lineage Differentiation Analyses**

Osteogenic differentiation was assayed by staining cells with 1% Alizarin Red solution. Medium was removed from cells grown on 6-well plates and cells were washed 3 times with 1X DPBS. Cells were fixed in 4% PFA in 1X DPBS at 4°C for 15 min prior to incubation with 1% alizarin red at RT. Alizarin red solution was aspirated and the cells were imaged using a standard inverted microscope.

Adipogenic differentiation was assayed by staining cells with 0.1% Bodipy solution. Medium was removed from cells grown on 6-well plates and cells were washed 3x with 1X DPBS. Cells were fixed in 4% PFA in 1X DPBS at 4°C for 15 min and washed twice with 1X DPBS prior to a 10 min incubation with 0.1% Bodipy at RT. Bodipy solution was aspirated and the cells were washed with 1X DPBS prior to acquiring phase contrast and fluorescence images with a Zeiss Axio Observer Z1 microscope and Hamamatsu EM CCD camera (Cat. # C9100-02).

Chondrocyte differentiation was assayed by staining chondrogenic spheroids and spheroid sections with 0.1% Safranin O solution. For staining whole spheroids, the medium was removed from the spheroids and they were fixed in 4% PFA for 15 min at RT. The spheroids were washed twice with 1X PBS, pH 7.4, before 15 min incubation with 0.1% Safranin O solution at RT. The stained spheroids were washed twice with 1ml water and transferred by serological pipette to a 48-well dish for imaging. For spheroid section staining, spheroids were fixed in 10% formalin for 18 h, washed twice in water, and placed in 70% ethanol. Spheroids were microtome sectioned by the UCLA Translational Pathology Core Laboratory, tissue placed on microscope slides. Sections were deparaffinized and rehydrated by washes in xylene, ethanol, and water. Unstained sections were stained with hematoxylin and eosin or 0.1% Safranin O for 10 min at RT prior to washing in ethanol. Sections were imaged under a standard inverted microscope.

**UPLC-MS Metabolomics Processing**

Ultra-high-performance liquid chromatography mass spectrometry (UHLPC-MS) was performed as described previously (Xiao et al., 2018) to quantify metabolites from ~7x10⁵ cells. Briefly, cells were rinsed with cold 150 mM ammonium acetate, pH 7.3, followed by addition of ice-cold 80% methanol. Cells were detached with scrapers, transferred into microcentrifuge tubes, and 1 nmol D/L-norvaline added. After vortexing, the suspension was centrifuged at 4°C for 15 min prior to the addition of ice-cold 80% methanol. Cells were lysed with Urea/Lysis Buffer, pH 7.5. After centrifugation, pellets were re-suspended in 58 mM Tris-HCl, pH 6.8, 5% glycerol, and 17 mg/ml sodium dodecyl sulfate and quantified by BCA protein assay (ThermoFisher, Cat. # 23225).

Metabolites were separated on a Luna NH2 (150 mm x 2 mm, Phenomenex) column using 5 mM NH₄AcO, pH 9.9 (buffer A), acetonitrile (buffer B), and the following gradient: initially at 15% buffer B, 18 min gradient to 90% buffer B, 9 min isocratic at 90% buffer B, 7 min isocratic at 15% buffer B. Samples were analyzed with an UltiMate 3000RSLC (Thermo Scientific) coupled to a Q Exactive mass spectrometer (Thermo Scientific) run with polarity switching (+3.50 kV / −3.50 kV) in full scan mode and m/z range of 65-975. Metabolites were quantified with TraceFinder 3.3 using accurate mass measurements (≤ 3 ppm) and retention times of pure standards.

**RNA Extraction**

Fibroblasts, iPSCs, and MSCs were grown in biological triplicates and technical duplicates to 70 – 80% confluence and purified using the RNeasy Mini Kit (QIAGEN, Cat. # 74104) and RNase-free DNase (QIAGEN, Cat. # 79254) following the manufacturer’s protocols. All samples showed a A260/280 ratio > 1.99 (Nanodrop; Thermo Scientific). Prior to library preparation, quality control of the RNA was performed using the Advanced Analytical Technologies Fragment Analyzer (Advanced Analytical, Inc.) and analyzed using PROSize 2.0.0.51 software. RNA Quality Numbers (RQNs) were computed per sample between 8.1 and 10, indicating intact total RNA per sample prior to library preparation.
RNA-Seq Library Preparation
Strand-specific ribosomal RNA (rRNA) depleted RNA-Seq libraries were prepared from 1 µg of total RNA using the KAPA Stranded RNA-Seq Kit with Ribo-Erase (Kapa Biosystems, Roche). Briefly, rRNA was depleted from total RNA samples, the remaining RNA was heat fragmented, and strand-specific cDNA was synthesized using a first strand random priming and second strand dUTP incorporation approach. Fragments were then A-tailed, adapters were ligated, and libraries were amplified using high-fidelity PCR. All libraries were prepared in technical duplicates per sample (n = 60 samples, 120 libraries total), and resulting raw sequencing reads merged for downstream alignment and analysis. Libraries were paired-end sequenced at 2x150 bp on an Illumina NovaSeq 6000.

Restriction-Fragment Length Polymorphism Heteroplasmy Assay
To quantify relative levels of mtDNA containing the A3243G substitution, total DNA was isolated from cells using the DNeasy Blood and Tissue kit (QIAGEN, Cat. # 69504). PCR amplification of the MELAS region to generate a 634 bp product was performed using the following primers: forward – CCTCGGAGCAGAACCCAACCT and reverse – CGAAGGGTTGTAGTAGCCGT. ApaI digestion (NEB Biolabs, Cat. # R0114S) of the PCR product was performed according to manufacturer’s protocol for 2 h at 25°C, and deactivated at 65°C for 20 min. Sample was separated on a 2.5% agarose gel at 100V for 1 h.

QUANTIFICATION AND STATISTICAL ANALYSIS
mtDNA Depletion and qPCR Verification
Statistical details are provided in each figure legend.

Mitochondrial Oxygen Consumption Rate Measurements
Statistical details are provided in each figure legend.

Metabolomics Data Analysis
Data analysis, including principal components analysis (PCA) and clustering, was performed using the statistical language R v3.6.0 and Bioconductor v3.9.0 packages (Huber et al., 2015; R Core Team, 2017). Metabolite abundance was normalized per µg of protein content per metabolite extraction, and metabolites not detected were set to zero. Metabolite normalized amounts were log transformed and then scaled and centered into Z-scores for relative comparison using R base function scale() with parameters “scale = TRUE, center = TRUE”. Heatmaps and Euclidean distance similarity plots were created using the Z-scores in R package pheatmap v1.0.12, and hierarchical clustering was performed using the Euclidean distance measure.

Pathway-level metabolite set enrichment analysis was performed using R Bioconductor package GSVA v1.32.0 (Hänzelmann et al., 2013). Metabolite normalized abundances were standardized using a log2(normalized amounts + 1) transformation, and metabolites per sample were converted to a pathways per sample matrix using function gsva() with parameters “method = gsva, maseq = FALSE, abs.ranking = FALSE, min.sz = 5, max.sz = 500”. GSVA pathway enrichment scores were then extracted and significance testing for multiple transfer conditions was calculated using R Bioconductor package limma v3.40.6, as described above. Pathway metabolite sets were constructed using the KEGG Compound Database and derived from the existing Metabolite Pathway Enrichment Analysis (MPEA) toolbox (Kanehisa et al., 2012; Kankainen et al., 2011).

RNA-Seq Pre-Processing
Fibroblasts, iPSCs, and MSCs were each sequenced in biological triplicates and technical duplicates (n = 60 total samples) to account for variation in extraction and culturing. Raw sequencing reads were converted into fastq files and filtered for low quality reads and Illumina sequencing adaptor contamination using bc2faatfq (Illumina). Reads were then quasi-mapped and quantified to the Homo sapiens GENCODE 28 (GRC38.p12, Ensembl 92, April 2018) transcriptome using the alignment-free transcript level quantifier Salmon v0.9.1 (Harlow et al., 2012; Mudge and Harrow, 2015; Patro et al., 2017). A quasi-mapping index was prepared using parameters “salmon index -k 31 -type quasi”, and comprehensive transcript level estimates were calculated using parameters “salmon quant -A –seqBias –gcBias –discardOrphansQuasi”. Transcript level counts were collapsed to gene level (HGNC) counts, transcripts per million abundances (TPM) and estimated lengths using R Bioconductor package tximport v1.12.3 (Soneson et al., 2015).

Differential Gene Expression Analysis
The resulting sample gene count matrix was size factor normalized and analyzed for pairwise differential gene expression using R Bioconductor package DESeq2 v1.18.1. Expression changes were estimated using an empirical Bayes procedure to generate moderated fold change values with design “~ Batch + Sample,” modeling batch effect variation due to day of RNA extraction (Huber et al., 2015; Love et al., 2014). Significance testing was performed using the Wald test, and resulting P values were adjusted for multiple testing using the Benjamini-Hochberg procedure (Benjamini and Hochberg, 1995). DEGs were filtered using an adjusted false discovery rate (FDR) q value < 0.05 and an absolute log2 transformed fold-change > 0.05.
**MitoXplorer Analysis**

Differential expression analysis was performed using DESeq2, specifying an absolute log2 transformed fold change threshold > 0.5. Result lists including all genes were uploaded to the MitoXplorer 1.0 pipeline (http://mitoxplorer.ibdm.univ-mrs.fr/index.php) comparing SIMR-fibroblasts, -iPSCs, or -MSCs to the corresponding cell fate of the BJ control. DEGs were filtered using a log2 transformed fold-change threshold of 0.05. Subsequently, the number of upregulated and downregulated DEGs for each mitochondrial process were counted.

**Gene Expression PCA and hierachical clustering**

Variance stabilized transform (VST) values in the gene count matrix were calculated and plotted for PCA using R Bioconductor packages DESeq2, FactoMineR, and factoextra, as described in the metabolomics methods (Huber et al., 2015; Love et al., 2014). For PCA of nucleus-encoded mitochondrial protein and mtDNA transcripts, relevant transcripts were extracted using localization evidence derived from MitoMiner v4.0, subsetting VST matrices using genes listed in MitoCarta 2.0 (Calvo et al., 2016; Smith and Robinson, 2016). Clonal heatmaps were prepared using R Bioconductor packages heatmap v1.0.8 and gplots v3.0.1 (Warnes et al., 2016; Kolde, 2015). Hierarchal clustering was performed using R based function hclust and plotted using the dendextend package.

**Metabolic Transcript Gene Set Variation Analysis (GSVA)**

GSVA on metabolic transcripts was performed similarly to metabolomics data as noted above. Pathway-level metabolic gene set enrichment analysis was performed using R Bioconductor package GSVA v1.32.0 function gsva() with parameters “method = gsva, naseq = FALSE, abs.ranking = FALSE, min.sz = 5, max.sz = 500” using a log2(TPM + 1) transformed gene expression matrix (Hänzelmann et al., 2013). GSVA pathway enrichment scores per sample were extracted and assessed for significance using R Bioconductor package limma v3.40.0, as described above except with a Benjamini-Hochberg adjusted P value threshold = 0.01. Pathway metabolite sets were constructed using the KEGG PATHWAY Database, utilizing gene sets annotated to the metabolic pathways overview map HSA01100 (Kanehisa et al., 2012). Significance testing across clones and conditions for each gene set were calculated using Kruskal-Wallis ANOVA.

**Gene Set Overrepresentation Analysis (ORA)**

DEGs were extracted and analyzed for pathway/gene ontology (GO) term overrepresentation using the R Bioconductor package clusterProfiler v3.12.0 and ReactomePA v1.28.0, using a background gene set of all genes expressed with at least one read count in the sample gene count matrix (Yu and He, 2016; Yu et al., 2012). Overrepresented Reactome/KEGG pathways and GO terms were identified across DEG lists and conditions using clusterProfiler function compareCluster() with significance testing cutoffs of p < 0.05, and an adjusted FDR < 0.25.

**HLA Class I Genotyping**

MHC Class I HLA genotypes were identified using OptiType v1.3.1(Szolek et al., 2014). All raw RNA-Seq sample FASTQs were aligned to the HLA Class I reference transcriptome packaged in OptiType using BWA MEM v0.7.17 with standard parameters (Li and Durbin, 2010). HLA subset reads were then analyzed for Class I genotype using OptiType in paired-end RNA mode with standard parameters.