Replication fork rescue in mammalian mitochondria

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Replication stalling has been associated with the formation of pathological mitochondrial DNA (mtDNA) rearrangements. Yet, almost nothing is known about the fate of stalled replication intermediates in mitochondria. We show here that replication stalling in mitochondria leads to replication fork regression and mtDNA double-strand breaks. The resulting mtDNA fragments are normally degraded by a mechanism involving the mitochondrial exonuclease MGME1, and the loss of this enzyme results in accumulation of linear and recombining mtDNA species. Additionally, replication stress promotes the initiation of alternative replication origins as an apparent means of rescue by fork convergence. Besides demonstrating an interplay between two major mechanisms rescuing stalled replication forks – mtDNA degradation and homology-dependent repair – our data provide evidence that mitochondria employ similar mechanisms to cope with replication stress as known from other genetic systems.

DNA replication is an inherently complex cellular event, where any fault in the processes can have dire consequences for the cell. In contrast to test tube conditions, DNA synthesis in a cell is not just a polymerization reaction with standard kinetic properties. Instead, replication forks stall frequently as the replisome encounters template damage, transcription complexes, positive superhelical torsion and protein-DNA complexes 1,2. If replication fork arrest is not relieved in a controlled manner, the condition will eventually result in a double-strand break and the loss of the non-replicated part of the genome in the absence of repair. Consequently, both prokaryotic and eukaryotic cells have evolved a vast arsenal of tools and strategies to remove the various obstacles blocking replisome progression as well as to deal with the DNA damage caused by replication failure.

Compared to nuclear genomes, the mitochondrial DNA (mtDNA) is tiny, 16–19 kb in most metazoans, and exists in hundreds to thousands of copies per cell 3. For a long time it has been thought that mitochondria lack most of the DNA repair pathways existing in the nucleus, therefore preferring mtDNA maintenance quantity over quality, and damaged molecules would be targeted for degradation rather than investing in their repair 4. While the latter could be true for post-mitotic or ephemeral cells in some organisms 5, exhaustive mutation pattern analyses from mouse have concluded that replication errors – not damage – are the main sources of mtDNA mutations 6,7. Similar to nuclear replication forks, mtDNA replication arrests at sites of template damage 8,9, on naturally occurring replication pause sites controlling transcription complex bypass 10,11 or due impaired enzyme function caused by pathological mutations in the replicative helicase TWNK or mitochondrial DNA polymerase Pol γ 12–14. Double-strand breaks resulting from replication stress have been suggested to be the main culprits behind the pathological mtDNA rearrangements observed in many human conditions 15–18.

In contrast to nuclear genome replication, mtDNA replication in mammals is mostly strand-asynchronous, with the lagging strand synthesized with a considerable delay compared to the leading strand 19,18. Replication of the leading strand (heavy or H-strand, due to its higher G to C ratio) initiates at one major origin (OH), resulting in a displaced single-stranded DNA (ssDNA) intermediate, protected by mtSSB 18 and/or active hybridization with preformed RNA 19. Compared to the H-strand, the initiation of the lagging strand (light or L-strand) is considerably more flexible. The major L-strand origin (OL) 16,20 is complemented by more-or-less frequent lagging strand initiation at undefined sites 21,22. The uncoupled leading and lagging strand synthesis means that potential template damage has very different consequences for the synthesis of the two strands. Both in prokaryotes and in the eukaryotic nucleus, damage in the lagging strand template does not significantly hamper replication progression 12, as

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expected from the discontinuous nature of its DNA synthesis – new Okazaki-fragments are constantly primed independently from the preceding ones. If the replisome stalls at a damaged site during leading strand synthesis, a dedicated translesion polymerase can be recruited to the replication fork to synthesize across the damage1. Alternatively, replication of the two strands can be uncoupled with the replicative helicase continuing to unwind while the lagging strand is synthesized. This will result in lesion skipping by upstream priming or fork regression and reset by origin-independent restart1,2. Both, regressed forks as well as broken, partially replicated molecules resulting from the cleavage of stalled replication fork, can be repaired by recombination-dependent replication3,4.

We have previously shown that the archaic primase-polymerase PrimPol is required for lesion skipping and promiscuous lagging strand priming in mammalian mitochondria5,6,7. The involvement of PrimPol in lagging strand priming is highly interesting as it helps to understand some aspects of basic mtDNA maintenance mechanisms, such as the occurrence of fully double-strand DNA (dsDNA) mtDNA replication intermediates that have been previously associated with strand-coupled replication24–26. While it is impossible to differentiate between frequent lagging strand priming and true strand-coupled replication, this might not be mechanistically important. Recent insight into the genome replication of bacteria has demonstrated that DNA replication is often dynamic, with frequent uncoupling and recoupling of leading and lagging strand synthesis27. In essence, replication coupling is mechanistically arbitrary, as fully dsDNA mtDNA replication intermediates can arise when L-strand synthesis catches up with the H-strand synthesis.

Regardless of their origin, fully dsDNA (i.e. RNaseH and ssDNA cleaving enzyme resistant) replication intermediates of mtDNA exist in all tissues, being predominant in metabolically active post-mitotic tissues28,29. mtDNA replication intermediates can arise when L-strand synthesis catches up with the H-strand synthesis. Highly interesting as it helps to understand some aspects of basic mtDNA maintenance mechanisms, such as the occurrence of fully double-strand DNA (dsDNA) mtDNA replication intermediates that have been previously associated with strand-coupled replication24–26. While it is impossible to differentiate between frequent lagging strand priming and true strand-coupled replication, this might not be mechanistically important. Recent insight into the genome replication of bacteria has demonstrated that DNA replication is often dynamic, with frequent uncoupling and recoupling of leading and lagging strand synthesis27. In essence, replication coupling is mechanistically arbitrary, as fully dsDNA mtDNA replication intermediates can arise when L-strand synthesis catches up with the H-strand synthesis.

We have previously reported that UV and oxidative damage as well as inhibition of mitochondrial DNA polymerase Pol γ by ddC result in an increase in dsDNA intermediates concomitantly with the appearance of cruciform (x-form) mtDNA4. To distinguish these x-forms from resembling replication termination intermediates with two meeting replication forks that occur at the non-coding region (NCR) of mtDNA33, we focused on a region spanning nts 12,273–16,012 of mtDNA, generated by Dral restriction (Fig. 1A). The NCR, including the main leading strand origin (OH) and the site of termination, lies outside of this fragment and any strand-asynchronous replication entering the fragment will give rise to so-called slow moving y-arc intermediates (smy) due to the inability of Dral to cleave the displaced ssDNA strand (Fig. 1A,B). Fully dsDNA replication intermediates in contrast are cleaved by Dral to generate y-forms (y). Replication initiating within the analyzed fragment will give rise to replication bubbles (b). All cruciform DNA will be double-stranded and consist of recombination junctions with equally long arms (x) or trident-shaped DNA (rx), the latter generated by regression of the replication fork. Both x-forms can be enhanced by UV treatment (Fig. 1C, Supplementary Fig. S2) and are resistant to the single-stranded DNA-digesting S1 nuclease, which also removes the RNA:DNA hybrid containing intermediates34. Regressed forks and recombination junctions identical to these x-forms are known to accumulate during replication stalling also in other genomes30–32. Note also the relative increase in the dsDNA y-forms and the appearance of replication bubbles (b) after UV exposure (Fig. 1C, Supplementary Fig. S2).

Homology-dependent rescue of broken and reversed forks in prokaryotes as well as the eukaryotic nucleus requires the action of a recombinase5,6,7. Although no dedicated recombinases have been reported from mammalian mitochondria, the replicative helicase TWNK has substantial strand-exchange activity in vitro and is involved in x-form mtDNA induction in a transgenic animal model5,6,7,8 as well as in the human heart5,6. Overexpression of TWNK in cells under normal culture conditions results in an increase of dsDNA replication intermediates (y) and reduced levels of all x-forms (Supplementary Fig. S3A). Remarkably, TWNK overexpression almost completely prevents fork regression following UV exposure (no rx). As knockout of TWNK results in loss of mtDNA, we decided to use cells expressing the dominant-negative TWNK-LD mutation to see if fork regression and recombination junction formation are dependent on the helicase activity. TWNK-LD represents a clinically relevant patient mutation with a 13 aa duplication in the so-called linker region (352–364), resulting in severe impairment of the helicase activity, mtDNA replication stalling and depletion of copy number11. Interestingly, expression of the mutant TWNK did not prevent the formation of either x-form, but dramatically increased the amount of fully dsDNA intermediates, forming both x- and y-forms from resembling replication termination intermediates with two meeting replication forks that occur at the non-coding region (NCR) of mtDNA33, we focused on a region spanning nts 12,273–16,012 of mtDNA, generated by Dral restriction (Fig. 1A). The NCR, including the main leading strand origin (OH) and the site of termination, lies outside of this fragment and any strand-asynchronous replication entering the fragment will give rise to so-called slow moving y-arc intermediates (smy) due to the inability of Dral to cleave the displaced ssDNA strand (Fig. 1A,B). Fully dsDNA replication intermediates in contrast are cleaved by Dral to generate y-forms (y). Replication initiating within the analyzed fragment will give rise to replication bubbles (b). All cruciform DNA will be double-stranded and consist of recombination junctions with equally long arms (x) or trident-shaped DNA (rx), the latter generated by regression of the replication fork. Both x-forms can be enhanced by UV treatment (Fig. 1C, Supplementary Fig. S2) and are resistant to the single-stranded DNA-digesting S1 nuclease, which also removes the RNA:DNA hybrid containing intermediates34. Regressed forks and recombination junctions identical to these x-forms are known to accumulate during replication stalling also in other genomes30–32. Note also the relative increase in the dsDNA y-forms and the appearance of replication bubbles (b) after UV exposure (Fig. 1C, Supplementary Fig. S2).

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well as TWNK overexpression prevented replication initiation within the studied fragment, none of the manipulations influenced the UV-dependent increase in smy-forms upon UV exposure as reported for PrimPol22.

As MGME1 KO cells showed the most prominent recombination intermediates (x-forms) we have ever observed in cultured cells, we hypothesized these to be linked with the high levels of linear mtDNA forms
that appear as a consequence of MGME1 loss. These 11 kb subgenomic linear fragments map rather precisely to the region between between O2t and O1t and are similar to the ones observed in mice expressing exonuclease-deficient Pol γ (the so-called mutator mice). The O2t-O1t fragments have been suggested to originate from replication fork breakage at a nick left of the replication origin due to a ligation defect caused by incomplete processing of flap structures at these loci. The persistence of these subgenomic linear forms both in the mutator mouse as well as in the MGME1 knockout is explained by the fact that both the 5′-exonuclease activity of MGME1 and the 3′-exonuclease activity of Pol γ are required for the turnover of broken mtDNA. We reasoned that increased replication stress results in more frequent double-strand breaks, which are enriched in MGME1 knockout cells due their inability to turn over broken molecules. The accumulation of linear mtDNA forces the cells to deploy other means to their disposal, such as recombination-dependent repair, which is detectable by an increase of x-forms.

As expected, the MGME1 KO HEK293 cells had high levels of the linear O1t-O2t segment and 7S DNA (Fig. 2), similar to the MGME1 patients5 and knockout mice8. While the level of the subgenomic fragment did not rise upon mild stalling caused by UV treatment, enforced replication after ddC treatment increased it 40-fold compared to untreated cells (Fig. 2B: +96 h, 4 kb fragment in AflII + HindIII digest). Interestingly, a similar fragment as observed in the MGME1 KO cells also appeared in parental HEK293T cells upon ddC treatment, together with an additional band corresponding to a mtDNA dimer in size (Fig. 2C). Whereas the O1t-O2t fragment was lost in normal HEK293T cells during the recovery after the ddC treatment, high levels of the fragment remained in MGME1 KO cells.

To see if the MGME1 knockout had any effect on mtDNA topology, we analyzed native undigested mtDNA using 1D- and 2D-AGE (Fig. 3, Supplementary Fig. S4). Again, the most prominent difference between parental HEK293T and knockout cells were observed in the cells recovering from ddC. MGME1 KO cells had high levels of linear mtDNA forms, including a 33 kb species corresponding to dimeric mtDNA in size (Fig. 3C). Additionally, novel “eyebrow” forms (cb in Fig. 3C) were also observed (Fig. 3D). These forms start from the open circular species (oc) and form an arc that is growing in size in the first dimension, thus behaving similarly as replication intermediates. Further analysis of the non-linear forms revealed that in contrast to the control cells showing bubbles and circles, the MGME1 KO cells had mainly y-shaped replicative molecules (Fig. 3E,F), which arise from broken replication forks with one end attached to O2t or O1t (Supplementary Fig. S5). Although the MGME1 knockout had a relatively small (25% reduction) effect on steady-state mtDNA levels under normal conditions, the cells were nearly unable to regain normal mtDNA copy number levels after a ddC treatment (Fig. 4A). Concomitant with the failure to recover mtDNA copy number, MGME1 knockout cells were unable to increase replication initiation from O1t in a similar manner as the wild type cells, as seen by the absence of a bubble arc in the observed fragment (Supplementary Fig. S6). In addition, replication initiation from alternative origins as a response to stalling was abolished in MGME1 KO cells (Fig. 4B,C), as also seen after UV treatment (Fig. 1D).

2D-AGE is a powerful tool to achieve a snapshot of all molecular forms present in a sample, but is dependent on the correct interpretation of the observed patterns. To consolidate our findings, we analyzed the mtDNA from untreated and 72 h ddC treated wild type HEK293T as well as MGME1 knockout cells using transmission electron microscopy (Table 1). As expected, forced replication during recovery after ddC treatment caused a marked increase in replicative forms resulting from replication stalling, with a much stronger influence on MGME1 knockout cells (8% vs. 19% replicative forms, Table 1). The majority of these molecules were broken at one end of the replication bubble (Supplementary Fig. S7), as already evident in the 2D-AGE analysis (Fig. 3C-F, Supplementary Fig. S5). Although these molecules might have been broken during sample preparation, it is highly unlikely that the breakage would mainly affect the MGME1 KO cells (compare also the difference in the linear arc between control and knockouts in Fig. 3C). Concomitant with our observations using 2D-AGE, the MGME1 KO cells also had higher levels of complex mtDNA forms containing two or more molecules (Fig. 5A, Supplementary Fig. S8), representing either recombining or catenated DNA, which also increased under ddC treatment (Table 1, Supplementary Fig. S4). Although linear mtDNA fragments are impossible to differentiate from contaminating nuclear DNA, we observed linear molecules corresponding to the size of dimeric mtDNA complexed with circular molecules in ddC treated MGME1 knockout cells (Fig. 5B, see also Fig. 3C) as well as cruciform molecules (Fig. 5C). Other odd molecules observed from MGME1 KO cells included several circular forms of about 6 kb in size (Supplementary Fig. S8). Interestingly, in parental HEK293T cells ddC treatment induced a specific molecular class with two replication events in one molecule (Fig. 6), likely corresponding to the observed increase in non-conventional replication bubbles under replicative stress (Figs 1, 4, Supplementary Figs S2 and 3).

Discussion

Replication stress or global impairment of DNA replication results in the transient slowing down or stalling of replication forks. In mitochondria, this has been suggested to be a major contributor for pathological mtDNA maintenance, especially due to the mainly strand-asynchronous manner of mtDNA synthesis. Damage on the leading strand template will stall the DNA polymerase, allowing the by-pass by a re-priming event (Fig. 7C).
If the replicative helicase (Supplementary Fig. S3)\textsuperscript{12,13} or the synthesis of both strands is stalled, as in the case of ddC treatment (Figs 2–4), re-start is no longer an option. In the nucleus such stalled forks can be rescued by converging forks from replication origins nearby (Fig. 7D), where a subset of replication origins are activated only during replication stress conditions\textsuperscript{44}. Interestingly, also in mitochondria replication stress results in the appearance of replication bubbles outside of the canonical replication origins (Fig. 1, 4 and 7B, Supplementary Figs S2 and 3)\textsuperscript{8} and in mtDNA molecules having two replication events (Fig. 6). As replication stress seems to specifically cause replication initiation outside of NCR, mitochondrial must possess mechanisms to license and prime alternative replication origins, which enable the completion of replication if the stalled fork cannot be resolved by other means. In bacterial systems, replication initiation outside of the canonical origins is primarily controlled by a RecBCD complex-mediated strand-invasion event\textsuperscript{45}. As MGME1 is a RecB-type exonuclease\textsuperscript{39} and the loss of MGME1 activity abolishes stress-induced replication initiation (Figs 1D and 4D), existence of similar mechanisms in mitochondria should be investigated. This type of mechanism would also have considerable flexibility as it would not be dependent on the existence of specific origin sequences. Interestingly, the first step
Figure 3. Replication stress causes an excess of linear forms and results in topological modification of mtDNA. (A) Topology gel of undigested mtDNA from control and MGME1 knockout (KO) cells treated with ddC or UV; treatments as in Fig. 2. Under normal conditions, mtDNA in HEK293T cells exists as high molecular weight forms, including catenanes of two interlocked circles (cat), open circular monomers (oc) and supercoiled monomers (sc). Some mtDNA is linearized (lin) either naturally or due to extraction procedures. For the assignment of the different molecular forms see Pohjoismäki et al. and Supplementary Fig. S4. While ddC or UV treatment do not have a major influence on the different forms in control cells, MGME1 knockout (KO) cells recovering from ddC exposure have novel molecular forms (black arrowheads) in addition to the 11-kb linear fragment (open arrowhead). (B) Long exposure of a separate re-run of the 72 h ddC samples. Note the excess smearing in the MGME1 knockout cells, probably due to mtDNA breakage. (C) 2D-AGE of undigested mtDNA from untreated and ddC-recovering parental and MGME1 knockout cells. The one-dimensional gel pattern from (A) is given below each panel to provide an approximate landmark. Notice the marked increase in linear forms in MGME1 knockout cells recovering from the ddC treatment. The additional bands marked with black arrowheads in (A) represent dimeric circular forms (2nc) and 33 kb dimeric linear mtDNA. An unusual eyebrow-arc (eb) is present in the MGME1 knockout cells, being more prominent in the cells recovering their mtDNA levels. (D) As the eb begins from the open circles and grows as an arc to the 2nc, it most likely represents replication intermediates broken at one end. (E) These broken intermediates are visible on 2D-AGE of mtDNA fragments cut once upstream of O_h, such as a PvuII digestion produces. Most replicating forms in PvuII-digested mtDNA should consist of replication bubbles (b), converting to double-γ (dy) and termination
intermediates close to the end of the molecule. If the replication bubble is broken at one end, the resulting intermediates will migrate on the y-arc (y). Note that some recently replicated circular forms are not cut by PvuII, because they either are partially single-stranded (ssc) or have RNA incorporation at the restriction site (oc). For further details regarding these forms and their connection to mtDNA replication, see Torregrosa-Muñumer et al.22. Again, the majority of the molecules are non-replicative and linearized (ln) by the restriction enzyme. (F) While the control cells have almost solely full-length replication bubbles, MGME1 knockout cells have mainly y-form replication intermediates, consistent with frequent breakage of the replication bubbles (see also Supplementary Fig. S5). The difference in the abundance of the circular forms (oc, ss) between the wild-type and MGME1 knockout is likely caused by fewer molecules completing replication due to the premature arrest and breakage of the replication intermediates, resulting in depletion of the newly replicated molecules.

of the RecBCD-dependent replication initiation is the formation of a displacement loop (D-loop)46, a prominent but enigmatic feature of also mammalian mtDNA, that is modified in MGME1-deficient cells40. Our observations on stress-induced replication could also explain the biological significance of replication initiation outside of the canonical origins, which have been known for over a decade24,26,47, but whose significance has been heavily disputed16,21,48. In contrast to replication initiating at O₁, replication from the stress-induced origins is bidirectional and strand-coupled, as evident from the S1 nuclease-resistant bubble arcs that span the entire restriction fragment length in 2D-AGE analysis (Figs 1, 4 and 7B, Supplementary Figs S2 and 3). As pointed out before, this type of replication can be confused with frequent lagging-strand priming49, which also occurs under stress conditions22. Due to this similarity, separation of the two mechanisms is technically challenging. Similarly, nothing is known about the replication origin licensing or initiation in mitochondria and the experimental interconnection of the different origins is currently impossible. The fact that overexpression of TWNK can effectively repress this alternative origin initiation in mitochondria (Supplementary Fig. S3A) could simply indicate that excess helicase availability can forcibly push the stalled fork forward, allowing re-priming to take place.

If a stalled fork is fully duplex DNA, as expected once the lagging strand synthesis has caught up with the leading strand, the fork can be rescued by a mechanism involving fork regression and restart43. Fork regression results in a four-way junction – chicken foot – where the newly synthesized strands are annealed at the free end (Fig. 7E: ii). On 2D-AGE these structures migrate as typical spikes or cones originating from the y-arc31,32, and these spikes/features (rx) appear also in mtDNA after damage (Figs 1, 4 and 7D). As of note, x-forms are sensitive to branch migration and more likely lost than generated as an artefact during the DNA extraction (Supplementary Fig. S9). In the nucleus, the RAD51 recombinase is required for the first step of fork regression45, while bacterial systems use helicases such as RuvAB and RecG41.

TWNK helicase has been shown to possess strand-exchange activity in vitro52,53 and to be able to induce mtDNA recombination in vivo55. Interestingly, TWNK overexpression reduced the regressed forks (rx) and recombination junctions (x) during replication stalling (Supplementary Fig. S3A). This is in contrast to the effects in some solid tissues, where x-forms are specifically induced by TWNK overexpression54,55. It should be noted that this phenotype in transgenic mice is restricted only to tissues having mainly dsDNA mtDNA replication intermediates such as skeletal muscle and heart, indicating a fundamental difference in the mtDNA maintenance strategies in different tissues28. The effect observed in cultured cells is quite likely due to a more efficient lesion bypass instead of initiation of alternative origins (note the lack of b in Supplementary Fig. S3A). Alternatively, excess TWNK might increase the turnover of the regressed forks. In fact, TWNK has been shown to possess potent 5′ to 3′ DNA branch migration activity54, which might be required for the turnover of the rx-forms. In contrast, the expression of the catalytically impaired dominant-negative TWNK-LD did not suppress the appearance of the regressed forks, and the cells maintained replication initiation from alternative origins (Supplementary Fig. S3A). The experiment demonstrates that the helicase activity of TWNK is not required for fork regression (rx-arc), formation of recombination junctions (x-arc) or replication initiation from alternative origins.

In the nucleus, fork rescue continues with the repair and restart of the stalled fork by a homologous recombination-mediated mechanism12 (Fig. 7E). Fork regression will initially enable the nascent lagging strand to be used as a template to continue the leading strand synthesis, resulting in a chicken foot structure that stabilizes the stalled fork (Fig. 7E: i – ii). If the cell is unable to repair the damage on the leading strand template, the 5′-end at the chicken foot structure is enzymatically degraded, resulting in a free 3′-end that can strand invade upstream of the lesion (Fig. 7E: iii vs. iv). The initial strand-invasion event will eventually form true Holliday-junctions (x-forms) after the replication has finished (Fig. 7E: v). As of note, the required strand-invasion step is in principle identical to the replication template switching observed to occur upon double-strand breaks in human mitochondria and highly similar to the replication-dependent break repair seen in T7 phage56. In T7 phage, the gp2.5 single-strand DNA binding protein is responsible for the strand-invasion step56, whereas in mammalian mitochondria this reaction might be catalyzed by TWNK52,53, independently of its helicase activity. Molecular recombination is also the main rescue mechanism of broken replication intermediates and an important genome maintenance mechanism from bacteria to vertebrates56 (Fig. 7F).

The accumulation of linear subgenomic fragments, corresponding to broken replication intermediates, in patients with an MGME1 loss-of-function mutation39,40 or knockout mice38 are likely a consequence of two independent events. They are probably generated by defective ligation of nascent strands at the origin due to faulty processing of DNA flap structures42 and then accumulate due to the inability of MGME1-deficient cells to degrade linear mtDNA fragments57. We show these molecules to arise from site-specific stalling and breakage at either one of the main replication origins (Supplementary Fig. S5), resulting in tailed intermediates (Fig. 3D–F, Supplementary Fig. S7) that eventually break to form the linear 11 kb fragment. The fact that MGME1 overexpression resulted in a decrease of all x-forms after mtDNA damage (Figs 1D and 3–5) could be linked to an increased turnover of linear mtDNA. The free end of a regressed fork is essentially a
Figure 4. mtDNA copy number recovery is impaired by the loss of MGME1. (A) 72 h ddC treatment of HEK293T, MGME1 overexpressor and MGME1 knockout cells, followed by recovery for 48 and 96 h. As evident from the severe copy number depletion after 72 h, ddC treatment results in an almost complete block of mtDNA synthesis. Parental HEK293T cells recover their mtDNA copy number to normal levels within 96 h after the removal of the drug, whereas the recovery is severely impaired in the MGME1 knockout (KO) cells. All samples are normalized to the untreated HEK293T mtDNA copy number (100%) to show the differences in basal mtDNA levels. Results are presented as mean ± SD (n = 3, from at least two independent experiments) and p values based on one-way ANOVA with post-hoc Tukey HSD Test. (B) 2D-AGE analysis of the Dral 12,273–16,012 fragment from ddC treated (200 µM ddC for 72 h) parental 293 T-REx cells (control) and MGME1 T-REx cells induced with 5 ng/ml doxycycline. (C) Identical analysis for regular HEK293 cells (control) and MGME1 knockout (KO) cells. Treatment with S1 nuclease (right panels) reveals the fully double-stranded intermediates. The lower panels represent longer exposures of the upper ones and the exposures between adjacent panels.
are comparative. The treatment stalls all mtDNA replication intermediates and induces replication initiation outside of O₁ (b, see Fig. 1) in all cells except MGME1 KO. Regressed forks (rx’s) appear different in the ddC treated cells compared to the UV (rx in Figs 1 and 2) and merge with the x-arc. Note the lack of a bubble arc, the prominent x-forms and an unusual arc (arrow) arising from x’s in the MGME1 knockout cells.

double-strand break and its rapid turnover in MGME1 overexpressor cells might result in a repair defect, which could explain the delayed recovery from ddC treatment (Fig. 4A). Conversely, impairment of linear DNA disposal in the MGME1 knockout cells increases the x-forms (Figs 1D, 4 and 5, Supplementary Fig. S4, Table 1), likely indicating an active double-strand break repair. This observation suggests that cells primarily try to discard broken mtDNA, but if this does not work, homology dependent repair is used as a reserve mechanism. The importance of the two mechanisms likely varies between cell types, explaining the abundance of x-forms in some tissues but not in others28,29. As of note, even untreated MGME1 KO cells have a specific class of dimeric molecules, which are sensitive to the cruciform-cutting endonuclease T7gp3, but not to the decatenating Topoisomerase IV enzyme (Supplementary Figs S4 and S9). Interestingly, both T7gp3 as well as S1 nuclease can convert more complex MGME1 KO mtDNA forms into a 33 kb linear species, further demonstrating that these genomic dimers are generated by recombination in mitochondria, as previously suggested for the human heart mtDNA forms30,35.

While it is impossible to pinpoint to which extent these recombining forms arise from fork rescue (Fig. 7E) or double-strand break repair (Fig. 7F), it is reasonable to assume that both mechanisms work in mitochondria. First, replication stalling per se results in fork regression and x-formation in UV treated cells (Fig. 1) without increasing the levels of the O₁−O০ fragment (Fig. 2B). Secondly, MGME1 KO cells have high basal levels of x-forms (Fig. 1D), correlating with linear fragments (Fig. 2), dimeric mtDNA (Fig. 3C) and junctional molecules (Fig. 5), all consistent with double-strand break repair by recombination. The dimeric mtDNA forms are mostly head-to-tail dimers, as their cleavage with a single-cut restriction enzyme results in 16.6 kb linear (Fig. 3C vs. 3F). Similar rearrangements, including unicircular dimers to pentamers and strand-invasion intermediates, have been reported from cells expressing catalytically impaired TWNK helicase43, suggesting a common mechanism for their formation through replication-stalling-induced double-strand breaks.

The fact that the O₁−O০ fragment appears transiently also in control cells as a result of replication stalling or when mtDNA replication is boosted to recover the copy number (Fig. 2), is interesting. First, the type and persistence of the damage seems to be important. Although UV treatment causes a distinct replication stalling phenotype, it does not influence mtDNA copy number (Supplementary Fig. S10) or the abundance of the O₁−O০ fragment (Fig. 2B), probably due to the transient nature of the replication stalling compared to the ddC treatment, which leaves blocked 3’-ends that require an exonuclease for repair. Secondly, the increase in the sub-genomic fragment during mtDNA copy number recovery after ddC treatment demonstrates that O₁ and O০ are frequent breakpoints also under normal conditions and that even normal mtDNA replication could contribute to double-strand breaks and subsequent deletion formation. In fact, rearrangement hotspots around O০ are well known both from mice57,58 and humans59,60, further indicating that this replication origin is an important fragile site on mtDNA. Our experiments might also give clues how mtDNA deletions rise during post-natal development in some tissues. Recovery from ddC-induced mtDNA depletion, characterized by rapid amplification of mtDNA to return to the steady-state copy number61,62, emulates physiological conditions where mtDNA levels increase as a response to mitochondrial biogenesis. A boost in mitochondrial biogenesis, such as during muscle development60, could generate a first generation of pathological deletion mutants, which get clonally expanded over time63. Furthermore, antiretroviral therapy using ddC or other similar nucleotide analogues has been shown to result in expansion of mutant mtDNA forms with deletions spanning between O₁ and O০, providing a potential link between replication fork breakage observed in this study and the generation of pathological deletions.

As a conclusion, our study demonstrates that mitochondria show similar responses to replication stress as the nuclear genome, including initiation of alternative replication origins and fork regression. Interestingly, linear mtDNA resulting from breakage of replication intermediates seems to be highly recombinogenic. An efficient MGME1-dependent pathway to dispose broken mtDNA molecules could therefore be a first line of genomic defense against pathological rearrangements. On the other hand, our results also suggest an important, perhaps tissue-specific role for recombination-dependent DNA repair in mitochondria. While the molecular mechanisms to overcome replication stalling are well known from other genetic entities, their elucidation in mitochondria, including the interdependence of x-forms and replication bubbles, remains a challenge for future research.

Methods
An overview of the reagents, cell lines and key equipment used in the study is presented in Supplementary Table S1.

Cell culture, ddC and UV light treatments. HEK293T and HEK MGME1-KO cell lines were cultured in DMEM containing 4.5 g/l glucose, 2 mM L-glutamine, 1 mM sodium pyruvate, 50 μg/ml uridine and 10% fetal bovine serum at 37 °C in a humidified atmosphere with 8.5% CO₂ atmosphere. Flp-In™ T-Rex™ 293 MGME1 wild type (MGME1 OE)62 and TWNK wild-type (TWNK OE) and linker-duplication (TWNK LD) mutant19 cell lines were cultured similarly, and 5 ng/ml of doxycycline was used to induce transgene expression (Supplementary Fig. S1). After the indicated times, cells were harvested and DNA samples were extracted. To induce mitochondrial DNA replication stalling, cells were treated with 3 days with 200 μM of 2′,3′-dideoxyctydine (ddC) (Sigma-Aldrich). For UV damage, cells were exposed on the tissue culture dish with DMEM to a single dose of 30 seconds of 1.34 mJ/cm² at x UVB (302 nm), using a Benchtop 2UVTM transilluminator 302 nm (UVV) instrument4. UV light doses were controlled using a handheld UV radiometer (MINOLTA, UM-25).
Generation of HEK293T MGME1 KO cell line. MGME1 knockout (KO) HEK293 cells were generated using the Crispr-Cas9 approach, as described previously. The plasmid vector carrying a single guide RNA (sgRNA) sequence for MGME1 (5′-AGACCATTTGCAGGCAGCTC-3′) and coding for Cas9 were purchased from Origene. HEK293T cells were plated at a density of 10^4 cells per 96-well and transfected with the plasmid using the GeneJuice transfection reagent (Novagen) on the following day. 48 hours after transfection, cells were resuspended and plated on 96-well plates at 0.5 cells/well concentration. After 2 weeks of growth, cells from single

|                     | HEK untreated | HEK ddC | MGME1 KO untreated | MGME1 KO ddC |
|---------------------|---------------|---------|--------------------|--------------|
| N                   | 400           | 400     | 400                | 100          |
| Catenated or        | 9%            | 15%     | 11%                | 29%          |
| recombining         |               |         |                    |              |
| Dimeric circles     | 0.1%          | 0.2%    | 1.3%               | ND           |
| Replicating         | 5%            | 8%      | 5%                 | 19%          |
| of which broken     | 10%           | 40%     | 57%                | 32%          |

Table 1. Mitochondrial DNA molecule types observed in the transmission electron microscopic (TEM) analysis of parental HEK293T and MGME1 knockout (KO) cells, without or with 72h ddC treatment. N = number of counted molecules. Note that catenated and recombining circles cannot be distinguished from the TEM imagery. Only 100 molecules were counted from the ddC treated MGME1 KO due to high levels of linear mtDNA in these samples and therefore rare dimeric molecules were not detected (ND).

Figure 5. Examples of putative recombining molecules in MGME1 knockout cells. Image interpretation and detail of the boxed area given in panels on the right. (A) 72h ddC treatment of MGME1 KO cells resulted in marked increase in complexes including two or more mtDNA molecules (see Table 1 for quantifications). (B) Dimeric (2n) linear molecule attaching to a monomeric (1n) circle with a cruciform junction. The picture is a montage of two separate TEM images because of the size of the molecule. (C) Two 16.6 kb linear molecules attached to each other with a cruciform junction. The adjacent arms of the molecule are equally long, indicating that the junction is at a homologous locus.
colonies were resuspended in 200 μl medium. Half the volume was transferred to a new well for further growth, while cells from the other half were centrifuged and resuspended in 20 μl lysis buffer (1 mM CaCl₂, 3 mM MgCl₂, 1 mM EDTA, 1% Triton X-100, 10 mM Tris pH 7.5) containing proteinase K (0.2 mg/mL, Qiagen). Incubation at 65 °C for 10 min was followed by heat inactivation at 95 °C for 15 min. Lysates were directly used for PCR amplification and subsequent genotyping by Sanger sequencing. The obtained MGME1 knockout clone was homozygous for the c.37_41del5 frameshift deletion and carried a 197-bp insertion from the cloning vector on one allele.

**Protein isolation and western blotting.** To check protein expression of TWNK-WT-myc, TWNK-LD-myc and MGME1-WT-flag constructs in the respective cell lines (TWNK OE, TWNK LD and MGME1 OE), cellular proteins were extracted as described. 30–40 μg of total protein were separated by 8–12% SDS-PAGE and blotted onto nitrocellulose membrane (Amersham™ Protran™ 0.2 μm, GE Healthcare). Detection of transgenic proteins was performed using anti-flag or anti-myc primary antibodies (Supplementary Table S1), HRP-coupled secondary antibodies and chemiluminescent detection with the BioSpectrum ® 810 imaging System (UVP).

**Total DNA extraction, mtDNA topology and mtDNA species determination by Southern blotting.** Total cellular DNA was isolated using proteinase K and SDS lysis followed by phenol:chloroform extraction and ethanol precipitation. DNA pellets were re-suspended overnight at 37 °C in the presence of...
Figure 7. Possible mtDNA fork rescue mechanisms as interpreted from existing data. (A) Pol γ stalls at a lesion (red triangle) on the leading strand template. In cultured cells, stalling of the leading strand (marked in blue) results in accumulation of partially single-stranded (ss)DNA replication intermediates, increasing the abundance of the smy-forms seen as in the UV-treated control cells. (B) Stalling of the leading strand enables the lagging strand synthesis (pink) to catch up with the replication fork, resulting also in increase of fully double-stranded, S1 nuclease resistant replication forks (y-forms). (C) TWNK continues unwinding and PrimPol re-primes the leading strand synthesis downstream of the lesion. PrimPol also frequently primes the lagging strand synthesis, contributing to the fully dsDNA replication intermediates. Consequently, the accumulation of dsDNA y-forms at the expense of smy is especially pronounced when TWNK helicase is stalled (Supplementary Fig. S3), as also the lagging strand synthesis is halted. (D) Replication initiation from unconventional origins could initiate the rescue of the stalled forks by convergence, giving rise to replication bubbles (b in B, Fig. 6) outside of the canonical origins. (E) If the leading H-strand replication remains stalled for longer periods, L-strand replication can bypass it. In other systems, fork regression is initiated by peeling back the lagging strand (i–ii), enabling the filling in of the stalled leading strand using the nascent lagging
strand as a template and resulting in fork stabilization to a chicken-foot structure (ii). If the blocking DNA lesion is repaired, the regressed part of the fork is cleaved and the fork re-started from the leading strand 3'-end (iii). Alternatively, recession of the 5'-end of the filled-in fork will generate a free 3'-tail that can be used for homology-driven strand-invasion upstream of the lesion (iv), eventually resulting in Holliday-junction formation once replication is completed (v). (F) In cultured cells broken forks are mostly degraded in an MGME1-dependent manner. However, double-strand breaks can also be repaired by homology-dependent replication after 5'-end recession. The 3'-overhang at each end of the broken molecule will strand-invade an intact template and replicate the missing piece (break-induced replication) [2]. A Holliday-junction (x-form) will form once the replication is completed. Cleavage of Holliday-junctions can generate both monomeric as well as dimeric molecules (Figs 3, 5 and S4).

FastDigest BglII (ThermoFisher), which does not cut mtDNA, and the DNA concentration was determined with a NanoDropTM1000 spectrophotometer. For mtDNA topology analysis, 2 μg of BglII digested total DNA were separated on a 0.4% agarose gel in 1 x TBE at 1.2 V/cm for 16 h. Southern blotting and DNA hybridization were carried out as previously [69], using a PCR probe spanning nucleotides 12,992–13,670 (NDS). For mtDNA species (intact mtDNA, 11 kb linear fragment, duplication and 7S DNA) determination, 2 μg of total DNA were digested with FastDigest Aff III and HindIII (ThermoFisher) for 6 hours at 37°C, and then separated on a 0.8% agarose gel in 1x TAE buffer at 25 volts for at least 16 hours. For the visualization of 7S DNA, DNA samples were partially denatured for 10 minutes at 65°C before loading. Southern blotting was performed as before and the blot was probed with a 7S probe spanning nts 12,992–13,670. Probes were labeled using the Rediprime™ II random prime labeling kit (GE Healthcare) and [α-32P]-dCTP (PerkinElmer; 3,000 Ci/mmoll. The autoradiographs were captured on a KODAK storage phosphor screen SO230, detected using Molecular Imager FX (BioRad) and quantified using the associated QuantityOne software.

**mtDNA copy number determination by real-time quantitative PCR (qPCR).** mtDNA copynumber per cell was determined from total DNA using duplex Taqman quantitative PCR as described [68] on an AriaMx Real-Time system (Agilent Technologies).

**Mitochondrial DNA isolation, two-dimensional agarose gel electrophoresis (2D-AGE) and Southern blotting.** Mitochondrial DNA was isolated using 20 μg/ml cytochalasin (Sigma-Aldrich) treatment for 30 minutes prior to cell breakage, followed by differential centrifugation and sucrose gradient purification [63]. The 2D-AGE analysis was performed essentially as in Pohjoismäki et al. [65]. Briefly, 10 μg of mitochondrial nucleic acid were digested according to the manufacturer’s recommendation with FastDigest Dra I (Thermo Scientific) and run on a 0.4% agarose gel in 1xTBE until the fragments of interest had migrated 10 cm into the gel. The gel slab was rotated 90° and a 0.95% agarose gel with 0.5 μg/ml ethidium bromide (EtBr) was cast around it. The second dimension was run in the presence of the 0.5 μg/ml EtBr until the fragment of interest had migrated ~10 cm. Southern blotting and hybridization were performed as described before using the NDS probe. Where indicated, partially single-stranded DNA was degraded by adding 50 units of S1 nuclease (Thermo Scientific) to the digested DNA samples and incubated for 5 minutes at room temperature prior loading the samples onto the gel. The long-range PvuII 2D-AGE analysis was performed as in Torregrosa-Muñumer et al. Uncut long-range 2D-AGE analysis was performed with 10 μg of undigested total DNA, run as a normal long-range 2D-AGE and probed against NDS. The autoradiographs were exposed either on Kodak MS film (Sigma-Aldrich) or captured on a KODAK storage phosphor screen SO230 and detected using Molecular Imager FX (BioRad).

**Transmission electron microscopy (TEM).** Mitochondrial DNA was isolated as for 2D-AGE, treated with 50 units of RNase I (Thermo Scientific) at 37°C for 30 min, subsequently purified using phenol-chloroform extraction and ethanol precipitation and dissolved in TEM grade TE buffer (10 mm Tris-HCl, 0.1 mm EDTA, pH 7.6). mtDNA preparation was visualized by EM following the Kleinschmidt procedure [66]. Briefly, 25–50 ng of mtDNA were diluted in 50 μl aqueous solution of 250 mM ammonium acetate and 100 μg/μl cytochrome c, mixed gently, and the drop allowed to develop for 15–20 min on paraffilm. Cytochrome-c-coated DNA floating on the drop surface was captured by gently touching the drop with a parlodion-coated copper grid. The grids were rinsed in 70% and 85% EtOH solutions, 20 s. each. Following quick air-drying, the grids were placed in a high vacuum evaporator and a thin layer of platinum (80%): palladium (20%) was evaporated on the sample at an angle of 8 degrees at 2 x 10⁻⁶ torr. Finally, the grids were covered with a thin layer of carbon to stabilize the paraffilm drop. Samples were examined in an FEI T12 TEM equipped with a Gatan 2kx2k Orius CCD camera at 40 kV. Contour lengths of the molecules were measured using Digital Micrograph software. Monomeric mtDNA (16,569 bp) equals about 5,500 nm.

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Author Contributions
J.L.O.P. and S.G. conceived and designed the experiments, supervised the experiments and wrote the manuscript. RT-M performed the majority of the experiments, A.H. and J.D.G. conducted the TEM experiments and contributed to the writing of the manuscript. D.B., G.Z. and W.K. generated and characterized the MGME1 knockout cells as well as participated in the manuscript writing.

Additional Information
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