Stepwise maturation of the peptidyl transferase region of human mitoribosomes

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Mitochondrial ribosomes are specialized for the synthesis of membrane proteins responsible for oxidative phosphorylation. Mammalian mitoribosomes have diverged considerably from the ancestral bacterial ribosomes and feature dramatically reduced ribosomal RNAs. The structural basis of the mammalian mitochondrial ribosome assembly is currently not well understood. Here we present eight distinct assembly intermediates of the human large mitoribosomal subunit involving seven assembly factors. We discover that the NSUN4-MTERF4 dimer plays a critical role in the process by stabilizing the 16S rRNA in a conformation that exposes the functionally important regions of rRNA for modification by the MRM2 methyltransferase and quality control interactions with the conserved mitochondrial GTPase MTG2 that contacts the sarcin-ricin loop and the immature active site. The successive action of these factors leads to the formation of the peptidyl transferase active site of the mitoribosome and the folding of the surrounding rRNA regions responsible for interactions with tRNAs and the small ribosomal subunit.
Human mitochondrial ribosomes (mitoribosomes) are responsible for the synthesis of 13 oxidative phosphorylation (OXPHOS) proteins, encoded by the mitochondrial genome.1-3 Due to their unusual architectural features1-3 and the requirement to coordinate mitochondrial ribosomal RNA (rRNA) synthesis with the import of all ribosomal proteins, their assembly is anticipated to involve mitochondrial-specific pathways and participation of both conserved and mitochondrially-specific maturation factors, such as GTPases and methyltransferases.4 Production of functional mitoribosomes is critical since defects in the mitochondrial translation machinery lead to a range of severe human diseases.4

Structural analyses of the yeast and prokaryotic cytoplasmic large subunits (5S–23S) as well as kinetoplastid mitoribosomal large subunit (mt-LSU) assembly intermediates6-10, combined with the biochemical studies of yeast and human mitoribosomal assembly11,12, showed that the solvent exposed side of the mt-LSU matures early, whereas the conserved functional sites of the large subunit, such as the peptidyl transferase center (PTC), the GTPases-associated center (GAC) and the polypeptide exit tunnel, assemble last. At the final stages of the maturation, assembly factors bind to these functional regions to facilitate their maturation and prevent premature translation13. Although human mitoribosome assembly has been extensively investigated using a combination of biochemical and high-throughput approaches12,14-16, structural understanding of this process is currently limited to a late assembly intermediate where a MALSU1–L0R8F8–mt-ACP module is bound.17

To better understand the structural basis of human mitoribosome maturation, we isolated mt-LSU assembly intermediates and investigated their composition and structures using cryo-electron microscopy (cryo-EM). We determined structures of eight distinct states of the human mt-LSU where a total of seven assembly factors (MTERF4, NSUN4, MRM2, MTG2, MALSU1, L0R8F8, and mt-ACP) were bound in different combinations. Our results reveal the conformational changes that allow successive modification and maturation of the functionally important regions of rRNA. The structural data supported by biochemical evidence provide an explanation for the role of the essential NSUN4–MTERF4 heterodimer in the process and specifically in the maturation of the functionally important tRNA-interacting P loop through interplay with the conserved GTPase MTG2 (GTPBP5). The obtained results allow us to propose a stepwise maturation pathway of the functionally important regions in the human mitochondrial large ribosomal subunit (mt-LSU).

**Results**

**Structures of assembly intermediates of the human mitoribosomal large subunit.** To purify assembly intermediates of the human mt-LSU, we transfected human embryonic kidney cells with a tagged mitochondrial GTPase 1 (MTG1) (Supplementary Fig. 1), which is a homolog of bacterial RbgA18,19 and is essential for the production of functional mitoribosomes due to its involvement in the late stages of mt-LSU assembly prior to monosome formation.20,21 The affinity-purified sample was investigated using single-particle cryo-EM to reveal several mt-LSU-like structures containing the entire set of ribosomal proteins. Furthermore, density for several additional proteins was observed at the intersubunit side bound to the rRNA in an immature conformation. Using focused classification around those additional features, we were able to obtain reconstructions of eight distinct cryo-EM classes (overall resolution range between 2.9 and 3.5 Å) corresponding to assembly intermediate states of the mt-LSU (Supplementary Figs. 2 and 3). Two of the classes, states A0 and D, resemble the structures of the assembly intermediates described previously, whereas a MALSU1–L0R8F8–mt-ACP module is bound at the intersubunit face, whereas the rRNA is either disorder or in a nearly mature state, respectively.17 Other classes correspond to novel assembly intermediates (Supplementary Fig. 2). We characterized in detail three of these intermediates, resolved to 2.9, 3.1, and 3.1 Å, and referred to them as states A, B, and C, respectively (Fig. 1 and Supplementary Table 1). They represent the key steps in the late stages of mt-LSU maturation, during which functionally important regions of the rRNA progressively mature. The remaining classes (states C0, D0', and D0") correspond to structurally related, but less complete states that occur between the described intermediates.

**NSUN4–MTERF4 dimer stabilizes the rRNA in a conformation that exposes the active site region.** In addition to the previously characterized MALSU1–L0R8F8–mt-ACP module,17 state A contains at the intersubunit side, below the central protuberance (CP), a dimer of NSUN4 and MTERF4, which were both previously identified as mitoribosomal assembly factors22,23 and their structures determined by x-ray crystallography24,25 (Fig. 1a). Although tagged MTG1 was used for affinity purification of ribosomal assembly intermediates, the factor was not sufficiently ordered on the mt-LSU to be structurally interpreted in any of the observed states. The NSUN4–MTERF4 dimer forms extensive interactions with the rRNA and ribosomal proteins by contacting rRNA helices H66, H75, H81, H87, and H93, and keeps the C-terminal of uL2m and mL48 in an immature conformation.

![Fig. 1 Structures of the key human mitoribosomal large subunit assembly intermediates.](image-url)

The molecular structures of states A, B, and C are shown in surface representation from the intersubunit side. Ribosomal proteins are depicted in light gray, 16S rRNA in beige, and assembly factors NSUN4, MTERF4, MRM2, MTG2, and the MALSU1–L0R8F8–mt-ACP (ACP) module in teal, light green, green, red, and blue, respectively.
Interestingly, NSUN4 is an RNA m^5^C methyltransferase that has been implicated in the assembly of both small and large mitoribosomal subunits, however, biochemical data suggested that in vivo it only modifies the rRNA of the small subunit. Consistently, no RNA substrate was found in the active site of NSUN4 where a clearly visible S-adenosyl-methionine (SAM) cofactor was bound (Fig. 2b and Supplementary Fig. 4). Since NSUN4 lacks an RNA substrate recognition domain that is present in bacterial homologs, it was proposed that it relies on interactions with MTERF4 to be targeted to the mt-LSU to regulate ribosome maturation, as we now observe in the mt-LSU-bound state.

The MTERF4 protein folds into a bent α-solenoid that binds with its convex region to the surface of the immature subunit and exposes its positively charged concave region towards the outside, where we observe a segment of double-helical RNA bound (Supplementary Fig. 5). Although we could not assign the identity of RNA nucleotides in this region, continuous density can be traced from rRNA helix H67, suggesting that the additional density belongs to the H68–H71 region of the 16S rRNA domain IV. The connection from helix H67 (pink) to H68–H71 is shown as dashed lines. The A loop, which is modified by MRM2 in state B, is colored orange. A continuous 10–30% sucrose gradient was used to separate mitochondrial lysates from HEK 293T cells expressing MTERF4 and either the wild-type or a mutated version of NSUN4 to determine their distribution and co-migration with mitochondrial ribosomal fractions. The small and large ribosomal subunit and polysomes in mitochondria isolated from wild-type cells were followed by immunoblotting for mitochondrial ribosomal protein markers of the small (bS16m) and large (bL12m) ribosomal subunits. The input mitochondrial lysate was used as a positive control.

**Fig. 2 Interactions of NSUN4 and MTERF4 with the immature human mitoribosomal large subunit.**

a. The NSUN4–MTERF4 heterodimer (teal and light green, respectively) in state A forms a number of contacts with the mitochondrial large subunit via rRNA (beige spheres) and ribosomal proteins, resulting in rearrangements of the mL48 (yellow) and uL2m (light blue) C-termini (highlighted as circles), as well as stabilization of the H68–H71 region (violet) of the 16S rRNA domain IV. The connection from helix H67 (pink) to H68–H71 is shown as dashed lines. The A loop, which is modified by MRM2 in state B, is colored orange.

b. The NSUN4 (teal) active site with bound S-adenosyl-methionine (SAM) cofactor in state A is shown together with nearby rRNA helices H81, H87, P loop, and PTC loop (beige cartoon). MTERF4 is also shown for orientation (light green). Effects of mutations in NSUN4 on its association with mitoribosomal fractions were analyzed by immunoblotting. A continuous 10–30% sucrose gradient was used to separate mitochondrial lysates from HEK 293T cells expressing MTERF4 and either the wild-type or a mutated version of NSUN4 to determine their distribution and co-migration with mitochondrial ribosomal fractions. The small and large ribosomal subunit and polysomes in mitochondria isolated from wild-type cells were followed by immunoblotting for mitochondrial ribosomal protein markers of the small (bS16m) and large (bL12m) ribosomal subunits. The input mitochondrial lysate was used as a positive control. The blots are representative of results obtained with at least three independent biological experiments. Source data are provided as a Source Data file.

c. Effects of mutations in NSUN4 on its association with mitoribosomal fractions were analyzed by immunoblotting. A continuous 10–30% sucrose gradient was used to separate mitochondrial lysates from HEK 293T cells expressing MTERF4 and either the wild-type or a mutated version of NSUN4 to determine their distribution and co-migration with mitochondrial ribosomal fractions. The small and large ribosomal subunit and polysomes in mitochondria isolated from wild-type cells were followed by immunoblotting for mitochondrial ribosomal protein markers of the small (bS16m) and large (bL12m) ribosomal subunits. The input mitochondrial lysate was used as a positive control. The blots are representative of results obtained with at least three independent biological experiments. Source data are provided as a Source Data file.

d. The NSUN4 (teal) active site with bound S-adenosyl-methionine (SAM) cofactor in state A is shown together with nearby rRNA helices H81, H87, P loop, and PTC loop (beige cartoon). MTERF4 is also shown for orientation (light green). Effects of mutations in NSUN4 on its association with mitoribosomal fractions were analyzed by immunoblotting. A continuous 10–30% sucrose gradient was used to separate mitochondrial lysates from HEK 293T cells expressing MTERF4 and either the wild-type or a mutated version of NSUN4 to determine their distribution and co-migration with mitochondrial ribosomal fractions. The small and large ribosomal subunit and polysomes in mitochondria isolated from wild-type cells were followed by immunoblotting for mitochondrial ribosomal protein markers of the small (bS16m) and large (bL12m) ribosomal subunits. The input mitochondrial lysate was used as a positive control. The blots are representative of results obtained with at least three independent biological experiments. Source data are provided as a Source Data file.

To better understand the role of both MTERF4 and NSUN4 in the context of the mt-LSU assembly, we designed a series of mutants based on our structural results with deletions in key interaction regions. The mutant proteins were then investigated with respect to their ability to form dimers and associate with the mitoribosome using continuous sucrose gradients in cells.
expressing a FLAG-tagged MTERF4 and HA-tagged NSUN4. Wild-type MTERF4 and NSUN4 associate with each other and co-migrate with the large subunit of the mitoribosome, confirming previous findings22–25. Two different mutants were designed to disrupt NSUN4 interactions with the 16S rRNA or MTERF4. The first NSUN4 mutant, bearing a triple mutation K215A, H218A and R275A located at the interface between the rRNA and NSUN4 (Supplementary Fig. 5), was designed to test the contribution of NSUN4 to binding of the complex to the 16S rRNA of the immature mt-LSU. This mutation neither reduced the association of NSUN4 with MTERF4 nor their co-migration with the large subunit (Fig 2c), indicating that MTERF4 plays a predominant role in delivering the complex to the immature subunit. A second NSUN4 mutant carrying a quadruple mutation of residues V65R, R136A, I139R, and R141A (Supplementary Fig. 5), designed to break the dimer between NSUN4 and MTERF4, completely abolished their interaction as shown previously25, and also prevented binding of either of the two proteins to the large subunit (Fig 2c). We conclude that formation of a stable NSUN4–MTERF4 heterodimer is critical for their function in mitoribosome assembly.

**MRM2 methylates the U3039 in the A loop before it adopts mature conformation.** While state A described above contains only the MALSU1–L0R8F8–mt-ACP module and NSUN4–MTERF4 dimer, in state B we additionally observe the MRM2 methyltransferase bound to the rRNA in an optimal position for methylation of its target nucleotide U3039 within the A loop of the 16S rRNA26,27 (Fig. 1b). The A loop is repositioned such that the 2′-O-ribose of nucleotide U3039 faces the active site of MRM2, although we do not observe density for a SAM cofactor (Fig. 2d). The binding site for the MRM2 methyltransferase, as observed in state B, is occupied by helix H71 in the mature mt-LSU (Supplementary Fig. 6). This implies a temporal order of maturation events where U3039 methylation must take place before helix H71 assumes its mature conformation.

**MTG2 and NSUN4 interact with the P loop in a tweezer-like manner.** In state C (Fig. 1c), MRM2 is replaced by MTG2, a GTPase conserved from bacteria to eukaryotes. The bacterial homolog (ObgE) has been recently visualized on the native GTPase conserved from bacteria to eukaryotes. MTG2 is proposed to play a key role in stimulating GTP hydrolysis of translational GTPases10,31 and suggest that the divergent mitochondrial

**Stepwise maturation of the human mitochondrial large subunit.** Visualization of eight distinct assembly intermediates of the human mt-LSU reveal the interdependence of assembly factors and the role of the rRNA in the process allows us to propose a model for stepwise maturation of the ribosomal active site (Fig. 4 and Supplementary Fig. 8). Binding of the NSUN4–MTERF4 dimer to the immature, but compositionally complete mt-LSU, sequesters the flexibly disposed H68–H71 region to expose the active site and allow access of factors that modify the rRNA and check its conformation. This conformation is recognized by the methyltransferase MRM2 that modifies nucleotide U3039 in the A loop of the rRNA. After the methylation of U3039, MRM2 dissociates and the quality checkpoint GTPase MTG2 binds to the P loop in an NSUN4-dependent manner to facilitate maturation of the PTC and to check the functionality of the GAC of the ribosome. Once the rRNA assumes its native or nearly native conformation, the assembly factors dissociate, and a mature, translationally competent mt-LSU is formed. Our data provide the basis for understanding the late stages of human mt-LSU maturation at a structural level and reveal the key role of the NSUN4 and MTERF4 to induce a conformation of the subunit ready for subsequent modification and maturation steps. The described assembly intermediates show maturation and/or proof-reading of all functionally important regions of the large subunit. These results complement the discovery of a remarkably complex mitoribosomal assembly machinery in trypanosomal mitochondria8–10,31 and suggest that the divergent mitochondrial
Fig. 3 Mitochondrial GTPase MTG2 interacts with the functional regions of the immature human mitoribosomal large subunit. a Interaction of the MTG2 (red cartoon) with the 16S rRNA and ribosomal proteins in state C. The N-terminal Obg domain of MTG2 contacts the peptidyl transferase center (PTC) region of the 16S rRNA (light cyan, PTC helices depicted individually, P loop in pink), whereas the G domain associates with the ribosomal GTPase-associated center (GAC) components uL11m (green) and the sarcin-ricin loop (SRL). b Detailed view of G domain interactions with the GAC rotated by 180° relative to panel (a). The Switch loops I and II are schematically indicated as dashed lines. The color key is the same as in panel (a). c Specific interactions of the NSUN4 N-terminal tail (teal) and the MTG2 Obg domain (red) with the P loop (pink). Amino acid residues involved in coordinating the P loop are highlighted. d Effects of N-terminal deletion of residues 26–36 in NSUN4 on its association with mitoribosomal fractions and 55S monosome formation were determined by immunoblotting. A continuous 10–30% sucrose gradient was used to separate mitochondrial lysates from HEK 293T cells expressing the MTERF4 and NSUN4 mutant to determine their distribution and co-migration with mitochondrial ribosomal fractions. The small and large ribosomal subunit and polysomes in mitochondria isolated from NSUN4 ΔN 26-36 transfected cells were followed by immunoblotting for mitochondrial ribosomal protein markers of the small (bS16m) and large (bL12m) ribosomal subunits. The input mitochondrial lysate was used as a positive control. The blots are representative of results obtained with at least three independent biological experiments. Source data are provided as a Source Data file. e Cross-section of the mitochondrial large subunit state C assembly intermediate. The spatial arrangement of assembly factors NSUN4 (teal), MTERF4 (light green), and MTG2 (red) ensure probing of the P loop (pink) and the entrance to the nascent polypeptide tunnel in the mitochondrial large subunit assembly intermediate (gray). The mL45 (blue) N-terminal tail occupies the exit tunnel, contributing to an inactive state of the subunit.
ribosomes are likely to involve an equally diverse set of assembly factors across different species. Furthermore, while this work was in preparation, several manuscripts describing complementary results on the topic of mt-LSU maturation were published or uploaded in pre-print repositories. Taken together, these results now provide a comprehensive description of the late stages of mitoribosomal large subunit assembly in human mitochondria. Furthermore, these results will help us understand the structural basis of mutations in the mitochondrial translational apparatus that are associated with neurodegenerative diseases and diverse progressive and fatal genetic disorders.

**Methods**

**Transient expression of MTG1-3xFLAG in HEK 293 EBNA cells.** HEK 293 EBNA embryonic kidney cells, adapted to suspension growth in serum-free Ex-Cell medium, were obtained from the protein production and structure core facility at EPFL. The cell line was cultured at 37°C under 4.5% CO2 in EX-CELL 293 serum-free medium for HEK 293 cells (Sigma), supplemented with 4 μM t-glutamine. The cells were not tested for mycoplasma contamination.

The pcDNA3.1 (+) plasmid encoding the C-terminally 3xFLAG-tagged MTG1 (UniProt ID Q9B717) was obtained from GenScript. The inserted sequence was verified using a CMV forward primer at Microsynth. Cells with a concentration of 10^6 cells/mL were transfected with 1.5 μg of DNA per liter of the culture using transfection reagent 40 kDa PEI MAX (Polysciences, Inc.) in a 1:2 ratio. Cells were harvested and mitochondria were isolated 72 h post-transfection.

**Isolation of mitochondria from HEK 293 EBNA cells.** Mitochondria were isolated as previously described with few modifications. Briefly, the cell pellets were resuspended in ice-cold RSB hypo buffer (10 mM Tris-HCl pH 7.5, 10 mM NaCl, and 1.5 mM MgCl2) and allowed to swelling for 10 min. The swollen cells were opened with several strokes of a dounce homogenizer followed by immediate addition of 2.5x MS buffer (12.5 mM Tris–HCl pH 7.5, 525 mM mannitol, 175 mM sucrose, 2.5 mM EDTA, and 2.5 mM DTT) to a final concentration of 1x MS buffer (5 mM Tris–HCl pH 7.5, 210 mM mannitol, 70 mM sucrose, 1 mM EDTA and 1 mM DTT). The homogenate was clarified at 13000g for 4°C for 10 min, followed by another 10 min centrifugation at 75000g and 4°C. Finally, the supernatant and the crude mitochondria fraction were separated after centrifugation at 95000g and 4°C for 10 min. The mitochondria pellet was resuspended in 20 mM HEPS–KOH pH 7.6, 250 mM sucrose, and 1 mM EDTA and adjusted to a layered sucrose gradient, consisting of 15%, 25%, 32%, and 60% (w/v) sucrose solutions in 20 mM HEPS–KOH pH 7.6 and 1 mM EDTA. After 70 min ultracentrifugation at 70,000g for 4°C at 4°C, the clarified lysates were loaded on a continuous 10–30% sucrose gradient containing 100 mM KCl, 20 mM MgCl2, 10 mM Tris–HCl pH 7.5, 2% digitonin, 40 μM RNase inhibitor, and 1x chase protease inhibitor cocktail (Roche). After 20 min centrifugation at 80,000g for 45 min at 4°C, the clarified lysates were loaded on a continuous 10–30% sucrose gradient containing 100 mM KCl, 20 mM MgCl2, 10 mM Tris–HCl pH 7.5, and the aforementioned RNase and protease inhibitors, followed by centrifugation at 71,000g in an Optima Beckman Coulter preparative ultracentrifuge. Fractions were collected, and one-third of each fraction was precipitated with 0.02% sodium deoxycholate and 12% trichloroacetic acid, washed twice with acetone, and resolved by SDS–PAGE. Representative markers of the small and large ribosomal subunits were detected by immunoblotting as described below.

**Sucrose gradients of mitoribosomal ribosomes to analyze NSUN4 mutants.** Sucrose gradient fractionation was carried out on purified mitochondria as previously described. Briefly, isolated mitochondria were separated on a 10–30% sucrose gradient and lysed in 260 μM sucrose, 100 mM KCl, 20 mM MgCl2, 10 mM Tris–HCl pH 7.5, 2% digitonin, 40 μM RNase inhibitor, and 1x chase protease inhibitor cocktail (Roche). After 20 min centrifugation at 70,000g for 45 min at 4°C, the clarified lysates were loaded on a continuous 10–30% sucrose gradient containing 100 mM KCl, 20 mM MgCl2, 10 mM Tris–HCl pH 7.5, and the aforementioned RNase and protease inhibitors, followed by centrifugation at 71,000g in an Optima Beckman Coulter preparative ultracentrifuge. Fractions were collected, and one-third of each fraction was precipitated with 0.02% sodium deoxycholate and 12% trichloroacetic acid, washed twice with acetone, and resolved by SDS–PAGE. Representative markers of the small and large ribosomal subunits were detected by immunoblotting as described below.

**Immunoblotting.** Specific proteins were detected using rabbit monoclonal antibodies against MRPL12 (B12m) (16394-1-AP), MRPS16 (B16m) (16735-1-AP), and FLAG (Sigma, F7425); and mouse monoclonal antibodies against HA (Cell Signaling, 2367). All primary antibodies were diluted 1:1000 using the Odyssey blocking buffer (LI-COR). IR dye 800CW goat antibody against IG or IR Dye 680LT goat anti-mouse IgG (LI-COR) secondary antibodies (diluted 1:10,000) were used, and the immunoblots were visualized using the Odyssey Infrared Imaging System (LI-COR).

**Cryo-EM sample preparation and data acquisition.** Quantifoil R2/2 holey carbon copper grids (Quantifoil Micro Tool) were prepared by applying an additional thin layer of continuous carbon, followed by glow-discharging for 15 s at 15 mA using an easyGlow Discharge cleaning system (PELCO). For both datasets, 4 μl of

**Expression of NSUN4 variants in HEK 293T cells.** Expression cassettes for NSUN4 variants were synthesized from overlapping oligonucleotides and cloned into pCMV (Twist Bioscience). All NSUN4 variants were expressed as fusions to a C-terminal HA tag, while MTERF4 was C-terminally FLAG-tagged. Human embryonic kidney (HEK 293T) cells were cultured at 37°C in humidified 95% air with 5% CO2 in Dulbecco’s modified essential medium (DMEM) ( Gibco, Life Technologies) containing glucose (4.5 g/L), l-glutamine (2 mM), 1 mM sodium pyruvate, 50 μg/ml uridine, and 10% (v/v) fetal bovine serum (FBS). The cells were tested and shown to be free of mycoplasma contamination. HEK 293T cells were plated at 60% confluence in 15 cm plates and transfected with mammalian expression plasmids in OptiMEM media (Invitrogen). 138 ng/cm2 of NSUN4 and MTERF4 plasmid DNA, in equal ratios, were transfected using Fugene HD (Roche). Cell incubations were carried out for 72 h following transfection and mitochondria were isolated as described previously.

**Preparation of MTG1-3xFLAG-tagged mitochondrial ribosomes.** Upon thawing of the mitochondria, 15 volumes of lysis buffer (20 mM HEPS–KOH pH 7.6, 100 mM KCl, 20 mM MgCl2, 1.6% Triton X-100, supplemented with 1x complete EDTA-free protease inhibitor cocktail (Roche)) was added, and mitochondria were lysed using a dounce homogenizer. Membranes were further solubilized by stirring for 15 min at 4°C. The lysate was clarified by centrifugation at 20,800g for 4 min at 4°C. Next, the supernatant was incubated for 1.5–2 h at 4°C with anti-FLAG M2 affinity gel (Sigma) while gently mixing. The anti-FLAG M2 affinity gel (Sigma) was pre-equilibrated with three sequential column volumes of 0.1 M glycine HCl pH 3.5 and washed with 10 volumes of TBS and wash buffer (20 mM HEPS–KOH pH 7.6, 100 mM KCl, and 20 mM MgCl2). After collecting the flow-through, the beads were washed with 10 column volumes of wash buffer. Bound mitoribosomes were eluted 3 times using elution buffer (20 mM HEPS–KOH pH 7.6, 100 mM KCl, 20 mM MgCl2, and 100–200 μg/mL 3xFLAG peptide), each time preceded by a 10–15 min incubation at 4°C with gentle mixing. Eluted fractions were pooled and subjected to a 2.5 h ultracentrifugation at 135,500 x g at 4°C using a TLA-55 rotor (Beckman-Coulter). Finally, the mitoribosome pellet was resuspended in a 20 mM HEPS–KOH pH 7.6, 100 mM KCl, 20 mM MgCl2, and 1 mM DTT, yielding mt-LSU at a concentration of ~40 nM. Samples from the key steps of the purification were subjected to western blot analysis (anti-3xFLAG antibody A8592, Sigma) (Supplementary Fig. 1).
resuspended MTG1-3×FLAG-tagged mitoribosome sample was directly applied onto the grid mounted in the Vitrobot chamber (Thermo Fisher Scientific) and incubated for 1 min. Excess of buffer was blotted away in yeast, and the grid was immediately plunge frozen in 1:2 ethane/propane (Carbagas) at liquid nitrogen temperature. The Vitrobot chamber was kept at 4 °C and 100% humidity during the whole procedure. For the second dataset, NP-40 detergent was added to a final concentration of 0.001% just before applying the MTG1-3×FLAG-tagged mitoribosome sample onto the grid.

Both datasets were collected on a Titan Krios cryo-transmission electron microscope (Thermo Fisher Scientific) operating at 300 kV. For the first dataset, the microscope was equipped with a Falcon III E Direct Electron Detector (FEI) and the films were reassembled in integration mode with a pixel size of 1.06 Å/pix, 30 frames, and a total dose of 60 e−/Å², with defocus varying from −3.0 to −0.6 µm. The second dataset was collected on a microscope equipped with a K3 detector (Gatan), mounted to a GIF Quantum LS imaging filter operated with an energy filter slit width of 20 eV. The movies were collected in counting and super-resolution mode, with 40 frames and a total dose of 60 e−/Å² at a physical pixel size of 1.06 Å/pix (0.53 Å/pix in super-resolution) with defocus varying from −3.0 to −0.6 µm. The collection of both datasets was automated with the EPU software (Thermo Fisher Scientific).

**Cryo-EM data processing.** Both datasets were collected and processed independently. Unless stated otherwise, all processing steps were performed using cryoSPARC 3.1.2.

Movies of the first dataset collected on the Falcon III E Direct Electron Detector (FEI) were drift-corrected and dose-weighted using MotionCor2. The corrected 15,441 micrographs were imported into cryoSPARC. The CTF parameters for each micrograph were estimated using patch-based CTF estimation, and 100 randomly selected micrographs were used to pick initial particles using a Laplacian-of-Gaussian filter-based method. These particles were subjected to a 2D classification. Classes resembling the large subunit of the human mitoribosome were used as a reference for picking particles from the whole dataset. The resulting 2,214,072 particles were extracted at 5.6-fold binning and subjected to 2D classification. Good-looking classes were selected and used for an ab-initio reconstruction of the initial 3D model. The initial model was then used as an input in homogenous refinement with all 1,064,609 selected particles from the 2D classification. The resulting refined map was used to create a mask covering the intersubunit side of the large subunit in UCSF Chimera. Aligned particles, together with the mask, were used in a 3D Variability analysis with five modes to solvate and resolve the resolution filtered to 6 Å. The particles were then divided into 10 clusters using 3D variability display.

The second dataset was pre-processed during collection using cryoSPARC live. The pre-processing included 2-fold binning, drift-correction, dose-weighting, CTF estimation, and particle picking using a Laplacian-of-Gaussian filter-based method, resulting in 10,682 processed micrographs and 1,626,837 extracted particles that were exported into cryoSPARC. The particles were then subjected to 2D classification, and classes resembling the mitoribosomal large subunit were selected, resulting in 833,994 particles. All particles were used for ab-initio reconstruction, and the resulting cryo-EM map was used as an input model in homogenous refinement. The previously described intersubunit mask was resampled onto the new map using UCSF Chimera and together with the aligned particles was used for 3D variability analysis with five modes to solve and the resolution filtered to 10 Å. The particles were then divided into 25 clusters using 3D variability display.

Overall, the processing of both datasets resulted in similarly looking 3D classes. The only difference was states B and D0, which were present only in the first and the second dataset, respectively. The key classes found in both datasets were re-extracted at full-size and refined again, this time with per-particle defocus estimation. The local resolution was calculated using a locally windowed FSC method as described in ref. 42.

**Model building and refinement.** Published structures of the mature human mitoribosome (PDB 6ZM6) and of a late assembly intermediate containing the ACP module (PDB 5OOL) were used as initial models and docked into the cryo-EM maps using UCSF Chimera. Composite models were assembled in PyMOL Molecular Graphics System, Version 2.1.5 (Schrödinger, LLC), followed by manual rebuilding of the proteins and nucleic acids using COOT. For interpretation of the additional density features representing the assembly factors, the crystal structures of the human NSUN4-MTERF4 dimer (PDB 4FZV) and MRM2 (PDB 2NYU) were fitted and re-built. For MTG2, an initial model was obtained using the PyMOL2 modeling server based on a crystal structure of the E. coli homolog OgbE (PDB 5M0A). The N-terminal Oub and C-terminal G domains were docked individually and refined. For fitting the CMIN and SMAP co-factors, superimposed high-resolution structures were used as a guide.

The models were real-space refined for five cycles using Phenix version 1.19.131, while applying side-chain rotamer and Ramachandran restraints. The remaining discrepancies between the models and maps were detected and corrected using real space electron density validation tools implemented in COOT. The final model geometry was validated using MolProbity (Supplementary Table 1). To evaluate the quality of the fit of the refined models to the EM maps, real-space correlation coefficients (CCmask) as well as the model versus map FSCs at the FSC = 0.5 criteria were calculated. The resulting resolutions were close to those calculated from the map half-sets at the FSC = 0.143 criteria (Supplementary Fig. 3).

**Figure preparation.** Molecular graphics and analyses were performed with UCSF Chimera and UCSF ChimeraX, developed by the Resource for Biocomputing, Visualization, and Informatics at the University of California, San Francisco, with support from National Institutes of Health P41-GM103311 and R01-GM129325 and the Office of Cyber Infrastructure and Computational Biology, National Institute of General Medical Sciences. Detailed views of cryo-EM map densities were created using The PyMOL Molecular Graphics System, Version 2.1.5 Schrödinger, LLC.

**Reporting summary.** Further information on research design is available in the Nature Research Reporting Summary linked to this article.

**Data availability.** The data that support this study are available from the corresponding author upon reasonable request. The atomic coordinates were deposited in the RCSB Protein Data Bank (PDB) under accession numbers 7OJR (state A), 7ODS (state B), and 7ODT (state C). The cryo-EM maps were deposited in the Electron Microscopy Data Bank (EMDB) under accession numbers EMD-12845 (state A), EMD-12846 (state B), EMD-12847 (state C), EMD-12848 (state A0), EMD-12849 (state C0), EMD-12850 (state D0), EMD-12851 (state D0), and EMD-12852 (state D). Source data are provided with this article.

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Author contributions  

N.B. and T.L. designed the project. T.L., T.S., and M.J. prepared the sample for cryo-EM. T.S. and M.S. established the methods for transfection of HEK cells with affinity-tagged mitochondrial proteins. A.S. and T.L. collected the cryo-EM data. M.J. and T.L. processed the cryo-EM data. M.L. built and performed coordinate refinement of the atomic models. T.L. and M.J. interpreted the structures. O.R., R.L., and A.F. performed in vivo experiments in HEK cells. T.L., M.J., A.F., and N.B. drafted the initial manuscript, all authors contributed to the final version.  

Competing interests  

The authors declare no competing interests.  

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