Respiratory complex I with charge symmetry in the membrane arm pumps protons

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Energy-converting NADH:ubiquinone oxidoreductase, respiratory complex I, is essential for cellular energy metabolism coupling NADH oxidation to proton translocation. The mechanism of proton translocation by complex I is still under debate. Its membrane arm contains an unusual central axis of polar and charged amino acid residues connecting the quinone binding site with the antiporter-type subunits NuoL, NuoM, and NuoN, proposed to catalyze proton translocation. Quinone chemistry probably causes conformational changes and electrostatic interactions that are propagated through these subunits by a conserved pattern of predominantly lysine, histidine, and glutamate residues. These conserved residues are thought to transfer protons along and across the membrane arm. The distinct charge distribution in the membrane arm is a prerequisite for proton translocation. Remarkably, the central subunit NuoM contains a conserved glutamate residue in a position that is taken by a lysine residue in the two other antiporter-type subunits. It was proposed that this charge asymmetry is essential for proton translocation, as it should enable NuoM to operate asynchronously with NuoL and NuoN. Accordingly, we exchanged the conserved glutamate in NuoM for a lysine residue, introducing charge symmetry in the membrane arm. The stably assembled variant pumps protons across the membrane, but with a diminished $H^+/e^-$ stoichiometry of 1.5. Thus, charge asymmetry is not essential for proton translocation by complex I, casting doubts on the suggestion of an asynchronous operation of NuoL, NuoM, and NuoN. Furthermore, our data emphasize the importance of a balanced charge distribution in the protein for directional proton transfer.

Significance

Respiratory complex I is a central enzyme of cellular energy metabolism coupling quinone reduction with proton translocation. Its mechanism, especially concerning proton translocation, remains enigmatic. Three homologous subunits that contain a conserved pattern of charged and polar amino acid residues catalyze proton translocation. Strikingly, the central subunit NuoM contains a conserved glutamate residue at a position where conserved lysine residues are found in the other two subunits, resulting in a charge asymmetry discussed to be essential for proton translocation. We found that the respective glutamate to lysine mutation in \textit{Escherichia coli} complex I lowers the amount of protons translocated per electron transferred by one-quarter. These data clarify the discussion about possible mechanisms of proton translocation by complex I.

The universal cellular energy adenosine triphosphate (ATP) is mainly produced by oxidative phosphorylation (OxPhos). OxPhos couples the activity of respiratory electron transfer chains with that of ATP synthase (1–3). The enzyme complexes of the respiratory chains use the redox energy obtained by the oxidation of the respective substrates to generate a protonmotive force (PMF) across a membrane. The PMF is used in turn by the ATP synthase to drive ATP synthesis (1, 2). Energy-converting NADH:ubiquinone oxidoreductase, respiratory complex I, is the main entry point for electrons from NADH into many electron transfer chains. Complex I couples the transfer of two electrons from NADH to a quinone (Q) with the translocation of four protons across the membrane (4–9). It has a two-part structure, consisting of a peripheral arm catalyzing electron transfer and a membrane arm conducting proton translocation (10). Thus, the complex performs proton-coupled electron transfer over a distance of more than 300 Å. The mitochondrial complex of mammals comprises 45 subunits, 14 of which represent the catalytic core. Homologs of these 14 subunits are found in all species that contain an energy-converting NADH:Q oxidoreductase. Seven of these core subunits are most hydrophobic, and are accordingly all located in the membrane arm of the complex. These subunits are mitochondrially encoded in euukaryotes hampering their genetic manipulation (11), although this is easily done in bacterial complex I, such as that of \textit{Escherichia coli}. This enables the straightforward characterization of effects of mutations in the homologs of the mitochondrial-encoded subunits on the stability, assembly, and catalytic activity of the variant complex (12–17). \textit{E. coli} complex I comprises 13 subunits called NuoA to NuoN (from NADH:ubiquinone oxidoreductase) due to the fusion of the genes encoding subunit NuoCD. This complex represents a structural minimal model of an energy-converting NADH:Q oxidoreductase (18).

The mechanism of proton translocation by complex I is still under debate (19–22). The membrane arm contains four distinct, yet interconnected proton pathways that are linked to the Q binding and reduction site (the Q cavity) by an unusual central axis of polar and charged amino acid residues located approximately in the middle of the lipid bilayer (Fig. 1). This axis is oriented perpendicular to the membrane normal. The proton pathway closest to the Q cavity is called E-channel and is made up of subunits

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NuoH, NuoJ, and NuoK (23). The name stems from the presence of several conserved glutamate residues in these subunits that are most likely involved in proton transfer between the central hydrophilic axis and the Q cavity. Along the membrane arm, three further proton pathways line up provided by the antiporter-type subunits NuoN, NuoM, and NuoL (Fig. 1), which are arranged in an unusual face-to-back manner (23). The name derives from the homology of these subunits to those of multisubunit Na+/H+ antiporters. In complex I they are solely involved in proton translocation. These subunits are homologous to each other (24, 25), with each featuring two pseudosymmetrically arranged half-channels, of which the N terminus is open to the N-side and the C terminus to the P-side of the membrane (23). They comprise a conserved pattern of charged and polar residues that are thought to play an essential role in proton translocation along and across the membrane (23, 26–33). It is assumed that a defined charge distribution in the membrane arm is essential for proton translocation. Each of the antiporter subunits contains a conserved lysine/glutamate or lysine/aspartate ion pair, a conserved central lysine residue, and a distal lysine (NuoL and NuoN) or glutamate (NuoM) residue (Fig. 1). The distal lysine/glutamate residues

![Diagram](https://doi.org/10.1073/pnas.2123090119)
are bridged to the central lysine residue by at least one conserved histidine residue (23). The central lysine residue is proposed to be protonated when the ion pair is closed (9, 26). The closed state describes a strong salt bridge between the residues of the ion pair. It is further assumed that quinone chemistry leads to a proton transfer to NuoK, causing the opening of the ion pair on NuoN (Fig. 1). The central lysine residue then transfers its proton to the distal residue, which then opens the ion pair at the next antiporter subunit (9). Finally, the proton is released from D400L (the superscript refers to the name of the E. coli complex I subunit) to the P-side (Fig. 1). Proton release at D400L triggers the reprotonation of the central lysine residues from the N-side in sequence, coupled with the proton release of the distal residues to the P-side (Fig. 1).

Remarkably, the distribution of the conserved titratable amino acid residues within NuoL, NuoM, and NuoN is asymmetric because NuoL and NuoN include a conserved distal lysine residue (K399 and K395), while NuoM, located between NuoL and NuoN, contains a conserved distal glutamate residue (E407) (23). It was proposed that charge asymmetry within the membrane arm is needed to prevent electrostatic imbalance in the membrane arm and to achieve an asynchronous charge transfer within the antiporter subunits (31). Consequently, a variant with a symmetric charge distribution should not be able to catalyze proton translocation.

To test this hypothesis, we replaced the conserved E407 residue by a lysine residue. The resulting E407K variant was stably assembled and showed a slightly diminished electron transfer activity. However, when reconstituted into liposomes, proton translocation was clearly diminished resulting in an H+/e− stoichiometry of about 1.5, assuming a wild-type stoichiometry of 2 (33−35). Thus, the variant was capable of catalyzing redox-driven proton translocation although less efficient. This finding is consistent with two models of proton translocation and renders the recently proposed “ND5-only” (31) model unlikely, as explained in the Discussion.

**Results**

**Conservation of Amino Acid Residues.** Multiple sequence alignments showed the conservation of a glutamate residue at position 407 (E. coli numbering) within the NuoM homologs and the conservation of a lysine residue at positions 399 (NuoL) and 395 (NuoN) (SI Appendix, Fig. S1A) (23). Sequence comparison further showed that the conserved glutamate residue in NuoM is placed at the homologous position of the conserved lysine residue in NuoL and NuoM (Fig. 1 and SI Appendix, Fig. S1B) (23). It is assumed that each of these residues releases a proton to the P-side in the course of proton translocation (9, 21−23, 26−31). To answer the question of whether E407 is essential for proton translocation, it was replaced by a lysine residue in the sequence of E. coli NuoM.

**Generation and Growth of the E407K Mutant.** The point mutation E407K was introduced into the pBAD-ndh expression plasmid encoding the entire E. coli ndh operon under the control of the inducible PBAD arabinose promoter (36). Possible recombination with the chromosomal wild-type allele during cloning was prevented by using strain DH5αΔndh as cloning host (36). The expression strain BW25113Δndh-λ-red-pBAD/ΔFRT was transformed with expression plasmids either encoding the parental genes or the ones with the mutation. The expression strain lacks the chromosomal gene of the alternative NADH dehydrogenase (ndh) (37) and the chromosomal ndh operon encoding complex I is replaced by a resistance cartridge (nptII) by λ-red−mediated recombination (38). Consequently, NADH-induced activities of membranes exclusively reflect activities of complex I encoded on the plasmid.

The host strain transformed either with the expression plasmid encoding the parental ndh genes or with the one with the E407K mutation was grown in minimal medium with acetate as nonfermentable carbon source. A functionally intact complex I is required for fast growth to high OD values in this medium, because it maintains a low NADH/NAD+ ratio, which is essential to preserve the TCA cycle and contributes to the PMF (39, 40). The strain producing wild-type complex I grew the fastest, while the strain producing the E407K variant grew initially as slow as the host strain that was used as control. However, the mutant strain started to grow faster after ~5 h but still significantly slower than the parental strain (Fig. 2). Thus, the E407K mutation affects complex I activity in the membrane.

![Fig. 2. Growth of the mutant strain and stability of the E407K variant. (A) Growth of the host strain (gray) transformed either with the expression plasmid encoding the parental ndh genes (black) or with the one encoding the E407K mutation (red) in minimal medium with acetate as sole carbon source. The black arrow indicates the induction of gene expression after 3 h growth by an addition of 0.02% 1-arabinose. (B) Sucrose gradient of detergent solubilized membranes from wild-type (black) and the E407K (red) mutant strain. The NADH/ferricyanide oxidoreductase activity of each fraction is shown; the activities are normalized to 20-ng membrane protein extract applied on each gradient.](https://doi.org/10.1073/pnas.2123090119)
Assembly, Stability, and Activity of the Variant in Cytoplasmic Membranes. Cells were disrupted and cytoplasmic membranes were obtained by differential centrifugation. The amount of the complex in the membranes was assessed from the artificial NADH/ferricyanide oxidoreductase activity. This activity is not coupled with Q reduction and proton translocation (41, 42) and it solely depends on the presence of the flavin cofactor, the amount of which is not affected by the mutation. Indeed, the NADH/ferricyanide oxidoreductase activity of the membranes from the mutant strain did not significantly differ from that of the parental strain (Table 1). The physiological activity was determined by measuring NADH oxidase activity. In this assay, NADH is oxidized by complex I in a rate-limiting step and the determination was made by measuring NADH oxidase activity. In this assay, NADH is oxidized by complex I in a rate-limiting step and the amount of which is not affected by the mutation. Indeed, the NADH/ferricyanide oxidoreductase activity of the parental strain (Table 1). The physiological activity was determined by measuring NADH oxidase activity. In this assay, NADH is oxidized by complex I in a rate-limiting step and the amount of which is not affected by the mutation. Indeed, the NADH/ferricyanide oxidoreductase activity of the membranes from the mutant strain did not significantly differ from that of the parental strain (Table 1). The activity curve of the NADH oxidase activity of both strains was linear, demonstrating that substrate binding was not affected by the mutation, as expected (SI Appendix, Fig. S2A). Thus, the E407KM mutation did not influence the amount of complex I in the membrane and slightly diminished its electron transfer activity.

The assembly of the complex in the mutant strain and its stability in the presence of the detergent LMNG (2,2-didecyl-propane-1,3-bis-β-d-maltopyranoside) was investigated by sucrose gradient density centrifugation (Fig. 2B). Cytoplasmic membranes of cells grown in autoinduction-medium were incubated with LMNG and the cleared extract was loaded onto gradients from 5 to 30% (wt/vol) sucrose (43). Solubilized membrane proteins were separated from each other by centrifugation and the gradients were fractionated in 1-mL portions. The position of the complex and its variant in the gradient was determined by its NADH/ferricyanide oxidoreductase activity. The parental complex sedimented around fraction 16 (Fig. 2B). The variant protein sedimented at the same position of the gradient with approximately the same total activity as expected from the NADH/ferricyanide oxidoreductase activity of membranes (Table 1). Thus, the E407KM variant was fully assembled in the membrane and stably extracted with LMNG.

Preparation of Complex I and the E407KM Variant. Cells were grown in autoinduction-medium and cytoplasmic membranes were obtained by differential centrifugation (43). Membrane proteins were extracted with LMNG and the complex and the variant were prepared to homogeneity by affinity and size-exclusion chromatography (SI Appendix, Fig. S3 A and B) (17). From 50-g cells, 7- to 9-mg protein was obtained. The elution profiles of the complex and the variant showed no significant differences. The preparation of the parental complex and that of the variant contained all complex I subunits and the protein band pattern was virtually identical (SI Appendix, Fig. S3C).

Table 1. NADH oxidase and NADH/ferricyanide oxidoreductase activity of cytoplasmic membranes from E. coli wild-type and mutant strain

| NADH oxidase activity | NADH/ferricyanide oxidoreductase activity |
|-----------------------|----------------------------------------|
|                        | U/mg                     | %       | U/mg                     | %       |
| Wild-type              | 0.261 ± 0.020            | 100 ± 9 | 2.8 ± 0.3                | 100 ± 11 |
| E407KM                 | 0.213 ± 0.025            | 83 ± 12 | 2.7 ± 0.3                | 96 ± 11  |

The data were obtained from three measurements of two biological replicates.

Catalytic Activities of the Preparations. The preparations of the E407KM variant had an insignificantly higher specific NADH/ferricyanide oxidoreductase compared to the complex (Table 2). The NADH:decyl-Q oxidoreductase activity was measured in the presence of quinol bo3 oxidase used as a quinone regenerating system, as described previously (17). The kinetic traces showed no lag-phase, indicating an unrestricted substrate binding and product release (SI Appendix, Fig. S2B). The E407KM variant showed a specific activity of 80% of the activity of the parental strain (Table 2). This is in accordance with the NADH oxidase activity of the mutant membranes that was diminished to approximately the same extent (Table 1).

To measure proton translocation, complex I and the E407KM variant were reconstituted into preformed liposomes, as described previously (44). The orientation of the enzymes in the liposomes was determined by measuring the NADH/ferricyanide oxidoreductase activity of intact proteoliposomes and an aliquot that was disintegrated by an addition of 2% detergent. The share of the accessible NADH binding sites was similar in the proteoliposome preparation of the complex and the variant (Table 3). The NADH:decyl-Q oxidoreductase activity of the preparation reconstituted into liposomes was determined by following the decrease of the NADH concentration at 340 nm. Due to the lack of a quinone regenerating system, the activities were lower and amounted to 0.19 (± 0.02) U/mg for the complex and to 0.16 (± 0.04) U/mg for the variant. Thus, the variant showed overall 84% of the electron transfer activity of complex I. Correcting this for the accessible NADH-binding sites in liposomes (Table 3) yields a value of 76%. These proteoliposome preparations were used in two different assays (Fig. 3).

Proton pumping was measured by following fluorescence quenching of ACMA (9-amino-6-chloro-2-methoxyacridine). An addition of NADH led to an acidification of the proteoliposomes lumen due to proton pumping of the incorporated complex and the variant, respectively. However, the rate and the extent of fluorescence quenching of the proteoliposomes containing the E407KM variant was reproducibly lower compared to those containing complex I. The signals obtained with both preparations were sensitive to an addition of gramicidin, demonstrating the presence of a proton gradient. The variant showed 52% of the ACMA quench of the complex (Table 3). An addition of valinomycin accelerated the reaction, but did not alter the relative quench ratio observed with the complex and the variant (SI Appendix, Fig. S4).

The membrane potential was directly determined by following the decrease in the absorbance of the potential sensitive dye oxonol VI (Fig. 3). As with the ACMA measurements, the signal’s intensity and the rate of formation was approximately halved with the proteoliposomes containing the E407KM variant compared to the signal obtained with proteoliposomes containing complex I. Again, the signals obtained with both preparations were fully sensitive to an addition of gramicidin that abolishes the membrane potential. The oxonol absorbance of proteoliposomes containing...
Table 2. NADH:decy1-Q and NADH/ferricyanide oxidoreductase activity of the preparations of complex I and the E407KM variant

|                          | NADH:Q oxidoreductase activity | NADH/ferricyanide oxidoreductase activity |
|--------------------------|--------------------------------|------------------------------------------|
|                          | U/mg                           | %                                        | U/mg                           | %                                        |
| Wild-type                | 32.8 ± 1.9                     | 100 ± 6                                  | 107.0 ± 3.3                    | 100 ± 3                                  |
| E407KM                   | 26.2 ± 2.3                     | 80 ± 9                                   | 114.9 ± 4.7                    | 107 ± 4                                  |

The data were obtained from three measurements of two biological replicates.

The variant was decreased to 54% compared to those containing complex I (Table 3).

The data show that the E407KM variant is capable of catalyzing redox-driven proton-translocation. In proteoliposomes, the electron transfer activity of the variant amounts to 84% of that of the complex. However, the variant liposomes contain a larger share of active enzyme due to its orientation within the membrane and the membrane impermeability of NADH. Taking the different accessibility of the NADH binding sites into account (Table 3), the variant exhibits 76% of the electron transfer activity of the complex in liposomes. Correcting the values of proton translocation activity (Table 3) with the electron transfer activity, the ACMA fluorescence quench of the variant reaches 68% (SD: ± 10.1%) of that of complex I and the oxonol absorbance amounts to 71% (SD: ± 14.8%). As measurements with proteoliposomes tend to underestimate the magnitude of proton translocation due to an inherent leakiness (33), the E407KM variant presumably translocates 75% of the number of protons per electrons compared to complex I. Assuming a stoichiometry of 2 for the latter (33–35), the H⁺/e⁻ stoichiometry of the E407KM variant is diminished to 1.5.

Effect on pKα Values. The altered H⁺/e⁻ stoichiometry of the variant may be due to a change in the local pKα value at position 407M caused by the substitution of the glutamate by a lysine residue. We calculated the pKα value at this position in comparison to that of K265M, the proposed proton donor to E407M using the program PROPKA 3.0 (45, 46). The mutation E407KM was introduced in silico into the structure of the E. coli membrane arm (Protein Data Bank [PDB] ID code 3RKO) (23) followed by energy minimization and H-bond optimization. The pKα of K265M was determined to 6.8 in the original and the variant structure. The pKα value of the original E407M was calculated to be 7.6, which raises the question of whether the membrane arm actually exhibits charge asymmetry, as suggested previously (31), since the lysine residues at the homologous position in NuoN is very similar (K959; pKα = 7.4), while that in NuoL (K999; pKα = 6.2) is more acidic. The latter is most likely due to its close vicinity to D400L, the proposed proton exit. Beyond that, the pKα of the substituted K407M exhibits a pKα of 6.8. Consequently, the mutation shifted the pKα at position 407M by nearly one pH unit, which renders proton transfer from K265M to position 407M more unlikely in the variant. This shift is presumably caused by the overall negative electrostatic surface potential in the environment of the carboxyl group of E407M (Fig. 4). Due to charge repulsion, the protonation of the carboxylate group of E407M is promoted. Modeling the lysine residue into the available space brings the ε-amino group close to a small positively charged surface patch favoring its neutral state (Fig. 4). Proton transfer from K265M to position 407M might be facilitated by H322M (Fig. 4) and water molecules, one of which is found in the structure (22, 23). Remarkably, the E407KM mutation also had a strong impact on H322M and neighboring residues, lowering the pKα by approximately two pH units. This could contribute to a significantly disturbed proton transfer from K265M to K407M and, furthermore, that K407M is less suitable for storing protons than E407M.

Discussion

Although several structures of complex I from various species are available at resolutions ranging from 2.3 to 4.5 Å, the molecular mechanism of proton translocation is still not understood. Proposals have been made to distinguish the deactive from the active state of the complex (31, 47, 48), the binding sites of several Q-site inhibitors have been determined (49–51), and the structure of the complex was obtained in the presence of NADH, quinone, both substrates, and under turnover conditions (31, 52, 53). Several models for proton translocation were proposed mainly based on the combination of structural data and molecular dynamics simulations (9, 20–22, 30–32, 54). However, one has to keep in mind that proton translocation might be accomplished via several short-lived conformational states of higher energy, which might not be detectable by cryoelectron microscopy (cryo-EM). We can also not exclude that the presence of the biological membrane and the PMF are needed to detect such intermediate states. A significant complementary method to confirm or disprove proposals on the enzyme’s mechanism is therefore the kinetic investigation of mutants, in which key amino acids have been exchanged.

Position E407M is such a key amino acid because the conserved glutamate residue is thought to introduce charge asymmetry into the available space bringing the ε-amino group close to a small positively charged surface patch favoring its neutral state (Fig. 4). Proton transfer from K265M to position 407M might be facilitated by H322M (Fig. 4) and water molecules, one of which is found in the structure (22, 23). Remarkably, the E407KM mutation also had a strong impact on H322M and neighboring residues, lowering the pKα by approximately two pH units. This could contribute to a significantly disturbed proton transfer from K265M to K407M and, furthermore, that K407M is less suitable for storing protons than E407M.

Table 3. NADH-induced quench of ACMA fluorescence and decrease in oxonol absorbance due to the activity of complex I and the E407KM variant reconstituted into liposomes

|          | ACMA quench | %      | % of WT | Decrease in oxonol absorbance | ΔA₅88–625 | % of WT | Accessible NADH sites |
|----------|-------------|--------|---------|------------------------------|----------|--------|-----------------------|
| Wild-type| 80.0 ± 3    | 100 ± 4| 100 ± 4 | 0.0304 ± 0.001               | 100 ± 3  | 48 ± 5            |                       |
| E407KM   | 41.2 ± 4    | 52 ± 10|         | 0.0163 ± 0.006               | 54 ± 3   | 53 ± 5            |                       |

The data were obtained from three measurements of two biological replicates.
the membrane arm of complex I as the common charged distal residue is a lysine in NuoL and NuoN, while it is a glutamate residue in NuoM (Fig. 5) (23, 31). To the best of our knowledge, this residue has not been studied by mutational analysis so far. The E407KM mutation should introduce charge symmetry into the membrane arm and should therefore influence proton translocation.

The most recent proposal for proton translocation by complex I is a two-stroke mechanism, with one stroke initiated by the reduction and protonation of Q, leading to proton release from the central hydrophilic axis to the P-side, and a second stroke due to the movement of the QH2 in the cavity, leading to a proton uptake to the central axis from the N-side (31). Both strokes lead to conformational changes that are first transferred to the E-channel and finally lead to a proton transfer within the ion pairs of the antiporter subunits. Protons are proposed to be translocated in a uniform manner across all antiporter subunits by pure electrostatic interactions (31). Due to the change of charges within the K217/E133 ion pair, the charge of the aforementioned conserved amino acids and the bridging histidine residues changes accordingly (Fig. 5). This kind of change continues in the same manner throughout the entire membrane arm. The uneven charge distribution within the antiporter subunits is supposed to prevent an electrostatic imbalance in the membrane arm provided that NuoM operates asynchronously to NuoL and NuoN. The presence of the distal glutamate residue in NuoM was proposed to be causative for the “out-of-sync” operation of the antiporter subunits. According to this model, one proton per antiporter subunit is translocated across the membrane. Hampering proton transfer from NuoM to NuoL by the E407KM mutation would accordingly disconnect NuoL from changes in the charge of the conserved amino acid residues propagating along the membrane arm. Proton release triggered by the first stroke and proton uptake by the second stroke would no longer be transmitted to NuoL, thus diminishing the H+/e− stoichiometry from 2 (complex I) (33–35) to 1.5 (E407KM). This diminished stoichiometry of 1.5 H+/e− is thus in agreement with the measured stoichiometry reported here, although no asynchronous operation of NuoM, a central feature of the proposed mechanism, is needed as the E407KM variant with a symmetric charge distribution within the antiporter subunits is obviously capable of proton pumping.

The same authors also proposed a “ND5-only mechanism” (31). Subunit ND5 is the homolog of NuoL in mitochondrial complex I (Fig. 5). Here, the two-stroke mechanism triggering proton translocation, as described above, applies; but according to this mechanism, all protons are translocated across the membrane only by the most distal subunit NuoL (ND5). This proposal is based on the observation that the E-channel as well as NuoM and NuoN lack proton pathways to the P-side in cryo-EM structures. On the contrary, the distal conserved lysine residue, K399 (K392ND5), was found to be well connected to the P-side by conserved D400L (D393ND5) and by several water...
molecules (Fig. 5). Furthermore, no conformational changes of the antiporter subunits were observed when the sample was frozen for EM analysis in the presence of NADH and decyl-Q. It was concluded that the accessibility of protons from NuoM and NuoN toward the P-side is not altered during turnover. As protons may travel all along the central hydrophilic axis of the membrane arm and because NuoM and NuoN may not have contact to the P-side, it was proposed that all protons are pumped to the P-side by NuoL. Accordingly, four protons are taken up from the N-side, most likely by NuoL and NuoM, and are further distributed along the central hydrophilic axis to the distal NuoL, probably including a temporal storage on the conserved bridging histidine residues (31).

This proposal is in contrast to findings obtained with deletion mutants in *Yarrowia lipolytica* and *E. coli* complex I (55, 56). In *E. coli*, the deletion of *nuoL* led to the assembly of stable complex just lacking NuoL that was capable to pump protons with a $\text{H}^+/\text{e}^-/\text{C}_0$ stoichiometry of about 1 (55). Similar results were obtained with *Y. lipolytica*: deletion of a gene encoding a small accessory subunit led to the formation of a complex that lacked NuoL and NuoM (ND5 and ND4). This complex I variant also showed a reduced stoichiometry of 1 (56). However, it can be argued that deletion of one or two of the antiporter subunits may distort the well-balanced architecture of the membrane arm, leading to the presence of artificial proton pathways that are not existent in the holo-complex.

![Fig. 5. Putative proton pumping mechanisms of complex I. Subunits NuoL (yellow), NuoM (red), NuoN (blue), and NuoK (green) are shown as blocks, the titratable amino acid residues as dots. The name of the individual residues are given in the Top scheme. The ion pairs are shown in black, the opened and closed conformations are indicated by separated and fused dots, respectively. The central lysine residues are shown in blue, the distal lysine residues (NuoL and NuoN) in green and the distal glutamate residue (NuoM) in red. Putative proton pathways are indicated by black arrows, forward (or first) and reverse (or second) power strokes are indicated by orange arrows. (A) Two-stroke mechanism with the propagation of a forward and a reverse electrostatic wave, leading to proton uptake and release in all three antiporter subunits. (B) Asynchronous proton uptake and release by two strokes moving in the same direction. (C) Same as in B, but all protons are released to the P-side by NuoL (ND5-only model). See Discussion for details.](https://doi.org/10.1073/pnas.2123090119)
In contrast, it is highly unlikely that the E407KM point mutation described here will lead to significant structural rearrangements. The calculated pKa shift at positions 407 and H322 induced by the mutation might interrupt the postulated free distribution of protons along the central hydrophilic axis (31). If all protons were released to the P-side by NuoL, the mutation would lead to a complete loss of proton translocation. If, instead, the shift in pKa was not sufficient to interrupt the free distribution of protons along the central axis, the mutation would have no effect on the H+/e− stoichiometry according to the ND5-only mechanism. Remarkably, the mutation diminished the H+/e− stoichiometry, which is inconsistent with the predicted effects. Thus, our results obtained with the E407KM variant argue strongly against the proposed ND5-only mechanism.

Based mainly on computational work, an initial proposal suggests a one-stroke mechanism with each of the antiporter subunits and the E-channel pumping one proton per stroke (9). It assumes a progression of an "electrostatic wave" triggered by the movement of a QH2−species in the Q cavity that leads to the opening of the K217/E133N ion pair (Fig. 5). The protonated central K247N transfers its proton to the distal K395N, which leads to an opening of the lysine/glutamate ion pair in NuoM (E144M/K234M). Accordingly, a "wave" of changes in the electrostatic properties of the titratable amino acid residues propagates across the entire membrane arm. The distal K399N will pass on its proton to conserved D400 that releases this proton to the P-side. This de-protonation induces a "back wave" leading to a protonation of the central lysine residues from the N-side, closing of the lysine/glutamate ion pairs and a proton release from E407K, K395N, and probably from E72K or E36N to the P-side. This proposed mechanism involves the opening and closing of the conserved ion pairs in the antiporter subunits and a concerted opening and closing of the proton pathways leading, on the one hand, from the N-side to the central hydrophilic axis of the membrane arm, and on the other hand, from there to the P-side. As these structural changes have not been observed by cryo-EM, this mechanism was considered unlikely (31). However, it is also possible that these conformational changes represent short-lived conformational states that are thermodynamically unfavorable and might not be detectable by cryo-EM.

According to this mechanism, the de-protonation of the central K265 will close the proton pathway in NuoM to K265M. An alternative explanation is that K407H in NuoM might equilibrate via the open half-channel with K265M. According to this mechanism, the de-protonation of the central lysine residues and a proton release from E407M, K395N, and probably from E72K or E36N to the P-side. This proposed mechanism involves the opening and closing of the conserved ion pairs in the antiporter subunits and a concerted opening and closing of the proton pathways leading, on the one hand, from the N-side to the central hydrophilic axis of the membrane arm, and on the other hand, from there to the P-side. As these structural changes have not been observed by cryo-EM, this mechanism was considered unlikely (31). However, it is also possible that these conformational changes represent short-lived conformational states that are thermodynamically unfavorable and might not be detectable by cryo-EM.

**Materials and Methods**

**Strains, Plasmids, and Oligonucleotides.** A derivative of E. coli strain BW25113 (35) chromosomally lacking the gene ndh was used as host to overproduce complex I (17). In this strain, the chromosomal nuo operon was also replaced by a resistance cartridge (pRPT). E. coli strain DH5α Δnuo was used for site-directed mutagenesis (33). Oligonucleotides were obtained from Sigma-Aldrich (SI Appendix, Table S1). Restriction enzymes were obtained from Fermentas.

Plasmid pBADnuoΔ (57) was used to introduce the respective point mutation on nuoM by site-directed mutagenesis according to the QuickChange protocol (Stratagene). A silent mutation was introduced generating a new restriction site close to the point mutation to identify positive clones by restriction analysis. Primer pair nuoM_E407K (SI Appendix, Table S1) was used to generate the plasmid pBADnuoΔ/eNuoF E407KM. The PCR was performed using the KOD Hot Start DNA Polymerase (Novagen). Mutations were confirmed by DNA sequencing (GATC Eurofins).

**Cell Growth and Preparation of Cytoplasmic Membranes.** Strains were grown aerobically at 37 °C while agitation at 180 rpm. Cells were grown in baffled flasks containing minimal medium with 25 mM acetate as sole carbon source (58). Expression of the nuo operon was induced after 3 h of growth by an addition of 0.02% (wt/vol) arabinose. For protein preparation, cells were grown in a rich autoinduction medium (1% [wt/vol] peptone, 0.5% [wt/vol] yeast extract, 0.4% glycerol, 25 mM Na2HPO4·2H2O, 25 mM KH2PO4, 50 mM NH4Cl, 5 mM Na2SO4, 2 mM MgSO4·7H2O, 0.2% [wt/vol] arabinose, 0.05% [wt/vol] glucose, 30 mg/ml Fe·NH4·citrate, 0.5 mM l-cysteine, 50 mg·L−1 riboflavin) containing chloramphenicol (34 μg/ml) (59). Cells were harvested by centrifugation (5,700 × g, 15 min, 4 °C; Avanti J-26 XP, Beckman Coulter; Rotor JLA 8.100) in the exponential phase yielding between 6 and 7 g cells/L. All further steps were carried out at 4 °C. Sedimented cells were suspended in a fivefold volume of buffer A (50 mM MES/NaOH, 50 mM NaCl, pH 6.0) containing 0.1 mM PMSF and a few grains of DNaseI. They were disrupted by three passages through an 0.1 mm stainless steel French pressure cell. The supernatant (0.15 mL) was transferred to a 1-mL ultracentrifugation tube (16,000 × g, 20 min, 4 °C) and centrifuged (12,000 × g, 20 min, 4 °C; Rotor JA25.50, Beckman; Avanti J-26 XP, Beckman Coulter). Cytoplasmic membranes were obtained from the supernatant by ultracentrifugation (160,000 × g, 70 min, 4 °C; L8-M Ultrafuge, Beckman; Rotor 60 Ti). Sedimented membranes were suspended in an equal volume (1:1 [wt/vol]) of buffer A* (buffer A with 5 mM MgCl2 containing 0.1 mM PMSF).

**Electron Transfer Assays.** Activity assays were performed at 30 °C. The NADH oxidase activity of cytoplasmic membranes was determined by a Clarke-type oxygen electrode (DWO, Hansatech) monitoring the decrease in oxygen concentration in the buffer. The electrode was calibrated by an addition of a few grains sodium dithionite to air saturated buffer. The difference in current before and after reduction was attributed to 237 mM oxygen (60). The assay contained 5 μL cytoplasmic membranes (80 to 90 mg/ml) in 2 μL buffer A*. After equilibration, the reaction was started by an addition of 1.25 mM NADH. The NADH:ferriyancide oxidoreductase activity was determined as decrease in the absorbance of ferriyancide at 410 nm with a diode-array spectrometer (QS cuvette, 1 cm, Hellma; TIDAS II, J&M Aalen). The assay contained 2 μL membrane suspension or 0.1 μL complex I and 1 μM K2[Fe(CN)6] in buffer A*. The reaction was started by an addition of NADH (0.2 mM, final concentration). The NADH:decylic-O oxidoreductase activity was measured as decrease of the NADH concentration at 340 nm using an ε of 6.3 × 103 cm−1 mol−1 (QS cuvette, d = 1 cm, Hellma; TIDAS II, J&M Aalen). The assay contained 2 μL membrane suspension or 0.1 μL complex I and 1 μM K2[Fe(CN)6] in buffer A*. The reaction was started by an addition of 150 μM NADH (62).
added to the membrane suspension (∼70 mg/mL) in buffer A\textsubscript{vNADH}. After 1-h incubation at room temperature with gentle stirring, the suspension was centrifuged for 20 min at 160,000 × g at 4 °C (Rotor 60Ti, L8-M Ultrafuge, Beckman). The supernatant was filtered (Filtropur 50.45; Sarstedt), diluted to 150 mL, and adjusted to 20 mM imidazole, and applied to a 35-mL ProBond Ni\textsuperscript{2+} IDA column (Invitrogen) equilibrated in binding buffer (A\textsubscript{vNi2} with 20 mM imidazole, pH 6.8). The column was washed with binding buffer and with binding buffer containing 160 mM imidazole until the absorbance at 280 nm dropped below 500 mAU. Bound proteins were eluted with binding buffer containing 308 mM imidazole. Fractions containing NADH/ferriyanide oxidoreductase activity were pooled and concentrated by ultrafiltration in 10 kDa MWCO Amicon Ultra-15 centrifugal filter (Millipore) (43). The concentrate was applied onto a Superose 6 size-exclusion chromatography column (300 mL; GE Healthcare) equilibrated in buffer A\textsubscript{vNi2}. The fractions with highest NADH/ferriyanide oxidoreductase activity were pooled and concentrated as described above. The protein was either directly used or stored at −80 °C.

Preparation of Liposomes and Measurement of Gradients. E. coli polar lipids (25 mg/mL in CHCl\textsubscript{3}, Avanti) were evaporated and dissolved in five times the volume of lipid buffer (5 mM MES/NaOH, 50 mM NaCl, pH 6.7). The suspension was frozen in liquid nitrogen and thawed at 29 °C seven times to get unilamellar vesicles (44). The liposomes were extruded by at least 21 passes through an extruder (0.1 μm polycarbonate membrane; Mini Extruider, Avanti). For reconstitution, ~0.5 mg complex I was mixed with reconstitution buffer (1:3 [vol/vol]) (10 mM Bis-Tris-propane/MES, 100 mM KCl, 73 mM sucrose, 2.5 mM MgSO\textsubscript{4}, 0.05% [wt/vol] α-phosphatidylcholine, 1.1% [wt/vol] n-octyl-glucoside, 0.6% [wt/vol] sodium deoxycholate, 0.6% [wt/vol] sodium cholate, pH 7.5) and incubated for 5 min on ice. Next, 250 μL liposomes were mixed with 8 μL sodium cholate (20% [wt/vol]) and the liposomes and complex I in reconstitution buffer were mixed and incubated for 20 min at room temperature with occasional flicking of the tube. Liposomes were formed using a size-exclusion column (PD-10 Desalting Column, 8.3 mL, Sephadex G-25, GE Healthcare) equilibrated in lipid buffer to remove excess detergent. The eluate (1.2 mL) was centrifuged (4 °C, 200,000 × g, 30 min; Rotor, A-100, Airfuge, Beckman) and sedimanted proteoliposomes were gosenuspendes in 500 μL lipid buffer. They were used immediately in subsequent assays at 30 °C.

The generation of ΔpH was determined by monitoring the fluorescence quenching of the pH-sensitive dye ACMA (Sigma). The assay contained 100 μM decyl-Q (Sigma), 0.2 μM ACMA, and 50 μL proteoliposomes in ACMA buffer (10 mM Bis-Tris-propane/MES, pH 6.75, 100 mM KCl, and 2 mM MgCl\textsubscript{2}). The reaction was started by an addition of 130 μM NADH. Fluorescence was detected with a LS 55 Fluorescence Spectrometer (Perkin-Elmer) using excitation and emission wavelengths of 430 nm and 480 nm, respectively. An addition of 1 μg gramicidin (Sigma) was used to dissipate the ΔpH.

The generation of ΔΨ was determined by monitoring the changes in absorption of the potential-sensitive dye oxonol VI (Sigma). The assay contained 0.5 μM oxonol VI, 50 μM Q\textsubscript{10} and proteoliposomes in oxonol buffer (10 mM MES/KOH, pH 6.75, 2 mM MgSO\textsubscript{4}, 100 mM KCl, 10 mM NaN\textsubscript{3}). The reaction was started by an addition of 100 μM NADH. The absorbance difference at 588 minus 625 nm was measured with a diode-array spectrometer (QS cuvette, d = 1 cm, Hellma; TIDAS II, J&M Aalen). An addition of 1 μg gramicidin was used to dissipate ΔΨ.

In Silico Modeling of E. coli Complex I. A structural model of the holo-
complex I from E. coli was generated using AlphaFold 2.1.1 (63) in mutimer mode (64). The complex was modeled in two separate segments, one compris-
ing the membrane arm and the basal peripheral stem, with a total of 10 sepa-
rate chains [i.e., NuoAB(CD)HJKL]. In a separate model, we generated the peripheral arm in an assembly of six subunits, NuoB(CD)EFGI. The full complex was then assembled with the overlapping subunits as a guide, avoiding the inclusion of distortions of subunits that were created by the absence of adjacent protomers. Following the assembly of the modeled protein subunits, the iron-sulfur clusters and FMN cofactor of the peripheral arm were modeled by hand and the entire complex was geometry-optimized using the Maestro Suite (Schrödinger).

Other Analytical Methods. Protein concentration was determined according to the biuret method using BSA as a standard (65). The concentration of purified complex I was determined by the difference of absorbance at 280 to 310 nm (TIDAS II, J&M Aalen) using an ε of 781 mM\textsuperscript{−1} cm\textsuperscript{−1} as derived from the amino acid sequence (66). SDS/PAGE was performed according to Schägger and von Jagow (67) with a 10% separating gel and a 3.9% stacking gel. Multiple sequence alignments were performed using Clustal X 2.1 (68). Default values were used for the penalties for opening and extending gaps.

Data Availability. All study data are included in the main text and/or SI Appendix. Genetic material can be made available upon reasonable request.

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