Energetics and Dynamics of Proton-Coupled Electron Transfer in the NADH/FMN Site of Respiratory Complex I

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ABSTRACT: Complex I functions as an initial electron acceptor in aerobic respiratory chains that reduces quinone and pumps protons across a biological membrane. This remarkable charge transfer process extends ca. 300 Å and it is initiated by a poorly understood proton-coupled electron transfer (PCET) reaction between nicotinamide adenine dinucleotide (NADH) and a protein-bound flavin (FMN) cofactor. We combine here large-scale density functional theory calculations and quantum/classical models with atomistic molecular dynamics simulations to probe the energetics and dynamics of the NADH-driven PCET reaction in complex I. We find that the reaction takes place by concerted hydrogen atom (H⁺) transfer that couples to an electron transfer (eT) between the aromatic ring systems of the cofactors and further triggers reduction of the nearby FeS centers. In bacterial, Escherichia coli-like complex I isoforms, reduction of the N1a FeS center increases the binding affinity of the oxidized NAD⁺ that prevents the nucleotide from leaving prematurely. This electrostatic trapping could provide a protective gating mechanism against reactive oxygen species formation. We also find that proton transfer from the transient FMNH⁺ to a nearby conserved glutamate (Glu97) residue favors eT from N1a onward along the FeS chain and modulates the binding of a new NADH molecule. The PCET in complex I isoforms with low-potential N1a centers is also discussed. On the basis of our combined results, we propose a putative mechanistic model for the NADH-driven proton/electron-transfer reaction in complex I.

INTRODUCTION

Complex I (NADH:ubiquinone oxidoreductase) functions as the electron entry point in aerobic respiratory chains. It catalyzes electron transfer (eT) between nicotinamide adenine dinucleotide (NADH) and quinone (Q), which the enzyme employs to pump protons across a biological membrane.1-5 This generates a proton motive force (pmf) that thermodynamically drives synthesis of adenosine triphosphate (ATP) by F₁Fₐ-ATPase and active transport.6-2 The eT process is fully reversible, and under high pmf conditions, complex I catalyzes the reverse eT from quinol (QH₂) to NAD⁺.8,9 Such operation modes are physiologically relevant under hypoxic conditions, where electrons leak out from complex I.10 The eT in complex I takes place in the hydrophilic domain, which extends ca. 100 Å above the membrane plane, whereas the proton pumping machinery comprises the ca. 200 Å wide membrane domain. It currently remains unclear how these two processes are coupled across such large distances, but mechanistic models have recently started to emerge (see ref 5 and refs therein).

The NADH binding site is located in a solvent-accessible cavity of the Nqo1 subunit (Thermus thermophilus nomenclature) at the “upper edge” of the hydrophilic domain (Figure 1).11,12 This cavity contains a flavin mononucleotide (FMN) cofactor that is surrounded by an unusual Rossmann fold with four parallel β-strands in contrast to the six strands typically found in such motifs. This modification could have evolved to allow for binding of the two nucleotides and to ensure a tight packing of the aromatic ring systems to facilitate the charge transfer between NADH and FMN that is supported by several conserved residues in the cavity.13

The first step in complex I catalysis is the oxidation of NADH at a high rate of ca. 600–15 000 s⁻¹,14,15 which involves transfer of two electrons via a proton-coupled electron transfer (PCET) process to the FMN, most likely forming FMNH⁻ or FMNH₂, whereas no stable flavosemiquinone radical (FMNH⁺) species has so far been detected.16 The process effectively involves a hydride ion (H⁺ = H⁺/2e⁻) transfer, although the exact reaction intermediates remain unknown. The reaction is thus chemically similar to many organic and enzyme-catalyzed PCET reactions.17-20

After the flavin reduction, the two electrons enter the FeS chain to stepwise reduce Q to QH₂ that links to local proton transfer from His38 and Tyr87 of the Nqo4 subunit.21,22 The FeS centers are one-electron carriers in complex I,23 and the NADH/FMN site has been suggested as an electron coupling site24,25,26 that bifurcates the eT process in the FeS chain.25 To this end, N1a and N3, which are located at 6–10 Å from the NADH/FMN site, function as the initial electron acceptors (Figure 1A, left inset). The N3 center mediates the eT across the ca. 100 Å FeS wire that connects FMN with Q (Figure 1A, right inset).

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The exact role of N1a, which is located on the side of the main transfer pathway, remains unclear. The N1a redox potential is unknown for the mammalian complex I, but a redox potential as low as −400 mV has been indirectly suggested for some species,24,25 relative to the redox potential of the other FeS centers at ca. −320 mV and the terminal N2 center at ca. −200 mV.26,27 In contrast, reduction of N1a in E. coli complex I, with an \( E_{\text{m}} \) of −240 mV, has been observed in EPR experiments.

N1a could have a protective role against formation of reactive oxygen species (ROS) that is linked to development of mitochondrial disorders,51 by modulating the dissociation of NAD\(^+\) from the FMN site.31 Interestingly, the relative \( K_m \) of NADH and NAD\(^+\) depends on the direction of the e\(^{-}\)T process: the \( K_m \) of (NAD\(^+\)) is 2 orders of magnitude lower than the \( K_m \) of (NADH) in the 1 mM range during forward e\(^{-}\)T, whereas under reverse e\(^{-}\)T conditions from QH\(_2\) to NAD\(^+\), \( K_m \) of (NADH) is in the 10–20 \( \mu \)M range.31,32,33 Moreover, the oxidized NAD\(^+\) is released on the millisecond time scale.15

To probe the energetics and mechanism of the initial PCET process between NADH and FMN, and its coupling to subsequent e\(^{-}\)T reactions, we combine here large-scale density functional theory (DFT) calculations with classical molecular dynamics (MD) simulations and hybrid quantum mechanics/molecular mechanics (QM/MM) calculations. We propose a putative mechanism for the PCET reaction and discuss its possible implications in the prevention of ROS formation.

**RESULTS AND DISCUSSION**

**PCET Energetics between NADH/FMN and the Nearby FeS Centers.** In order to study the energetics of the PCET reaction between NADH and FMN, we built DFT and QM/MM models of the NADH/FMN site based on the crystal structure of complex I from *T. thermophilus*, shown in Figure 2A (see Methods). The stacked aromatic rings of NADH and FMN in the optimized reactant geometry are at a ca. 3.5 Å distance from each other, with the transferring hydrogen at ca. 2.1 Å from the acceptor nitrogen. The resulting energy profile for the PCET process is shown in Figure 2B.

We obtain a free energy barrier of ca. 12 kcal mol\(^{-1}\) for the PCET reaction, which is exergonic by ca. −6 kcal mol\(^{-1}\). The barrier is consistent with the experimental \( k_{\text{cat}} \) of ca. 600–15 000 s\(^{-1}\)14,15 which corresponds to a barrier of ca. 12–14 kcal mol\(^{-1}\) if a standard pre-exponential factor \( k_0 \) of 6.45 ps\(^{-1}\) and a transmission coefficient of \( \kappa = 1 \) are used in transition state theory. We find that the zero-point energy (ZPE) and entropic corrections at \( T = 310 \) K of the model affect the reaction barrier and energetics by <1 kcal mol\(^{-1}\).

The transferring hydrogen has at the transition state an imaginary frequency of ca. 1380 cm\(^{-1}\), which corresponds to a barrier of ca. 12–14 kcal mol\(^{-1}\) if a standard pre-exponential factor \( k_0 \) of 6.45 ps\(^{-1}\) and a transmission coefficient of \( \kappa = 1 \) are used in transition state theory. We find that the zero-point energy (ZPE) and entropic corrections at \( T = 310 \) K of the model affect the reaction barrier and energetics by <1 kcal mol\(^{-1}\).

In order to probe how the protein environment influences the PCET energetics, we also studied the reaction by hybrid QM/MM calculations. Consistent with the DFT models, we obtain a reaction barrier of ca. 17 kcal mol\(^{-1}\) and a driving force of ca. −4 kcal mol\(^{-1}\) (Figure 2B). The differences between the DFT and QM/MM models arise from small polarization and structural changes in the surroundings (Figure S15, SI).

Initially, the NADH and FMN headgroups are neutral, and during the PCET reaction, a net negative charge is transferred to the flavin cofactor (Figure 2B). The transferring proton carries spin but a nearly neutral charge, suggesting that an electron and a proton are transferred concertedly as a hydrogen atom, H*, while the second electron moves along the aromatic ring system of the nucleotides (Figures 2A,B and S4 (SI)). We observe a similar H* transfer process by employing different density functionals, suggesting that the obtained results do not arise from artifacts due to intrinsic DFT errors. This chemical picture is further supported by analysis of frontier orbitals (see Figures S3 and S5–S7 and Movie S1 and Movie S2, SI).

The resulting FMNH\(^+\) increases the proton affinity of the flavin, which could lead to formation of FMNH\(_2\) (cf. refs 38 and 39), for example, via Glu97. However, due to sterical
clashes, this process requires NAD\(^+\) to dissociate away from the binding site, which is expected to take place on the millisecond time scale.\(^{15}\) The FMNH\(^2\) state was therefore not considered further (but see Supporting Information).

We next probed the proton transfer energetics after one-electron oxidation of the NADH/FMN pair (Figure 2C), a state that could arise after initial e\(^T\) to N1\(a\) or N3 (see below). This reaction also takes place by H\(^+\) transfer but with a slightly higher reaction barrier of ca. 14 kcal mol\(^{-1}\) and an exergonicity of ca. –30 kcal mol\(^{-1}\). Our QM/MM calculations show a similar energy barrier of ca. 16 kcal mol\(^{-1}\), but a somewhat smaller driving force of ca. –22 kcal mol\(^{-1}\), which is overall similar to the DFT results.

On the basis of these observations, we cannot rule out that the second electron is transferred concertedly to N1\(a\) or N3 with the H\(^+\) transfer step. We therefore constructed combined models of the NADH/FMN and N1\(a\)/N3 subsystems (Figure 3). In the combined DFT models, we also obtain a reaction barrier of ca. 12 kcal mol\(^{-1}\) for the H\(^+\) transfer with driving forces of ca. –8 and –20 kcal mol\(^{-1}\) for the e\(^T\) to N1\(a\) and N3,
FMN–N1a and FMN–N3 distances are 11.3 and 6.4 Å (Figure 1), suggesting that the electronic couplings are strong between the NADH/N1a and FMN/N3 eT couples. On the basis of the computed eT parameters, a generic protein-packing density ($\rho = 0.76$), and dynamically averaged edge-to-edge distances between the cofactors (see SI and Figure S9), we obtain effective eT rates in the 1–10 ns range between the NADH/FMN and N1a/N3 couples, further supporting that both eT processes are fast and possibly competing in the bacterial complex I isoforms considered here.

Conformational Dynamics Coupled to the eT Process. To study how the PCET process is linked to binding of the NADH/NAD$^+$ cofactor, we performed classical MD simulations of the substrate-bound complex I in different redox states. In simulations of the NADH/FMN state with oxidized N1a and N3, the NADH remains tightly stacked with FMN at ca. 3.6 Å distance, similar to that in the DFT models and the crystal structure [Figures 2 and S10 (SI)]. After formation of the NAD$^+/FMNH^+$ state, the nucleotides also remain in close contact, but at a slightly higher distance of ca. 4.7 Å, most likely due to the electrostatically dominant interaction (Figure 4A).

We next probed the dynamics of the NAD$^+/FMNH^+$ state, which is expected to form after eT from NAD$^+/FMNH^+$ to one of the nearby FeS centers (see above). With both N1a and N3 oxidized, a state that could arise if the first electron is transferred via N3 to N2, the electrostatic interaction between the cofactors is lost, and NAD$^+$ rapidly dissociates from FMNH*. NAD$^+$ does not, however, fully leave the binding site but remains bound to conserved aromatic residues on the Nqo1 surface [Figures 4C and S1 (SI)]. We also observe similar dissociation dynamics with the N3 center modeled in the reduced state (Figure 4B), in which NAD$^+$ is fully released to the solvent. However, the lifetime of this state is expected to be transient due to rapid eT from N3 toward the N2 center, but it may increase upon steady-state turnover conditions when the FeS is partially reduced.$^{9,29,41}$ FMN(H)$^*$ can also form during reverse eT from Q to FMN. After NAD$^+$ dissociation, our MD simulations suggest that the flavin is accessible to water molecules (Figure S14, SI), and may thus provide a possible pathway for ROS production.

Our calculations suggest that the dynamics of the nearby charged residues may strongly modulate the energetics of the eT to the FeS centers. Arg104 and Arg135 of Nqo1, and Lys121 of Nqo2 move closer to N1a upon its reduction $^{42}$ during the 100 ns simulation upon reduction of N1a (Figure 4B). In these simulations, Glu97 moves away from N1a and forms an interaction with NAD$^+$/FMNH$. Arg104 and Arg135 of Nqo1 are fully conserved, whereas Lys121 of Nqo2 does not (Figures S1 and S2, SI), which could explain the suggested high variation in the N1a redox potential among the bacterial complex I isoforms, despite the latter process being thermodynamically preferred in the structural models.

eT rates also depend on the electronic coupling, which has an exponential dependence on the donor and acceptor distance separation, exp$(-\beta_0 R_{DA})$ (see the Supporting Information). We note that the edge-to-edge distances between NADH–N1a and NADH–N3 are 10.6 and 12.2 Å, respectively, whereas the
against formation of reactive oxygen species that could leak out from FMN.9,29,43

**Energetics of the Second eT Process.** NAD⁺ dissociation decreases the electron affinity of FMNH⁺ that in turn favors the transfer of the second electron to N3 by ca. −3 kcal mol⁻¹ (130 mV) in our DFT models. Experimentally, no flavosemiquinone species has been detected, and we thus expect that the second electron rapidly enters the FeS chain upon NAD⁺ dissociation. We observe that the oxidation of FMNH⁺ couples to proton transfer to the conserved Glu97 of Nqo1. After protonation of Glu97, the backward eT from N1a to FMN becomes endergonic by ca. −3 kcal mol⁻¹ (130 mV) in the DFT models, whereas we are not able to oxidize N1a with a singly reduced FMN and a deprotonated Glu97, suggesting that the state has a significantly higher energy. The subsequent reduction of N3 by FMN with a protonated Glu97 has a strong driving force of ca. −10 kcal mol⁻¹ (430 mV) in the DFT models. This suggests that deprotonation of FMN by Glu97 could allow for the eT from N1a back to FMN and further to N3, providing a pathway for the second eT toward the N2 center (Figure S12, SI).

When Glu97 is modeled in its protonated state, the barrier for the PCET reaction between NADH/FMN increases by ca. 7 kcal mol⁻¹ (Figure S11, SI), consistent with the experimentally observed decreased NADH-oxidation activity in the E97Q mutant.13 Interestingly, the Glu97 protonation alters the conformation of the active site, preventing the entry of a new NADH molecule (Figure 5C), by forming a hydrogen bond with the backbone of Gly67 of a Gly-rich loop (residues 66-69) of the Rossmann fold (Figure 5D) that blocks the exposed FMN face where NADH binds. The proton release from Glu97 to the solvent could therefore favor the initiation of the next catalytic cycle and also provide a mechanism to prevent over-reduction of the FeS chain.

Substitution of Glu97 by a neutral glutamine upshifts the redox potential of the FMN by ca. 40 mV,44 and mutation of its hydrogen-bonding partner, Tyr180, is linked with development of Leigh syndrome,45 a neurometabolic disorder with increased ROS production. In the light of our findings, we suggest that Glu97 has an additional role in modulating NADH binding/NAD⁺ release and the transfer of the second electron from N1a further to the FeS chain.

**PCET Energetics in Different Complex I Isoforms.** To study the PCET reaction energetics and coupled eT in other complex I isoforms, we built DFT models of both E. coli and mouse N1a centers with high- and low-potential forms of the cluster,24 respectively (see the SI). Our combined models suggest that eT from NAD⁺/FMNH⁺ to E. coli N1a has similar energetics as in our T. thermophilus models (Figure S13, SI), although the electron affinity is around 80 mV higher in the former model. These findings are consistent with the suggested electron bifurcation to N1a and N3 observed in the E. coli complex I.15 In contrast, the electron affinity in the mammalian N1a center is low, and we do not observe eT to N1a in the models (Figure S13 and Table S3, SI).
Interestingly, experiments suggest that the redox potential of N1a could affect whether the reduced flavin leads to production of H2O2 or superoxide. The bovine complex I with a low-potential N1a mainly produces superoxide, whereas the *E. coli* complex I with a high potential N1a results in H2O2 production. Birrell et al. did not observe changes in the superoxide/H2O2 ratio upon introduction of redox-tuning point mutations near the N1a cluster in the 24 kDa subunit of the *E. coli* complex I, whereas Gnad et al. observed a decreased ROS production upon N1a reduction. The molecular basis of these different conditions and isoforms are not fully understood, but taken together, they suggest that N1a reduction could prevent ROS formation, e.g., by increasing NAD+ binding affinity, at least in *E. coli* complex I (see above). The different complex I isoforms have indeed a variation in the Kd of NAD+/NAD+ which in turn could arise from subtle structural differences in the different complex I isoforms and/or from differences in the flavin redox state.

### PUTATIVE MECHANISTIC MODEL

On the basis of our quantum and classical molecular simulations, we propose a mechanistic model for the initial NADH-driven PCET reaction in different complex I isoforms (Figure 6). The PCET reactions are initiated by hydrogen atom transfer that couples to an eT between the NADH/FMN ring systems, forming NAD+ and FMNH+. The reaction is exergonic, and it thermodynamically drives the semiconcerted eT from FMNH+ to N1a in *E. coli*-like complex I isoforms, which kinetically competes with reduction of N3 (see below for discussion on low-potential N1a centers). The N1a reduction prevents NAD+ from dissociating prematurely from the binding cavity that is favored by formation of FMNH+ due to loss of the electrostatic interaction between NAD+ and FMNH+. The electrostatic trapping of NAD+ could be important during reverse eT from quinol toward N3 to prevent electrons from leaking from FMN to the solvent.

The second eT from FMNH+ to N3 couples to a proton transfer from the flavin to the conserved Glu97. This protonation helps to release the electron stored at N1a to FMN, and further along N3 to the FeS chain. Proton release from Glu97 to the solvent facilitates NADH binding due to conformational changes, favoring complex I to enter the next catalytic cycle. This protonation-triggered “gate” could prevent over-reduction of the FeS chain, which would also be expected to have a pH-dependence.

The transfer of the two electrons to the nearby FeS centers has been observed in the microseconds time scale in *E. coli*, whereas NAD+ dissociation is reported in the slower milliseconds time scales. We note that NAD+ dissociation favors the release of the second electron that could be triggered by a partial NAD+ dissociation from the flavin to the nearby protein surface, as observed in the classical MD simulations (Figure 4).

For complex I isoforms with low-potential N1a centers (see above), the first electron is expected to be transferred to N3 and further toward the N2 center. The resulting FMNH+ lowers the affinity for NAD+, which favors dissociation of NAD+ to the bulk. Although this also lowers the electron affinity of the flavin and favors release of the second electron, it could lead to formation of a transient dangerous intermediate that may leak electrons out to the solvent. As discussed above,
ROS is produced differently in mammalian and E. coli complex I.\textsuperscript{34,47}

During forward eT to Q, this flavo-semiquinone radical state would be transient due to the fast eT to N3. However, under reverse eT conditions with a highly reduced FeS chain, the occupation of this state is expected to increase, which in turn would lead to increased ROS generation.\textsuperscript{9,29,41} As discussed above, N1a reduction could prevent ROS formation by trapping NAD$^+$ to the binding site under such conditions in E. coli complex I. However, we also note that the reduced N1a decreases the electron affinity of the FMN by 0.3 eV in our DFT models (Table S4, SI) that could further protect against formation of this potentially dangerous state. Complex I variants with a low N1a redox potential may employ, for example, structural changes linked to the \textit{active-to-deactive} transition\textsuperscript{39–41} to modulate the eT kinetics and prevent ROS formation.

## CONCLUDING REMARKS

In this work, we proposed a mechanistic model for the initial PCET reactions in the NADH/FMN site of respiratory complex I. Our large-scale DFT and QM/MM models that focus here on bacterial complex I isoforms suggest that the PCET between NADH and FMN involves an exergonic hydrogen atom (H$^+$) transfer that couples to eT between the aromatic rings of the cofactors. In our models, the two electrons further bifurcate in the subsequent eT steps to the nearby FeS centers, N1a and N3, consistent with EPR experiments\textsuperscript{15} and with mechanistic analog to other flavoenzymes.\textsuperscript{52} The N1a reduction increases the binding affinity of NAD$^+$ that prevents the electrons from leaking out to the solvent. Proton transfer between the transiently formed FMNH$^+$ species and a conserved Glu97 was found to facilitate the release of the second electron from the flavin and to prevent a new NADH from entering the binding site by triggering conformational changes in the cavity. We suggest that this proton transfer process is also involved in releasing the electron stored on the N1a center via FMN back to the FeS chain. These findings also imply that the forward and backward eT reactions could be modulated by the external pH.

Our calculations showed that the protein dynamics may strongly influence the energetics of the eT between NADH/FMN and the N1a/N3 centers. Conformational changes in highly conserved charged residues of Nqo1 that move closer to the FeS centers facilitate their reduction, whereas the high variability of charged residues in the Nqo2 subunit, as also observed from large variations in the N1a redox potential in different species, i.e., from −240 mV in \textit{E. coli} to suggested values of ca. −400 mV in mammalian complex I,\textsuperscript{15} is consistent with our calculations on different complex I isoforms. In DFT models of low-potential N1a isoforms, we do not observe reduction of N1a, suggesting that eT follows a different reaction pathway, as highlighted in Figure 6. Modulation of the eT kinetics might couple to conformational changes, which are currently not well understood. Redox-triggered conformational changes have not yet been observed in available experimental structures, possibly due to their 3–4 Å resolution. Taken together, our combined results suggest that the long-range electron transfer process in complex I is initiated by coupled protonation and reduction steps in the NADH/FMN site that trigger electrostatic and conformational changes in the protein environment to prevent electron leaks and over-reduction of the FeS chain. The model has important mechanistic implications for the forward and reverse electron transfer process in complex I and for understanding the subsequent quinone reduction process that, in turn, activates the proton pumping machinery.

## METHODOLOGY

DFT models for the NADH/FMN, N1a, and N3 subsystems comprising first- and second sphere protein residues were built on the basis of the crystal structure of complex I from \textit{T. thermophilus} (PDB ID: 3IAM; see the Supporting Information).\textsuperscript{11} Additional N1a models from \textit{E. coli} and mammalian complex I were also built (see the SI). The individual subsystems comprised 150–170 atoms and the combined systems ca. 330–350 atoms. Spin energetics were treated using the spin-flip broken symmetry approach (Table S2, SI),\textsuperscript{51} and geometry optimizations were performed at the B3LYP-D3/def2-SVP/def2-TZVP(Fe)/ε=4 level, with single-point energies reported at the B3LYP-D3/def2-TZVP/ε=4 level.\textsuperscript{52–55} The electronic structures of the combined models were converged into localized “diabetic” electronic states by merging the molecular orbitals of the electron donor (NADH/FMN) or acceptor sites (N1a or N3) in their respective reduced/oxidized states together. Zero-point energies (ZPE) and entropic corrections were estimated at the B3LYP-D3/def2-SVP/ε=4 level using smaller model systems. Reaction pathways for the PCET reaction were optimized using a \textit{chain-of-state} method,\textsuperscript{60} and hydrogen/deuterium (H/D) kinetic isotope effects (KIE) were estimated on the basis of the ZPE differences between reactant and transition state structures. Approximate quantum effects were evaluated by the Wigner correction,\textsuperscript{61} and electron transfer rates were based on the Moser–Dutton rule.\textsuperscript{62} All QM calculations were performed with TURBOMOLE versions 6.6–7.2.\textsuperscript{63}

QM/MM models of the NADH/FMN subsystem with 132 QM atoms were constructed for computing the potential energy surface (PES) for the PCET reaction between NADH and FMN. The QM subsystem was treated at the B3LYP-D3/def2-SVP level, and the MM subsystem was described at the CHARMM36 level based on the classical model (see below).\textsuperscript{64} The reaction coordinate, R = r1 − r2, where r1 = (NADH:C4−H) and r2 = (FMN: N5−H), was optimized from reactants, R = −1.5 Å, to products, R = +2.5 Å, using a harmonic potential restrain. During optimization, the QM region and the MM surroundings within a 15 Å sphere centered on FMN were allowed to relax. All QM/MM calculations were performed with the CHARMM/TURBOMOLE interface.\textsuperscript{65}

Classical atomistic molecular dynamics (MD) simulations were performed on the hydrophobic domain of complex I by including subunits Nqo1 with the NADH/FMN cofactors, Nqo2, Nqo3, and Nqo15. The protein was solvated and neutralized in a TIP3P water/100 mM NaCl solution box and comprised ca. 170 000 atoms. MD simulations were performed at p = 1 bar and T = 310 K in an NPT ensemble, using a 2 fs integration time step and treating long-range electrostatic interactions by the Particle Mesh Ewald (PME) method. The CHARMM36 force field\textsuperscript{66} and our in-house DFT-derived parameters for N1a and N3 in both oxidized and reduced states were used. The MD simulations were performed with NAMD\textsuperscript{67} and VMD was used for analysis.\textsuperscript{68} All simulations are summarized in Table S1 (SI). See the Supporting Information for a detailed computational description.

## ASSOCIATED CONTENT

\* Supporting Information

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Detailed computational methods; figures showing the multiple sequence alignment of Nqo1/NuoF/NDUVF1 and Nqo2/NuoE/NDUFV2 subunits of complex I, energetic of the NADH/FMN PCET process, radial distribution of the spin density on the transferred hydrogen atom, spin density on the transition state...
structures, spin density and HOMO orbitals of the reactant, the transition state, and the product structure, distances for N1a in Nqo1 and Nqo2 subunits, edge-to-edge distances between NADH–N1a/N3 and FMN–N1a/N3 during MD simulations, dissociation energy profiles, energetics of the PCET reactions, the energetics of the second electron transfer from the flavin to the FeS centers from DFT models, energetics of the PCET reaction between NADH/FMN and N1a in complex I, the FMN binding cavity, comparison of the energetics of the NADH/FMN PCET reaction between QM and QM/MM levels; tables summarizing the simulations, optimized broken-symmetry spin-flip states for the oxidized and reduced FeS centers, electron affinities of the N1a FeS in DFT models, electron affinities of FMN with oxidized and reduced N1a; and captions describing Movies S1–S4 (PDF)

Movie S1: Spin density in the PCET reaction between NADH and FMN (MPG)

Movie S2: Highest occupied molecular orbital (HOMO) in the PCET reaction between NADH and FMN (MPG)

Movie S3: Spin density in the PCET reaction between NADH and FMN that couples eT to N1a (MPG)

Movie S4: Spin density in the PCET reaction between NADH and FMN that couples eT to N3 (MPG)

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**Notes**

The authors declare no competing financial interest.

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