

Site-directed nanoparticle labeling of cytochrome *c*

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Edited by Kenneth A. Dill, University of California, San Francisco, CA, and approved January 16, 2009 (received for review July 28, 2008)

Although nanoparticle-protein conjugates have been synthesized for numerous applications, bioconjugation remains a challenge, often resulting in denaturation or loss of protein function. This is partly because the protein-nanoparticle interface is poorly understood, which impedes the use of nanoparticles in nanomedicine. Although the effects of nanoparticle ligand and material on protein structure have been explored, the choice of the labeling site on the protein has not yet been systematically studied. To address this issue, we label cytochrome *c* site-specifically with a negatively charged Au nanoparticle via a covalent thiol-Au bond. The attachment site is controlled by cysteine mutations of surface residues. The effect of labeling on protein structure is probed by circular dichroism. Protein unfolding is the most severe when the nanoparticle is attached to the N- and C-terminal foldon, the core motif of cytochrome *c*. Also, when the nanoparticle is attached in the vicinity of charged residues, the amount of structural damage is greater because of salt-dependent electrostatic interactions with charged ligand bis(*p*-sulfonatophenyl) phenylphosphine on the nanoparticle. Molecular dynamics simulations also elucidate local to global structural perturbation depending on labeling site. These results suggest that the labeling site must be considered as one of the main design criteria for nanoparticle-protein conjugates.

bioconjugation | protein folding | circular dichroism | molecular dynamics simulation | nanoscale interface

Nanomaterials hold much promise for biomedical applications such as multiscale imaging (1–3), novel therapeutic approaches (4), and biomolecular sensing (5). The emerging field of nanomedicine is based on exploiting the unique size- and material-dependent properties of nanomaterials. However, the biggest barrier in effectively using them is their biological interface. Historically, biological-inorganic interfaces suffer complications such as surface fouling in medical devices and nonspecific adsorption of proteins on sensors, inhibiting practical use. The problem significantly worsens for nanoscale systems because of their high surface-to-volume ratio. Nanoscale surfaces are physically and chemically different from the bulk, requiring nontraditional tools to effectively probe them. These factors suggest a pressing need for deeper understanding of the “nano-bio” interface.

One particular issue related to the interface that limits the potential of nanoparticles (NPs) is conjugation to a biomolecule. Covalently linking a NP to a protein can affect both its structure and function. Even if protein structure is retained, the NP can sterically hinder substrate binding. NPs are large compared with dye molecules, and their interface with proteins is highly complex. NPs are not hard spheres but crystals with facets, edges, and vertices, and their passivating ligands can change conformation or come on and off the NP. Consequently, there are numerous nonspecific interactions between the NP and protein, where amino acid side chains noncovalently bind to the NP or ligand in undesirable ways. This often leads to protein denaturation, rendering the NP-protein conjugate useless.

Despite these complications, proteins on nanostructures are often assumed to be fully folded, an oversimplification that can severely obfuscate quantitative data in fluorescence resonance energy transfer (FRET) (6) and activity measurements (7), or

lead to fundamental obstacles in comprehending biological side effects and cytotoxicity. To date, efforts have focused predominantly on exploiting material properties of NPs (8, 9), but ultimately these endeavors are limited by a poor understanding of the NP-protein interface. To realize the full potential of NPs in biological applications, understanding and ultimately controlling these interactions is critical.

To this end, efforts have begun to elucidate how NP labeling affects protein structure and function. For covalently linked proteins, the structural effects of ligand charge (10, 11) and NP material (12) have been studied. However, a key factor not yet explored is the effect of the labeling site, which may affect the protein conformation and activity, and thus would be crucial for obtaining quantitative information from NP-protein conjugates. Whereas the advances in conjugation chemistry have enabled site-specific labeling, the choice of labeling site must be strategized. Proteins have different domains of varying structural and functional importance, and charge and hydrophobicity are rarely uniform on protein surfaces. Obviously, labeling sites necessary for protein function, such as binding or active sites, must be avoided. Furthermore, each domain on a protein can stabilize the structure differently, so labeling sites must be carefully chosen to minimize destabilization. We have previously shown that protein structure in NP-protein conjugates is dominated by interactions with amino acid side chains in local vicinity of the conjugation site (10, 12). However, to date, there is no study that systematically investigates the effect of labeling site on protein structure, and the choice of labeling site has been largely overlooked.

Here, we conjugate Au NPs to the protein cytochrome *c* (cyt *c*) at different sites and study the effect on protein structure. We show that electrostatic interactions of amino acid side chains with the NP ligand are partially responsible for protein behavior, but more importantly, there are key motifs on the protein that must not be labeled to avoid denaturation. Conversely, labeling a protein with a NP can be used to induce local denaturation. Based on this, we propose a general guideline for NP functionalization: To preserve protein structure, flexible and loosely folded motifs should be labeled instead of nucleation centers for folding. Fundamental understanding of the effect of labeling site on protein structure, in combination with material, size, and ligand dependence, will enable strategic labeling of diverse proteins with a variety of biomedical importance.

Results

Choice of Protein and NP. Cyt *c* is ideal for this study because its structure (13, 14) and folding have been extensively characterized, both experimentally (15, 16) and computationally (17, 18). Cyt *c* from *Saccharomyces cerevisiae* was used because its unique

Author contributions: M.-E.A.-T., W.H., and K.H.-S. designed research; M.-E.A.-T. performed research; M.-E.A.-T. analyzed data; M.-E.A.-T., W.H., and K.H.-S. wrote the paper.

The authors declare no conflict of interest.

This article is a PNAS Direct Submission.

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This article contains supporting information online at www.pnas.org/cgi/content/full/0807299106/DCSupplemental.

Table 2. α -Helical content of NP-cyt *c* and change in α -helicity in 0.1 M NaCl

Conjugate	No salt, %	0.1 M NaCl, %	$\Delta\alpha$ -helicity, %
H39C-NP	7.3	15.6	8.3
D50C-NP	16.2	17.5	1.3
E66C-NP	7.7	20.7	13.0
K99C-NP	4.8	6.1	1.3
C102-NP	5.6	6.9*	1.3

CD spectra were deconvoluted by CDSSTR algorithm (37).

* α -helicity reported previously for same NPs is higher (10) because of different labeling conditions (pH, the cyt *c*/NP labeling ratio, and DTT wash).

MD simulations. The NP was modeled as a 1.5-nm diameter dummy atom with 21 BPS ligands distributed on its surface. Inductively Coupled Plasma Optical Emission Spectrometry (ICP-OES) measured an S molar concentration $2.2\times$ lower than Au, confirming a BPS coverage comparable to that reported for triphenylphosphine on 1.5-nm Au NPs, 21 BPS/NP (22). We postulated that the protein displaces one BPS when forming a thiol bond with Au (16).

After building the NP-cyt *c* conjugates and performing initial minimization, we calculated distances of charged residues from the NP surface (Fig. 3 *A* and *B*). Compared with other conjugates, D50C-NP has significantly fewer charged residues near the NP surface. Other conjugates have a similar distribution of negatively charged residues. E66C-NP has slightly more positive charges in the vicinity of BPS. During MD simulations at 300 K, slight structural disturbance was observed. Because unfolding of proteins on NPs can take up to hours (23), we raised the temperature to 450 K to accelerate unfolding (24). We chose 450 K as unlabeled yeast cyt *c* remained folded over the course of the simulation. Snapshots of protein conformations at the end of the simulations at 300 K and 450 K (NP removed for clarity; Fig. 4) show that side chains of positively charged residues interact with BPS ligands (e.g., Fig. 5), and the secondary structures are perturbed differently depending on NP attachment site (Fig. 4).

Structural disturbances observed in MD simulations agree with secondary structure changes observed experimentally. Root-mean-square deviation (rmsd) of C_α atoms in the WT conformation (Fig. 6) reveals that NP labeling of residue 66 perturbs the 60's helix, whereas labeling residues 99 and 102 perturbs both the N- and C-terminal helices that are in contact. Labeling residue 39 perturbs the β -bridge at positions 38–39 and 58–59 (Fig. 4). This shows that the NP generally disturbs the structure to which it is attached. These results based on rmsd are also supported by analysis of α -helicity of each of the mutant conjugates and the WT (Figs. S2–S4).

To study the role of Coulombic interactions, the charge on each BPS molecule was increased from $-2e$ to $-4e$ ($e = 1.6 \times 10^{-19}$ C). This had a pronounced effect on the C102-NP conju-

gate, causing the N helix to unfold at 300 K within the 8-ns simulation time (Fig. 6).

Discussion

We find that the choice of labeling site for the NP can have a profound influence on protein stability. This conclusion was based on conformations of cyt *c* site-specifically labeled with NP at various positions, which were probed by using CD spectroscopy and further analyzed by MD simulations.

Electrostatic Effects. Confirming earlier observations (10), electrostatic interactions between the protein and the NP ligand play a prominent role in the protein stability. Addition of 0.1 M NaCl results in an improvement of folding for all labeled mutants (Fig. 2C, Table 2). Thus, the degree of denaturation can be tuned by salt, especially for H39C or E66C. Addition of 0.1 M NaCl increases charge screening, and enables the protein to retain structure. However, the amount of structural recovery for each conjugate is different, with $\Delta\alpha$ ranging from 1.3 to 13.0% (Table 2). The Debye length at 0.1 M NaCl is 0.96 nm. A large portion of charged residues are screened at 0.1 M NaCl, because they are located further than 0.96 nm from BPS O atoms (Fig. 3 *A* and *B*). Results of MD simulations are consistent with the secondary structural measurements from CD, and yield insight into molecular interactions involved. Interaction of the amino acid side chains with BPS molecules is observed (Fig. 5), underscoring the importance of modeling the NP ligand explicitly.

D50C-NP has significantly fewer charges near BPS molecules (Fig. 3 *A* and *B*). This conjugate is the most folded both experimentally and in simulations, as evidenced by the least α -helical disturbance (Figs. 4 and 6). The present analysis confirms local electrostatic effects previously observed in NP labeling of WT yeast cyt *c* at position 102 (10). It also suggests that based on electrostatics alone, the labeling of sites with the least number of charged residues in the vicinity results in the least structural perturbation.

NP Destabilizes the Whole Protein when Attached on Structures Essential for Folding.

However, the different amounts of denaturation and recovery on charge screening cannot be explained solely by electrostatics. In both CD and MD, NP labeling of K99C and C102 results in strong denaturation that is barely improved by salt addition ($\Delta\alpha = +1.3\%$). This contrasts with E66C-NP, which also has the surface Cys on an α -helical motif with a similar number of charged residues as K99C, but retains significantly more of its structure in presence of salt (7.7% \rightarrow 20.7%). Thus, local electrostatics are not the only factor influencing protein structure. When the NP is on positions 99 or 102 in the C-terminal α -helix, both the N- and C-terminal helices are substantially distorted (Figs. 4 and 6). This indicates that the mere presence of the NP is a large enough perturbation to the folding of these helices. Folding studies of cyt *c* have shown that

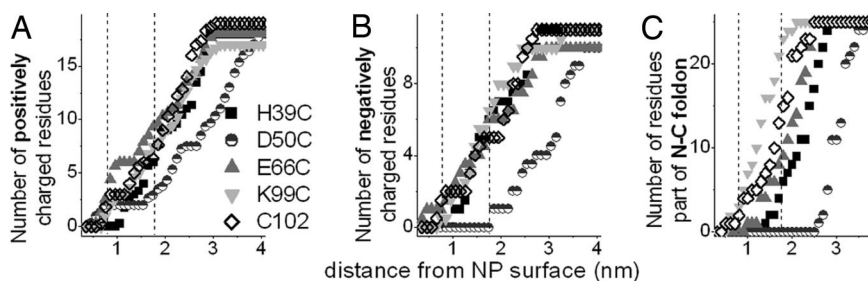


Fig. 3. Distribution of positive (*A*) and negative charges (*B*), and residues (*C*) part of the N-/C-terminal helices around the NP. Distances are between Au surface and amino acid charged end groups (*A* and *B*) or C_α atoms (*C*), before MD when the protein structure is folded. Dashed lines: Average distance of BPS O atoms from NP surface (8 Å) and one Debye length (9.6 Å) away from BPS O at 0.1 M NaCl.

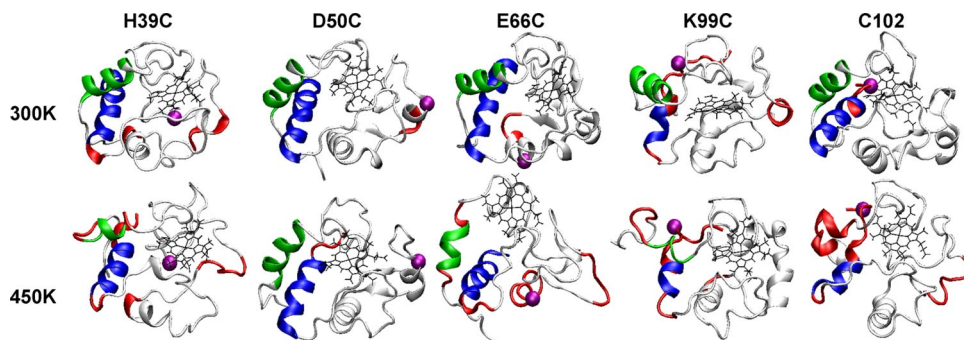


Fig. 4. Snapshots of protein structure at the end of 8-ns MD simulations at 300 K and 450 K. Green/blue: N-/C-terminal helices; red, loss of >50% α -helicity from WT structure (compare Fig. S2); purple sphere, NP attachment site. The program VMD was used for visualization (44).

these positions are on a motif critical for folding. Hydrogen exchange experiments on *cyt c* have shown that the N- and C-terminal helices, known as the N-C foldon, fold first (15) and bind together by hydrophobic residues at the interface. Four residues at the N-C interface, Gly-6, Phe-10, Leu-94, and Tyr-97, are the only non-heme-binding residues conserved among known *cyt c* sequences (25), underscoring the importance of this interface. Once the N-C foldon is locked in place, it guides the folding of the rest of the protein. Simulations also support this sequential stepwise folding pathway (17, 18). The N-C foldon is crucial for the stability of the whole protein and mutations in either the N- or C-terminal helix significantly affects folding (26, 27). It thus acts as a “safety pin” for the entire structure, while other motifs such as the 60’s helix, are less stable and thus can denature more locally. We monitor N-C foldon conformation in MD simulations via the solvent-accessible surface area (ASA) of its hydrophobic cluster (28). Although no significant change is observed for simulations with normal charges on BPS, ASA increases for K99C-NP and C102-NP when charges on BPS are enhanced (Fig. S5). Indeed, our contact analysis reveals that the N-C helices form a “hub” that is linked to other multiple domains that constitute additional foldons of *cyt c* (Fig. 7A) (29). The level of denaturation can also be characterized by changes in the contact map (Fig. 7 B and C and Fig. S6). Thus, the NP prohibits not only formation of the C-terminal helix but also the N-C helix contact, affecting the stability of the entire protein.

More moderate unfolding is observed when the NP is attached further from the N-C foldon (Figs. 2 and 3C). MD simulations of H39C-NP and E66C-NP show that the N-terminal amino group and Lys-2 (compare Table 1) are close enough to the BPS molecules in these conjugates to form salt bridges, which eventually lead to the disruption of the N helix (Fig. 5). In these conjugates, most of the N-C foldon is away from NP negative charges by more than the Debye length at 0.1 M NaCl (Fig. 3C),

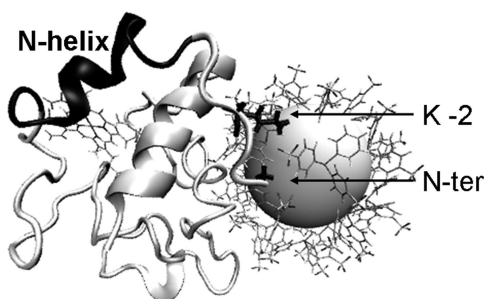


Fig. 5. Interaction of positive charges of K-2 and the N terminus with BPS. The snapshot (in “perspective” view) is from MD simulation of E66C-NP at 450 K.

which explains the considerable improvement in folding by salt. In contrast, NP labeling at position 50 does not affect the folding of the rest of the protein as dramatically, consistent with CD results. The N-C foldon is still maintained in this conjugate and contacts are more stable (Figs. 4, 6, and 7). Thus, induced disorder is more localized in D50C-NP.

To verify our simulations performed in an implicit solvent environment (*Materials and Methods*), for D50C-NP and K99C-NP, we performed explicit water MD simulations, where each production run lasted 4.5 ns at 450 K. A stronger structural disturbance in K99C-NP was again observed from the beginning of the simulation, albeit deformation occurred more slowly because of solvent viscosity (Figs. S7–S10).

Ramifications for NP-Labeling Strategies. The result here indicates that choice of labeling site is crucial for protein structure. It has been suggested that the network of contacts among amino acids in protein structures exhibits the “small-world” feature, in which there are a few key residues that play the role of hubs in the network, stabilizing the structure (30). Labeling motifs that contain these key residues result in the greatest amount of protein denaturation. A very large family of proteins has the N- and C-terminal motifs in contact, which has been suggested to control initial folding, native state stability, and turnover (29). Incidentally, protein termini are the most common NP labeling sites. His tags are predominantly put at N or C termini because of convenience in cloning, labeling chemistry, and conjugate purification. However, if the protein is part of the large family that possesses the N-C contact (29), it could potentially be the worst site for labeling, resulting in denaturation that cannot be rescued by salt screening, and ultimately loss of biological function. For these systems, labeling of low-stability motifs

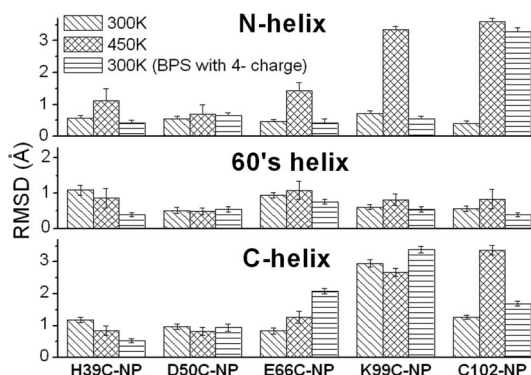


Fig. 6. rmsd of C_{α} atoms from the WT conformation averaged over the last 500 ps of MD. Error bar: standard deviation over the time period. Fig. S4 shows rmsd trajectories.

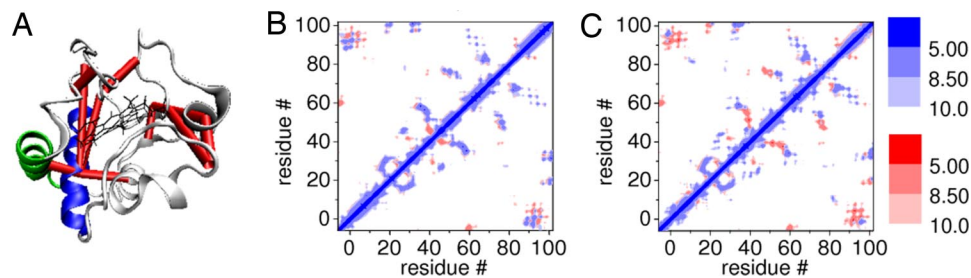


Fig. 7. Contact map of cyt c. (A) WT contact network. Centers of mass of 2 domains are connected by a red cylinder if there are residues with C_{α} atoms closer than 8.5 Å. Domains connected to the N–C foldon are known to fold before the network on the right (β -bridge and 50's helix) (45). Contact maps of D50C-NP (B) and K99C-NP (C), averaged over the last 500 ps of MD at 450 K (blue) on top of the WT contact map (red). Scale corresponds to distances between C_{α} atoms (Å). Wider red area in C indicates more denaturation. (See also Fig. S6.)

would be optimal. In addition to choosing the “least stable” motifs for labeling, sites should also have minimally charged residues in the vicinity to reduce electrostatic interactions. We find that the level of denaturation correlates with the number of charged amino acids in the local vicinity of the NP labeling site.

Furthermore, by inducing localized perturbations with a careful choice of labeling sites, there is potential for exploiting NP labeling as a tool for studying protein structure and folding dynamics. Obviously this effect strongly depends on NP size, because the amount of interaction with the NP ligand will vary with NP curvature (31, 32); 1.5-nm NPs are small enough to introduce local denaturation effects, because only residues close to the attachment site will interact with the NP and ligands. For larger NPs, disorder in the protein might not be so localized. Dependence on NP size and the effect on redox potential are the subject of future studies, in addition to application to other proteins.

Materials and Methods

Gold Nanoparticle Synthesis. The 1.5-nm diameter Au NPs were synthesized in toluene with triphenylphosphine ligands (22). Negatively charged BPS (Strem Chemicals) was used to passivate NPs to permit water solubility. Two-phase ligand exchange was used. NPs were drawn into aqueous phase in the presence of BPS. BPS-NPs were purified from excess BPS by size-exclusion HPLC (Varian ProStar) with TSK-gel G4000PW column (TOSOH Bioscience). TEM showed a $d = 1.5 \pm 0.2$ nm ($n = 298$). Optical absorption showed no plasmon peak, confirming a $d < 2$ nm and no aggregation. NP concentrations were determined by using $\epsilon_{420} = 110.0 \text{ mM}^{-1} \text{ cm}^{-1}$ (33). BPS and NP molar ratios were measured by ICP-OES (SPECTRO model Ciroc-Vision) assuming 100 atoms/NP.

Expression of Cyt c Mutants. WT cyt c from *S. cerevisiae* was purchased (Sigma). Mutant plasmids were obtained from the groups of Harry Gray and Jay Winkler (19) (Table 1, sequences with mutations in bold). C102S mutation is for removal of the WT Cys. Because *S. cerevisiae* cyt c expressed in *Escherichia coli* lacks the trimethylation at Lys-72 found in the native protein, a K72A mutation was necessary for preventing the heme from binding to Lys-72 (34). Pletneva et al. (20) determined that the mutations (H39C/K72A/C102S, D50C/K72A/C102S, E66C/K72A/C102S and K72A/K99C/C102S) do not significantly perturb folding.

Plasmids were sequenced (MIT Biopolymers Facility) to verify mutations, and cloned into *E. coli* BL21DE3 Star cells (Invitrogen). To increase yield, cyt c maturation gene cassette pEC86 with Chloramphenicol (Chl) resistance (35) and cyt c plasmid with Ampicillin (Amp) resistance were cotransformed. Colonies from LB agar plates with 100 $\mu\text{g}/\text{mL}$ Amp and 34 $\mu\text{g}/\text{mL}$ Chl were screened for best cyt c expression in 5 mL of TB medium containing 100 $\mu\text{g}/\text{mL}$ Amp and 34 $\mu\text{g}/\text{mL}$ Chl grown for 24 h shaking at 37 °C. When possible, colonies were selected by the pink color of the cell pellet (due to presence of cyt c) and saved for frozen stocks.

Cells from frozen stocks were used to inoculate 10 mL of TB medium containing 100 $\mu\text{g}/\text{mL}$ Amp and 34 $\mu\text{g}/\text{mL}$ of Chl. After overnight growth, large TB cultures (≈ 5 –10 L) containing 100 $\mu\text{g}/\text{mL}$ Amp and 34 $\mu\text{g}/\text{mL}$ of Chl were inoculated. Cells were grown for 36 h at 37 °C, harvested by centrifugation, and resuspended in 10 mM NaPi buffer (pH 7.0).

To isolate the protein, cells were lysed by French Press at 28 PSI (Constant

Systems Basic Z Model). 20 nM DNase, 1 mM phenylmethanesulfonyl fluoride (PMSF), and 1 mM DTT were added and the suspension was centrifuged at 6000 rpm for 30 min and the supernatant was collected. To purify the protein, the solution was loaded onto a Fast Flow CM Sepharose Econo-column equilibrated with 20 mM NaPi buffer with 1 mM DTT and 1 mM PMSF (pH 7.0). Cyt c was eluted with a stepwise salt gradient (0–0.5 M NaCl). Protein was further purified by ion-exchange chromatography with a Mono S Sepharose column by using a FPLC system with the same buffer, and then by size exclusion chromatography with TSK-gel G4000PW column on a HPLC system (Varian-ProStar). Protein purity was confirmed by SDS/PAGE. Cyt c concentrations were determined by using $\epsilon_{410} = 106.1 \text{ mM}^{-1} \text{ cm}^{-1}$.

Conjugation of Cyt c with NPs. Cyt c were washed from excess DTT with 5-kDa centrifugal filters at 4 °C and resuspended in 10 mM NaPi buffer at pH 7.3 with or without 0.1 M NaCl. NPs were attached to cyt c by incubation overnight in the same buffer at 4 °C and then studied by CD and absorption spectroscopy. Cyt c/NP incubation ratio was 1:1.5 to ensure that there is at most 1 protein per NP so as to remove effects of protein–protein interactions on the NP surface that may influence structure (12). To show site-specific labeling, WT yeast cyt c and horse cyt c (Sigma) were incubated with NPs in 0.5 \times TBE with and without 1 M NaCl for 1 h and loaded in a 1% agarose gel for electrophoresis in 0.5 \times TBE with a field strength of 10 V/cm. Gels were blue stained for protein (SimplyBlue SafeStain, Invitrogen). Samples were extracted from an identical gel for absorption spectroscopy with 0.2- μm centrifugal filters (Nanosep, Pall) at $10^4 \times g$.

Spectroscopic Analysis. Absorption spectra were obtained (Varian Cary 50 Spectrophotometer) at room temperature with a 1-cm pathlength cuvette. Scan rate was 5 nm/s. Far-UV CD spectra of proteins, NPs, and conjugates were obtained (Aviv model 202) at 25 °C in a 1-mm-pathlength cuvette. NP contribution to CD signal was subtracted. An isodichroic point at θ_{206} was found for heat-induced cyt c denaturation, as reported for glycerol-induced denaturation (36). Because cyt c concentration in NP:cyt c solutions could not be computed with high precision from absorption spectra because of NP absorbance in the Soret region, CD spectra of NP-cyt c conjugates were normalized to $\theta_{206} = -9.5 \times 10^3 \text{ deg cm}^2 \text{ dmol}^{-1}$. We used the CDSSTR algorithm (37) to deconvolve the spectra into % α -helicity.

Molecular Dynamics Simulations. MD simulations were performed by using CHARMM version 32 (38), with the generalized Born with simple switching (GBSW) implicit solvent (39). Although we did not retain 4 internal water molecules in the crystal, WT remained folded at 300K in simulations with GBSW. A 20-Å cutoff was applied for nonbonded interactions with a switch function between 16 Å and 20 Å. To verify results from simulations by using GBSW, explicit water simulations were performed for D50C-NP and K99C-NP (see *SI Text* and Figs. S7–S10). The param22 all-atom force field (40) was used and modified to include NP parameters. The NP core was modeled as one dummy atom with Lennard–Jones parameters ($\epsilon = 0.02$ and $r_{\text{min}}/2 = 14$ Å). Each BPS ligand was modeled explicitly with partial charges estimated from known parameters for the benzene ring and methyl sulfate. The BPS molecule was constructed from coordinates of a triphenylphosphine molecule linked to an Au atom (41). Twenty-one BPS molecules were uniformly distributed on the NP surface. Cyt c were constructed from Protein Data Bank (PDB) ID code 2BCN (42) and mutated accordingly. The heme was covalently linked to Cys 14 and 17. Covalent bonds were maintained between the heme iron and Met-80 and His-18, which are assumed to remain formed during the timescale of our simulations (17). Cyt c was linked to the NP surface by replacing one BPS ligand

by the labeled Cys. To prevent formation of close contacts, this was done by approaching the protein sequentially in steps of 1 Å, and performing an energy minimization between each step until the S of the thiol reached the target coordinate. During energy minimization and dynamics, the position of the S of the labeled Cys and the P atoms of BPS ligands were fixed at 10 Å from the NP center, based on reported P–Au and S–Au bond lengths (41, 43).

For MD, the leap-frog method was used with a 2-fs time step and the SHAKE algorithm to constrain bonds involving hydrogen atoms. The system was heated and equilibrated from 48 K to 300 K for 20 ps followed by a production run at 300 K for 8 ns, or from 48 K to 450 K for 40 ps and maintained at 450 K for 8 ns. During production runs, temperature fluctuated by 1.3%, and no

energy or temperature drift was observed. rmsd of C α atoms from energy-minimized WT configuration were calculated for the N, C, and 60's helices.

ACKNOWLEDGMENTS. We thank Harry Gray, Jay Winkler, and Jennifer Lee for generously providing plasmids and helpful advice on mutant expression and purification; Center for Materials Science and Engineering for the use of the transmission electron microscope; Computational and Systems Biology Initiative (CSBi) Biophysical Instrumentation Facility for CD spectrophotometer; the Van Vliet group for UV-vis spectrophotometer; MIT Nuclear Reactor Laboratory for Inductively Coupled Plasma Optical Emission Spectrometry; the CSBi Computational facility; the Thomas Schwartz lab for use of French Press; Ky Lowenhaupt for FPLC purification; and Viviana Serra and Sisi Zhu for help with experiments.

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