Immobilization of two organometallic complexes into a single cage to construct protein-based microcompartments†

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Natural protein-based microcompartments containing multiple enzymes promote cascade reactions within cells. We use the apo-ferritin protein cage to mimic such biocompartments by immobilizing two organometallic Ir and Pd complexes into the single protein cage. Precise locations of the metals and their accumulation mechanism were studied by X-ray crystallography.

Compartmentalization is a common strategy used by living organisms for controlling reaction environments within the cell such as the reactions of metabolic pathways. The carboxysome is a unique example of a protein-based microcompartment that encapsulates diverse enzymes into a single cage to promote CO2 fixation. Mimicking such compartment systems as new biomimetic materials is important for understanding complex cellular process as well as catalytic applications. In efforts to develop such compartment systems, foreign proteins are incorporated into the protein cage. In particular, Douglas and coworkers have reported an approach where multiple enzymes are packed into a single protein cage of virus-like particles (VLP). There remains the problem that encapsulated enzymes in the cages have lower activity than in free solution as a result of reduced stability, variable structure and other factors.

In one of the simpler ways to address this issue, such systems are mimicked by employing artificial metalloenzyme technology. Protein scaffolds provide useful reaction space when occupied by non-natural metal cofactors or metal ions and can promote various catalytic reactions. Recently, artificial metalloenzymes have been used to avoid mutual deactivation of catalysts when combined with biocatalysts for cascade reactions. Ward and coworkers have developed synthetic cascades by combining artificial transfer hydrogenase (ATHase) with oxidase enzymes. Protein cages that include metal nanoparticles are available for designing cascade reactions in combinations with native enzymes or metal complexes in buffer solutions. Although such combinations of biocatalysts and artificial metalloenzymes promote cascade reactions, it remains challenging to isolate each catalytic reaction within a single cage in a manner similar to reactions occurring in natural protein-based microcompartments. In this work, our objective was to construct a single protein cage containing multiple metal complex catalyst species to promote tandem reactions. We selected ferritin, a natural iron storage protein, because this self-assembled 24-mer protein provides a spherical cavity with an internal diameter of 8 nm that is stable over a wide pH range (2–11) and at temperatures up to 80 °C. Individual incorporation of various metal complexes into the apo-ferritin cage have been investigated in terms of the metal accumulation process, catalysis and drug delivery. We are now interested in whether two different organometallic complexes can be immobilized simultaneously on the interior surface of a single apo-ferritin cage to develop an artificial protein compartment (Fig. 1). Here we describe the preparation of apo-ferritin containing two different IrCp* and Pd(allyl) organometallic moieties and structural characterization of the resulting composite by X-ray crystallography. Catalytic reactions such as Suzuki–Miyaura coupling and hydrogenation reaction by the single protein cage containing dual organometallics are also described.

Immobilization of two different metal complexes within the single protein cage was performed by introducing the [IrCp*Cl2]2 complex into the recombinant L-chain apo-ferritin from horse liver (apo-rHLFr) followed by introduction of the [Pd(allyl)Cl]2 complex. The IrCp* immobilization reaction was carried out by stirring apo-rHLFr (10 μM in 50 mM Tris-HCl (pH 8.0), 0.15 M NaCl) with 85 equiv. of [IrCp*Cl2]2 for 1 h at 50 °C. After reaction, the mixture was dialyzed against 0.15 M NaCl to remove unbound IrCp* moieties and finally the ferritin protein cage containing IrCp* complexes ([IrCp*apo-rHLFr] was purified by gel filtration (Superdex G200). The number of IrCp* moieties accumulated in [IrCp*apo-rHLFr] was estimated using inductively coupled plasma-mass spectrometry (ICP-MS) and the bicinchoninic acid (BCA) assay. These analyses suggest that a total of 37 ± 4 Ir atoms...
are accumulated per apo-rHLFr cage. To determine the precise locations of the IrCp* moieties in the apo-rHLFr cage, we crystallized the \textit{IrCp*}-apo-rHLFr species using the hanging drop vapor diffusion method as reported previously.\textsuperscript{16,26} The crystal structure was refined to 1.56 Å resolution as shown in Fig. 2a. Selected crystallographic data and refinement statistics are listed in Table S1 (ESI†). The positions of the IrCp* moieties were determined from the anomalous difference Fourier maps (see ESI†). In the metal accumulation centre, two different IrCp* binding sites are present. A large ellipsoidally formed anomalous density is observed near His49. It is expected that several conformers of Ir–N\textsuperscript{(His49)} exist at the binding sites in the 24-mer cage of apo-rHLFr. Three different IrCp* binding positions (Ir1A, Ir1B and Ir1C) were modelled with three possible orientations of the imidazole moiety with a reasonable Ir–N distance range of 2.08–2.45 Å (Fig. 2c and Fig. S1, ESI†). The adjacent His173 was found to coordinate Ir1C with an Ir1C–N\textsuperscript{(His173)} distance of 2.40 Å. The total combined occupancy of Ir1A, Ir1B and Ir1C is 0.8, although the electron density corresponding to the Cp* ligand was not observed in the vicinity of the Ir ions due to the low occupancy of each Ir metal. The other IrCp* moiety (Ir2) was observed as bound to S\textsuperscript{(Cys48)} with a bond distance of 2.35 Å and an occupancy of 0.4 (Fig. 2e). At the 3-fold channel, there are three IrCp* binding sites from 3 different monomers (Fig. S2, ESI†). Ir3 is ligated by His114 with a bond distance of 2.13 Å and occupancy of 0.3. Phe128 coordinates to Ir4 (occupancy 0.2) with an Ir4–C\textsuperscript{(Phe128)} distance of 2.12 Å, which is comparable to the literature (Fig. S3, ESI†).\textsuperscript{34} The other binding site for Ir5 was observed at the 4-fold axis channel with occupancy of 0.2 (Fig. S3, ESI†). Although the total number of Ir binding sites observed in the crystal structure is 120, the sum of Ir atoms estimated from all of the occupancies is 45 in a single cage of apo-rHLFr. This value is consistent with that obtained from quantitative analyses (ICP/BCA) (37 ± 4). Thus, IrCp* moieties are expected to tend to coordinate His residues at the accumulation center and the 3-fold channel.

After immobilization of IrCp* into the apo-rHLFr cage and structural characterization, we focused on introducing organometallic [Pd(allyl)Cl]\textsubscript{2} complexes into the \textit{IrCp*}-apo-rHLFr structure to accumulate different metal complexes inside the single protein cage. The deposition of Pd(allyl) was carried out by stirring 100 equiv. of [Pd(allyl)Cl]\textsubscript{2} with \textit{IrCp*}-apo-rHLFr (5 μM) in 0.15 M NaCl at RT for 1 h. The mixture was purified by dialysis in 0.15 M NaCl followed by gel filtration (Superdex G200) to remove excess unbound Pd(allyl). The ICP-MS/BCA analyses indicate a total of 35 ± 3 Ir and 95 ± 6 Pd atoms present per apo-rHLFr cage. Interestingly, the number of IrCp* moieties remained almost the same as that of \textit{IrCp*}-apo-rHLFr. This indicates that Pd(allyl) binds at different locations than IrCp*. The crystal structure of ferritin cage containing IrCp* and Pd(allyl) complexes (\textit{IrCp*/Pd(allyl)}-apo-rHLFr) was refined to 1.87 Å resolution as shown in Fig. 2d. Selected crystallographic data and refinement statistics are listed in Table S1 (ESI†). The overall spherical structure of the apo-rHLFr cage is conserved in the presence of both IrCp* and Pd(allyl) complexes because of the root mean square deviation (rmsd) of the C\textsubscript{α} atoms from apo-rHLFr (0.23). The positions of Ir are distinguished from those of Pd metals by comparing the anomalous maps (see ESI†). The Ir binding sites are identical to those of \textit{IrCp*}-apo-rHLFr with similar occupancy values, except for the absence of IrCp* bound to S\textsuperscript{(Cys48)}. The Pd(allyl) moieties were identified by comparing our previously reported structures of \textit{Pd(allyl)}-apo-rHLFr.\textsuperscript{26} At the accumulation centre, there is one IrCp* binding site and two Pd(allyl) binding sites (Fig. 2f). The coordination of IrCp* at His49 remained the same as in \textit{IrCp*}-apo-rHLFr with total occupancy of 1.0. The electron density corresponding to His173, which coordinates to Ir1C in \textit{IrCp*}-apo-rHLFr, was not determined due to insufficient density (Fig. 2e and f). The observed Ir–N\textsuperscript{(His49)} bond distances are 2.07–2.61 Å. The IrCp* binding sites...
site at Cys48 in IrCp*/apo-rHLFr was replaced by two Pd(allyl) moieties in IrCp*/Pd(allyl)apo-rHLFr with occupancies of 1.0 (Pd1) and 0.7 (Pd2), respectively. This suggests that Pd(allyl) tends to coordinate to Cys. Glu45 showed dual conformations that coordinate to Pd(allyl). The Pd(allyl) moieties maintain a S*(Cys48) bridged dinuclear structure supported by the carboxyl group of one of the adjacent Glu45 conformers (Fig. 2f). The other conformer of Glu45 coordinates to Pd1, which has higher occupancy than Pd2. Both the Pd1 and Pd2 maintain a square-planar geometry with an allyl ligand, S*(Cys48) and O*(Glu45). The structure is similar to the reported dinuclear Pd(allyl) complexes.26,35 The observed Pd1–Pd2 distance was 2.95 Å. The structure is similar to the reported dinuclear Pd(allyl) and 2.76 Å, respectively, whereas the Pd1–O*(Glu45) and Pd2–O*(Glu45) distances are 2.40 Å and 2.51 Å, respectively. The Pd1–O*(Glu45) distance of alternate Glu45 conformer was 2.29 Å. The anomalous intensity map at the 3-fold channel suggests that both Ir2 and Pd3 are present with occupancies of 0.4 and 0.6, respectively (Fig. S2 and S4, ESI†). His114 coordinates to Ir2 at an Ir–N*(His114) distance of 2.71 Å. The short distance between the imidazole (His114) and Pd3 indicates the possibility of double conformations of His114 because both metals have occupancy of ~0.5 (Fig. S4, ESI†). The observed distance between the Pd3 and N* of the alternate conformation of His114 is 2.42 Å. The structure indicates that Cys126 coordinates to Pd3 when it is ligated by the respective His114 conformation (Fig. S4, ESI†). The observed Pd3–S*(Cys126) and Ir2–S*(Cys126) distances are 2.55 Å and 3.19 Å, respectively. The Ir binding sites at Phe128 and the 4-fold channel remained the same as in IrCp*/apo-rHLFr (Fig. S3, ESI†). The total number of Ir and Pd binding sites per apo-rHLFr cage observed from the crystal structures is 96 and 72, respectively. The sum of the Ir ions estimated from all occupancies is 40 and this is consistent with the number (35 ± 3) obtained from the ICP/BCA assay. However, the number of Pd ions (95 ± 6) determined from the quantitative analysis is higher than the total number of Pd binding sites (72). This suggests that several Pd(allyl) complexes are located at nonspecific binding sites as reported previously.36

Interestingly, although we attempted to prepare the heterometallic composite in a reversed procedure with [Pd(allyl)Cl]2 incorporated first, followed by [IrCp*Cl]2, and the reaction was unsuccessful because of the precipitation. We suspect that because Pd(allyl) forms a stable dinuclear structure involving coordination to His49 or His114 in Pd(allyl)apo-rHLFr, the IrCp* deposition in the second step may not be able to displace Pd and, as a result, random immobilization and precipitation occur.26 This observation suggests that the particular reaction sequence is important for immobilization of different metal complexes into a single protein cage. The process of accumulation of Pd(allyl) was understood when 50 equiv. of [Pd(allyl)Cl]2 was reacted with IrCp*/apo-rHLFr and determined the intermediate structure [IrCp*/Pd(allyl)]apo-rHLFr (Fig. S5, ESI†). The positions of the IrCp* and Pd(allyl) in the intermediate structure remained the same as in IrCp*/Pd(allyl)apo-rHLFr except for the coordination structures and occupancies of Pd.
coordination for catalytic hydrogenation reaction. We also performed the reaction using the mixture of \textit{IrCp}\textsuperscript{*}-apo-rHLFr and Pd(allyl)-apo-rHLFr, which gives similar yield for biphenyl ethanol as for \textit{IrCp}\textsuperscript{*}/Pd(allyl)-apo-rHLFr. It is expected that the apo-rHLFr cage is not diffusion limiting for small substrates and the intermediate between the cages as observed for VLP containing enzymes.\textsuperscript{7} The results demonstrate the possibility of achieving tandem reactions within a single protein cage by immobilizing different metal catalysts. Optimization of the catalytic reaction to improve activity and selectivity is now underway.

In summary, we have demonstrated immobilization of two different organometallic \textit{IrCp}\textsuperscript{*} and Pd(allyl) complexes into a single apo-rHLFr cage for promotion of tandem reactions. The X-ray crystal structure analyses suggest that \textit{IrCp}\textsuperscript{*} tends to bind to His whereas Pd(allyl) forms a dinuclear structure at Cys48. Both metal complexes are catalytically active within the cage. There is scope for improving catalytic efficiency. Incorporation of multiple complexes within a cage based on principles of coordination chemistry is expected to be a useful strategy for mimicking protein-based micromachinations.

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Notes and references