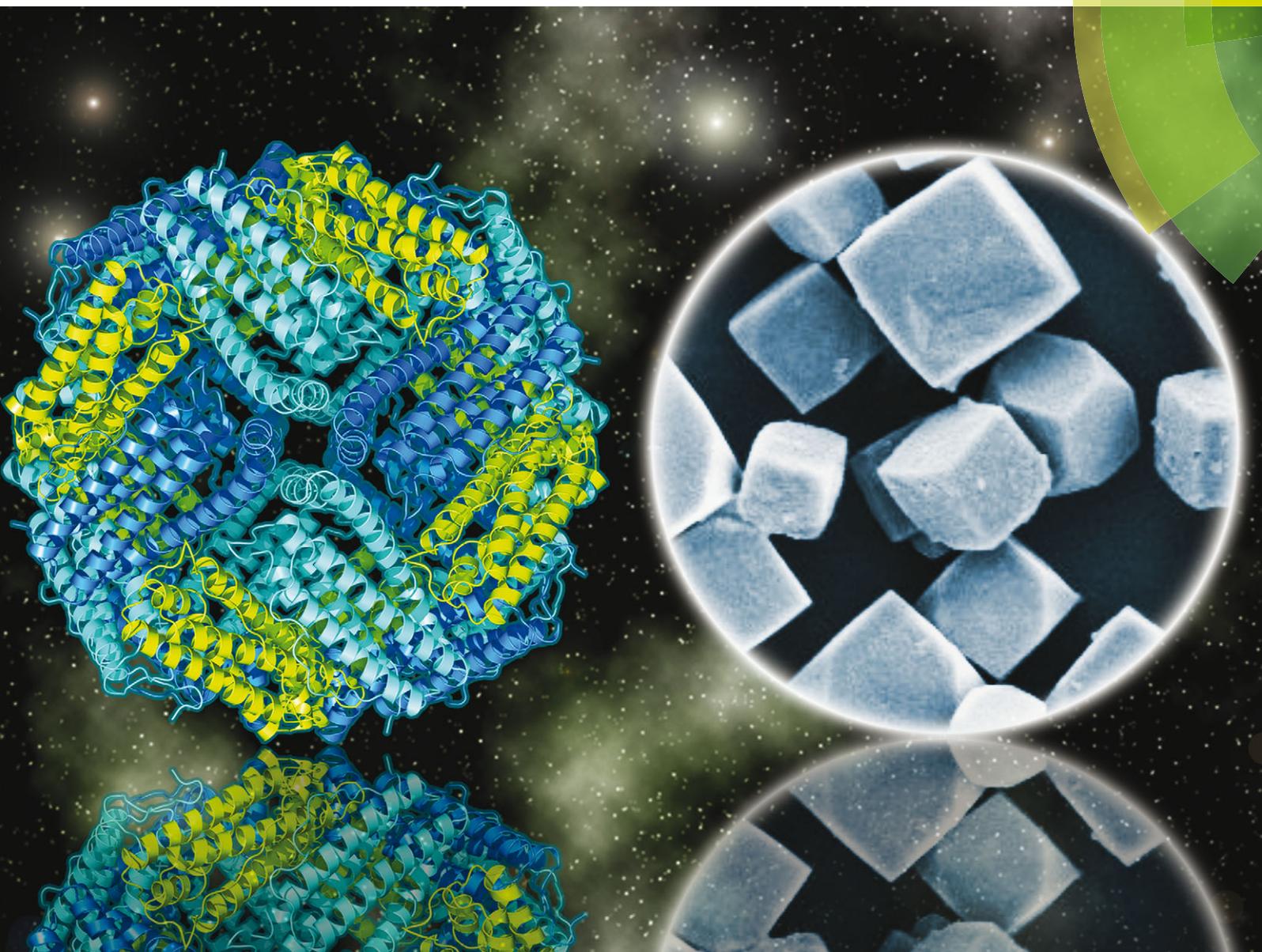


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FEATURE ARTICLE

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Design of a confined environment using protein cages and crystals for the development of biohybrid materials

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Design of a confined environment using protein cages and crystals for the development of biohybrid materials

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There is growing interest in the design of protein assemblies for use in materials science and bionanotechnology. Protein assemblies, such as cages and crystalline protein structures, provide confined chemical environments that allow immobilization of metal complexes, nanomaterials, and proteins by metal coordination, assembly/disassembly reactions, genetic manipulation and crystallization methods. Protein assembly composites can be used to prepare hybrid materials with catalytic, magnetic and optical properties for cellular applications due to their high stability, solubility and biocompatibility. In this feature article, we focus on the recent development of ferritin as the most promising molecular template protein cage and *in vivo* and *in vitro* engineering of protein crystals as solid protein materials with functional properties.

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1. Introduction

In recent years, strategies for the development of biomaterials using protein assemblies have attracted considerable attention in the fields of materials science and bio-nanotechnology due to their wide variety of applications.^{1–6} Highly ordered self-assembled proteins provide unique chemical environments for preparation of uniform nanomaterials and functionalization of metal complexes for applications in catalysis, imaging, and drug delivery, for example. Various protein assemblies, such as cages, tubes, and crystalline proteins, are utilized for such purposes.⁷

Although various synthetic templates, such as polymersomes, liposomes and dendrimers, are also known to have similar properties to nanomaterial scaffolds,^{8–11} the monodispersity, solubility and stability of protein assemblies under biological conditions are significantly distinct from those of synthetic materials. Among the various known classes of protein assemblies, we have particular interest in cages and crystalline protein assemblies for the following reasons; (i) the confined environments of proteins can accommodate various functional molecules, (ii) incorporation of the foreign substances into the interior space can be achieved by assembly/disassembly reactions, genetic manipulations, and protein crystallization methods, and (iii) these protein assemblies can be used in both *in vivo* and *in vitro* systems. In this feature article, we focus on recent significant development of ferritin cages as well as *in vivo* and *in vitro* protein crystal engineering.

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2. A protein cage for controlling cellular functions

Protein cages are formed from self-assembling proteins which provide a confined environment. Examples include virus capsid, heat shock protein (hsp), ferritin and DNA-binding protein from starved cells (dps) (Fig. 1a).¹² Nature uses such confined protein compartments to protect interior molecules from the surrounding environment as observed in carboxysomes, in a manner similar to the function of organelles.^{13,14} The concept has been utilized to synthesize nanomaterials which can avoid self-aggregation.^{15,16} Similarly, small molecules or enzymes are incorporated into the cages.^{2,17} The advantages of using a protein cage include increased aqueous solubility, stability and biocompatibility of functional molecules when loaded into the cage.¹⁸ Virus capsids and ferritin are the two widely used protein cages for cellular applications. The size of the virus capsids or virus-like particles is larger (>20 nm) than that of the ferritin cage (12 nm) and often used for encapsulating larger substances like enzymes, polymeric materials, *etc.* for biological applications.^{17,19–23} For example, the interior cavity of the P22 capsid was used to incorporate Gd complexes embedded in the polymeric matrix for developing magnetic resonance imaging (MRI) contrast agents.^{24,25} Similarly, a CCMV capsid was used to incorporate a photosensitive self-immolative polymer which undergoes a head-to-tail depolymerization into its monomeric subunits when irradiated with light, resulting in the slow release of the molecular cargo.²⁶ Chemotherapeutic drug taxol was functionalized in MS2 virus capsids which release the drug in MCF-7 cells.²⁷ In contrast, the ferritin cage with a smaller cavity (8 nm) has specific metal binding sites and can be functionalized with a number of synthetic metal complexes or small organic molecules for biomedical applications. Importantly, the ferritin has specific cell targeting ability and internalizes through the transferrin receptor mediated endocytosis pathway.²⁸ This is the important distinction of ferritin from other protein cages. Ferritin cages have potential applications

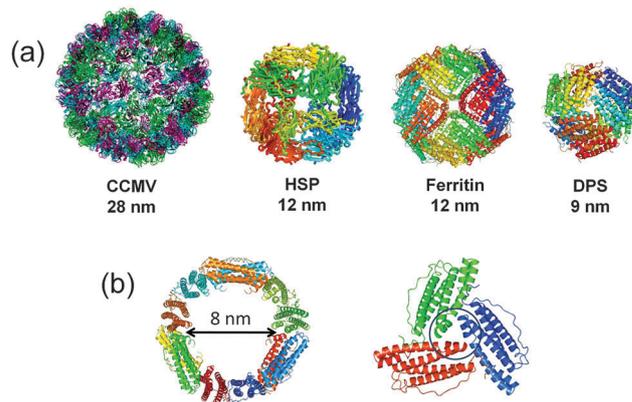


Fig. 1 (a) Library of protein cages found in nature such as cowpea chlorotic mottle virus (CCMV; pdb ID: 1ZA7), heat shock protein (HSP; pdb ID: 1SHS), ferritin (pdb ID: 1DAT) and DNA binding protein from starved cells (DPS; pdb ID: 1QGH). (b) The intersection of the ferritin cage and a 3-fold symmetric pore.

in designing new materials, imaging, drug delivery, and catalysis.²⁹ Several reviews have been published on the use of the protein cages such as ferritin cages and virus capsids as nanoreactors.^{3,18,30} Ferritin is a highly stable 24-mer self-assembled spherical protein cage.³¹ The outer diameter of the cage is 12 nm and the inner diameter is 8 nm (Fig. 1b). The natural function of ferritin is to store iron in a compact mineral form. The small 3-fold axis channels present in the ferritin cage provide the gateway for iron transport into the interior of the cage. It is known that a number of metal ions, metal complexes and organic molecules can be immobilized on the interior surface of apo-ferritin (apo-Fr) (Table 1). Due to their excellent biocompatibility, functionalized ferritin cages are used as carrier molecules for various biological applications (Table 1). The unique advantages of the ferritin cage are harnessed for the targeted delivery of small molecules and nanomaterials. We have focused our efforts on the ferritin cage which we believe is the most promising natural carrier protein cage. We discuss the recent advances in efforts to functionalize the ferritin cage for cellular applications.

2.1 Nanomaterials in the ferritin cage

The uniform size of the ferritin cage is utilized to prepare monodisperse nanoparticles which have several applications in catalysis and imaging, among others. For example, uniform Pd nanoparticles have been prepared in a ferritin cage and used for catalytic olefin hydrogenation and an aerobic alcohol oxidation reaction.^{32–34} Biocompatible nanomaterials can be prepared using the ferritin template in one step. In contrast, conventional methods require additional capping or targeting ligands. The ferritin-nanoconjugates are easily taken up by cells expressing transferrin receptors and have been demonstrated to reach their targets. This is significant because targeting often represents a problem for most types of nanoparticles. Considering such advantages, several ferritin-nanocomposites have been developed for various cellular applications such as enzyme mimetics and bioimaging, among others.^{36,41,42,69}



Takafumi Ueno

Takafumi Ueno received his PhD from Osaka University, Japan, in 1998. He held a position as an assistant professor at the Institute for Molecular Science in Okazaki, Japan. From 2002 to 2008, he worked as an assistant professor at Nagoya University and then took a position as an associate professor at iCeMS at Kyoto University. In March 2012, he moved to the Tokyo Institute of Technology as a full professor.

His current research interests include the molecular design of artificial metalloproteins and the exploitation of meso-scale materials using protein assemblies.



Table 1 Recent functionalization of the ferritin cage

Composites	Functions	Ref.
PdNP	Olefin hydrogenation	32
	Aerobic alcohol oxidation	33
Au/Pd (core/shell)	Olefin hydrogenation	34
AuNP	Catalysis: nitro phenol reduction	35
Au nano cluster	Kidney specific targeting nanoprobe	36 and 37
Co doped ferrite NP	Theranostic agent	38
Fe ₃ O ₄	Targeting/imaging tumor tissue	39
Fe ₃ O ₄ /Cy5.5	Imaging	40
CeO ₂ NP	Artificial redox enzyme activity	41
PtNP	Catalase and peroxidase activities	42 and 43
	Anticancer drug	44
PbS quantum dot	Antitumor activity	45
[Rh(nbd)Cl] ₂ ^a	Polymerization of phenyl acetylene	46
[Pd(allyl)Cl] ₂	Suzuki-Miyaura coupling	47
[Ru(CO) ₃ Cl ₂] ₂	CO releasing molecule	48
	NF-κB activation	
[Mn(CO) ₅ Br]	CO releasing molecule	49
	Light controlled NF-κB activation	
Catecholamide ^b /FeCl ₃	Siderophore mimic: enterobactin	50
Zn-phthalocyanine	Drug delivery: photodynamic therapy	51
Oxaliplatin	Drug delivery	52
K ₂ PtCl ₄	Drug delivery	53
Carboplatin		
Cisplatin		
Gd chelate complex	MRI contrast agent	54–56
Curcumin/Gd-chelate	Theranostic agent	56 and 57
Mn ²⁺	MRI sensor of melanin formation in melanoma cells	58
	T ₂ contrast agent	59
RGD-4C, ^c Cy5.5 and ⁶⁴ Cu	Multimodal imaging	60
NTTA ^d -Eu ³⁺	Bioprobe for time-gated luminescence bioimaging	61
PTTA ^e -Tb ³⁺	Nitric oxide probe	62
Doxorubicin	Drug delivery	63–65
Methylene blue	Drug delivery: photodynamic therapy	66
Near-infrared dye IR820	Multimodal imaging and photothermal therapy	67
AF350/AF430 ^f	Fluorescence resonance energy transfer	68

^a nbd = norbornadiene. ^b Catecholamide = *N*-(2-(2,5-dioxo-1*H*-pyrrol-1-yl)ethyl)-2,3-dihydroxybenzamide. ^c RGD-4C = Cys-Asp-Cys-Arg-Gly-Asp-Cys-Phe-Cys. ^d NTTA = *N,N,N',N'*-[40-(1-naphthyl)-2,20:60,20,0-terpyridine-6,60,0-diyl]bis(methylenenitrilo)tetrakis(acetic acid). ^e PTTA = *N,N,N'(1),N'(1)-[[4'-phenyl-2,2':6',2'-terpyridine-6,6'-diyl]bis(methylenenitrilo)tetrakis(acetate)diyl]bis(methylenenitrilo)-tetrakis(acetate)*. ^f AF = Alexa Fluor.

SOD mimicking nanoceria (CeO₂) particles have been incorporated into the ferritin cage *via* a dissociation–reconstruction route (Fig. 2a).⁴¹ The protein scaffold provides biocompatibility and higher cellular uptake *via* a receptor mediated pathway. The cage also manipulates the electron localization at the surface of nanoparticles. The ROS (reactive oxygen species) scavenging activity of apo-Fr encapsulated CeO₂ is 3.5 times higher than that of the natural SOD enzyme. The activity was tested in HepG2 cells using DCFH-DA (2,7-dichlorofluorescein diacetate) dye which is used to measure the ROS level. The effective quenching of the green fluorescence of DCFH-DA suggests that the system has excellent ROS scavenging activity in living cells. The iron oxide core was prepared in the apo-Fr cage to develop ferritin containing magnetic particles (magnetoferritin) which has applications in targeting and visualizing tumor tissues.³⁹ Since the iron core is formed inside the ferritin cage, the nanoparticles in magnetoferritin are easily taken up by overexpressed transferrin receptor 1 (TfR1) of tumor cells.

The composite catalyzes the oxidation of peroxidase substrates such as 3,3,5,5-tetramethylbenzidine (TMB) or di-azaminobenzene (DAB) in the presence of hydrogen peroxide to

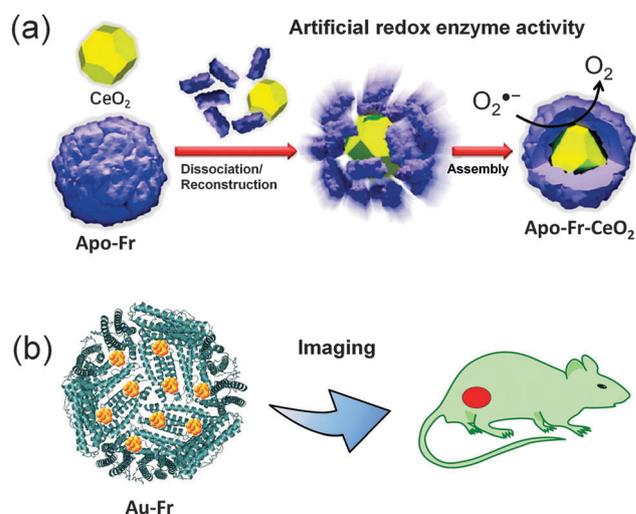


Fig. 2 Applications of nanomaterials prepared in the ferritin cage. (a) The SOD mimicking apo-Fr-CeO₂: preparation and functions. Reproduced from ref. 41 with permission from RSC. (b) Use of ferritin templated gold nanoclusters for bioimaging.



produce colored products which are utilized for visualizing the tumor tissues. The formation of hydroxyl radicals on the surface of the iron core is thought to be responsible for the catalytic oxidation reaction. Magnetoferritin can be used as an efficient diagnostic tool for cancer cell detection in a number of clinical samples. In a recent report, it was demonstrated that iron oxide can be loaded into the ferritin cage and functionalized with near-infrared emitting dye (Cy5.5).⁴⁰ The hybrid nanocomposite exhibits extremely high transversal relaxivity (r_2) up to $224 \text{ mM}^{-1} \text{ s}^{-1}$) and can cross the endothelium, epithelium and blood brain barrier layers. The composite can specifically target tumors overexpressing TfR1. Thus the ferritin based nanoprobe is used for targeted and ultra-sensitive imaging. There are several other reports describing the incorporation of Pt nanoparticles into the ferritin cage to mimic catalase and SOD activity.^{42,43}

PbS quantum dots (QDs) can be prepared inside the ferritin cage for effective inhibition of growth of colorectal carcinoma cells *via* generation of ROS.⁴⁵ The PbS-ferritin QDs are well tolerated by mice and no adverse effects were observed. Interestingly, the PbS-ferritin QDs were found to have NIR photoluminescence properties which are useful in combination with anticancer activity for cancer diagnostics, imaging and treatment.

The unique structure of the ferritin cage provides a suitable template for preparing small nanoclusters which have interesting fluorescence properties and can be utilized for cellular imaging. Nie and coworkers have developed an interesting strategy to assemble two gold clusters at the ferroxidase site of L-chain ferritin containing 2 H-chain subunits.³⁶ The coupling interaction between the paired Au nanocluster gives enhanced fluorescence and is red shifted compared to the free clusters. The emission of the paired Au clusters can be tuned from green to far-red fluorescence. This emissive property was used for whole body imaging and it was found that the composite specifically targets kidney tissue (Fig. 2b). In a similar method, multiple Au clusters were prepared in H-chain ferritin which exhibits near-infrared (NIR) fluorescence with high quantum yield about 63% and used as a kidney targeting NIR imaging agent.³⁷

2.2 Metal complexes in the ferritin cage

In addition to the preparation of nanomaterials, the ferritin cage provides suitable coordination environments for metal complexes for the generation of composites used in various applications. We have recently reported immobilization of organometallic $[\text{Pd}(\text{allyl})\text{Cl}]_2$ and $[\text{Rh}(\text{norbornadiene})\text{Cl}]_2$ complexes into the ferritin cage to promote catalytic C–C coupling and polymerization reactions and determined X-ray crystal structures of the composites.^{47,70,71} The ferritin cage has also been functionalized with several different metal complexes with the aim of controlling cellular functions, developing imaging agents and for drug delivery. The potent anticancer drug cisplatin and its analogues have been incorporated into the ferritin cage *via* an assembly/disassembly route (Fig. 3a).^{52,53} To enhance the loading efficiency, Pt particles were synthesized inside the ferritin cage by incubating K_2PtCl_4 in ammonia buffer (pH 8.5). The ferritin-encapsulated cisplatin has higher uptake efficiency

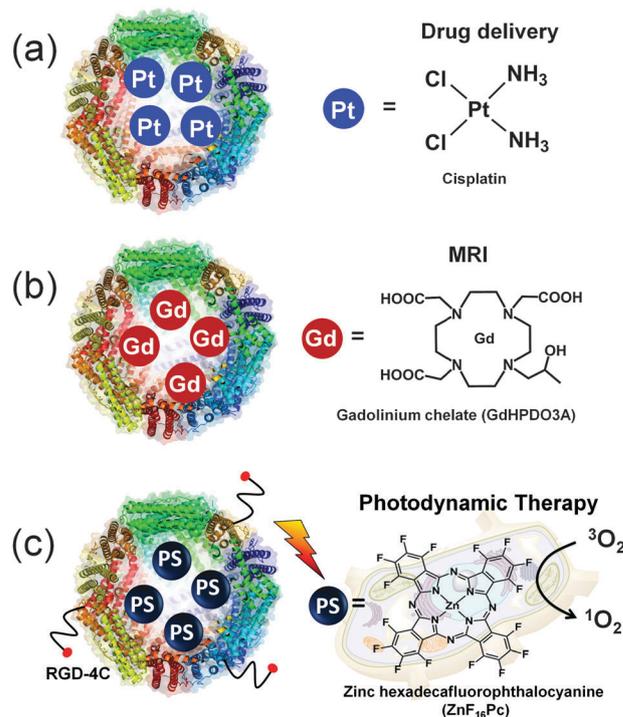


Fig. 3 Applications of the ferritin cage in drug delivery and imaging. (a) Incorporation of cisplatin into the ferritin cage for drug delivery.⁵³ (b) Gadolinium chelate complex in the ferritin cage for magnetic resonance imaging (MRI).⁵⁴ (c) Accommodation of zinc-phthalocyanine in the RGD-4C modified ferritin cage for targeted photodynamic therapy.⁵¹

than the naked drug which suggests that the system could overcome the cisplatin-related resistance problems. Receptor mediated endocytosis is thought to be responsible for the higher uptake of ferritin loaded platinum drugs. The cytotoxicity was found to be less than that of naked cisplatin due to limited release from the cage. However, the results indicate that ferritin has potential to be developed as a promising carrier for platinum-based anticancer drugs.⁵²

A Gd(III)-chelate complex has been incorporated into the ferritin cage for the development of a high relaxivity contrast agent for magnetic resonance imaging (Fig. 3b).⁵⁴ The cage can accommodate about 10 paramagnetic units and provide long-term stability with a low leaching rate. The resulting composite has 20 times higher longitudinal relaxivity than the naked chelate complex in aqueous medium. It is expected that the interaction between the protein interior cage and free exchangeable protons is responsible for the higher relaxivity. The Gd chelate has been loaded with curcumin, a polyphenolic compound for MRI guided treatment of cancer cells overexpressing ferritin receptors.^{56,57}

In a similar manner, Mn-loaded ferritin has been developed as an MRI sensor of melanin formation in melanoma cells.⁵⁸ The Mn^{2+} in the engineered ferritin cage exhibits high T_2 relaxivity indicating its potential for use as an ultrasensitive T_2 contrast agent.⁵⁹

For targeted delivery, the surface of the ferritin cage was modified with the RGD-4C (Cys-Asp-Cys-Arg-Gly-Asp-Cys-Phe-Cys)



peptide which selectively binds to integrin $\alpha_v\beta_3$, a tumor angiogenesis gene which is overexpressed in tumor cells.⁷² The RGD-4C-modified ferritin cage was then used to load a photoactive drug for selective delivery and controlled activation (Fig. 3c).⁵¹

Photoactive zinc hexadecafluorophthalocyanine ($ZnF_{16}Pc$) has poor pharmacokinetics due to its aqueous insolubility. The solubility of this compound was found to be significantly improved by loading it into the RGD-4C modified ferritin cage. Despite the high loading (60 wt%) of $ZnF_{16}Pc$, the overall particle size of apo-Ferritin remains the same. The photoactive ferritin nanocarrier has high tumor uptake efficiency in U87MG cells through RGD-integrin interaction and toxicity due to singlet oxygen generation when irradiated with red light. Recently, the $ZnF_{16}Pc$ loaded ferritin has been functionalized with folic acid as a tumor targeting ligand.⁷³ The nanoconjugate enters cells *via* a folic acid receptor-mediated endocytosis pathway and suppresses tumor cell growth only in the presence of light while minimizing effects on normal cells. Interestingly, the conjugate exhibits PDT-stimulated suppression of lung tumor metastasis *in vivo*. Thus, the functionalized ferritin carrier has potential applications in targeted photodynamic therapy.

Recently, Ueno *et al.* used the apo-ferritin cage to deliver carbon monoxide (CO) into living cells (Fig. 4).⁴⁸ CO in living cells acts as a signaling molecule to produce cytoprotective effects that counteract inflammation, proliferation and apoptosis.⁷⁴ The organometallic $[Ru(CO)_3Cl_2]_2$ complex used as a carbon monoxide releasing molecule (CORM) was incorporated into the cage of recombinant horse L-chain apo-ferritin (apo-rHLFr) (Fig. 4a). The resulting composite was found to have better activity than $Ru(CO)_3Cl(glycinate)$ (CORM-3), a typical CORM.⁴⁸ The RuCO in the ferritin cage was substantially immobilized at the metal accumulation site and the 3-fold axis channel as determined by X-ray crystal structure analysis (Fig. 4b). The Fr-CORMs have ATR-IR CO stretching frequencies near 2040 and 2060 cm^{-1} indicating the presence of a *cis*- $Ru(CO)_2$ coordination structure. The protein cage stabilizes the *cis*- $Ru(CO)_2$ structure and reduces the rate of CO release compared to CORM-3. The slow release of CO is important for cellular applications as it is toxic to cells. The uptake of Fr-CORMs in living cells is 4-fold higher than that of CORM-3 and it has been confirmed that the composite releases CO inside the cell (Fig. 4c). Fr-CORMs were found to activate the nuclear factor κB (NF- κB), a transcriptional regulator of a number of pro-inflammatory and anti-apoptotic genes and considered as a potential therapeutic target of CO. The NF- κB activity of the non-toxic Fr-CORMs in the presence of tumor necrosis factor α (TNF- α) was found to be up to 4-fold higher relative to CORM-3. This shows the potential of a protein cage for controlling cellular functions.

After utilization of Fr-CORM, a photoactive CORM was constructed inside the ferritin cage for dose-regulated delivery in living cells.⁴⁹ Photoactive $Mn(CO)_5Br$ was incorporated into the ferritin cage and characterized by X-ray crystal structure analysis (Fig. 4d). Although the CO ligands were not observed in the crystal structure, the ATR-IR was found to have three different CO stretching frequencies at 2028, 2011 and 1917 cm^{-1} corresponding

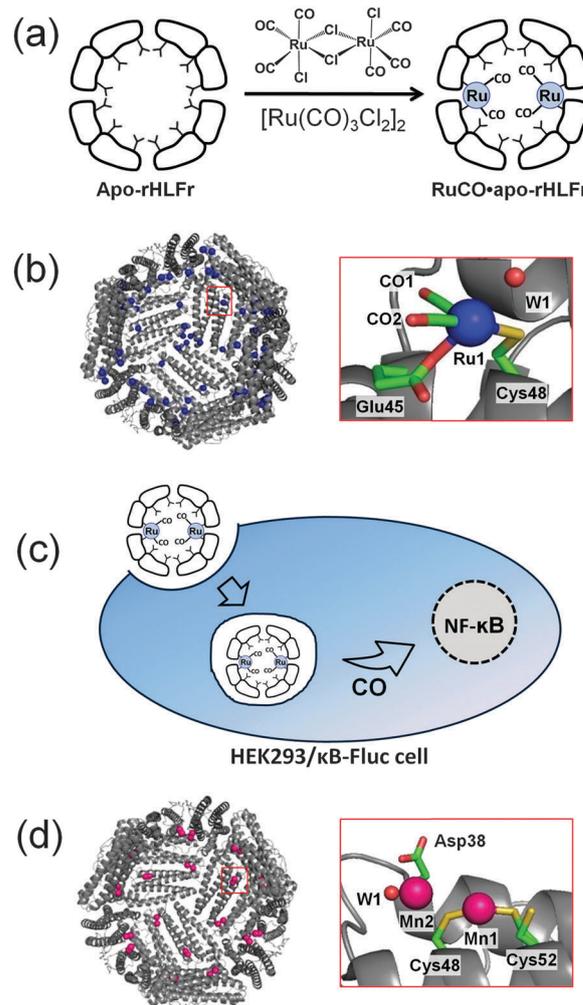


Fig. 4 Carbon monoxide releasing ferritins. (a) RuCO incorporation into the ferritin cage.⁴⁸ (b) Crystal structure of RuCO-ferritin showing the whole structure and Ru binding at the metal accumulation site. (c) Cellular uptake in the HEK293/ κB -Fluc cell and activation of NF- κB by CO releasing RuCO-ferritin. (d) Crystal structure of photoactive MnCO-ferritin showing the whole structure and Mn binding at the metal accumulation site.⁴⁹

to the *fac*- $Mn(CO)_3$ coordination structure. Irradiation of visible light of 465 nm activates the MnCO-ferritin composite and releases CO in a time-dependent manner. The composite has uptake efficiency similar to CORM-1 ($Mn_2(CO)_{10}$) in HEK293/ κB -Fluc cells and can be activated inside living cells in a controlled manner using light. The CO released from MnCO-ferritin in the HEK293/ κB cell activates NF- κB in the presence of tumor necrosis factor α (TNF- α). In contrast, the naked CORM-1 does not show any activity despite similar uptake efficiency and large amounts of released CO relative to MnCO-ferritin. Therefore, the protein cage specifically stabilizes the photoactive MnCO moiety for light controlled activation of NF- κB .

In addition to applications in therapeutics and control of cellular functions, the ferritin cage can be engineered with neutral metal coordination moieties for biomimetic applications. The symmetric 3-fold axis channels formed by 3 monomer subunits of apo-ferritin are known to accommodate and transfer



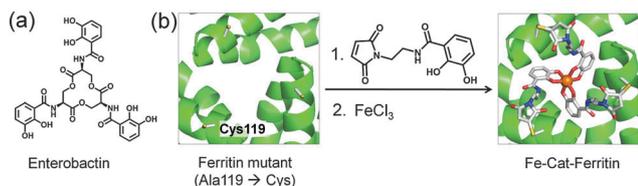


Fig. 5 (a) Molecular structure of Enterobactin. (b) Construction of an enterobactin analogue at the 3-fold channel of the apo-ferritin cage.

Fe ions into the cage. Such C_3 axis channels are engineered to introduce a catecholamide derivative, *N*-(2-(2,5-dioxo-1*H*-pyrrol-1-yl)ethyl)-2,3-dihydroxybenzamide to mimic the enterobactin, a siderophore of *Escherichia coli* which is known as the strongest chelator of ferric ions (Fig. 5).⁵⁰ In the presence of FeCl_3 , the modified ferritin exclusively produces the $\text{Fe}(\text{III})$ -tris-catecholato complex, as confirmed by the appearance of a characteristic LMCT band at 498 nm for the tris complex. The crystal structure analysis shows that $\text{Fe}(\text{III})$ forms an octahedral $\text{Fe}(\text{III})$ -tris-catecholato complex at the entrance of the 3-fold axis channel and blocks access of other molecules into the interior cavity. This work has broadened the scope of use of the ferritin cage for developing new biomimetic materials.

2.3 Organic molecules in the ferritin cage

Several medicinally important organic molecules were incorporated into the ferritin cage to improve pharmacokinetics, activity and selectivity. Doxorubicin (Dox) was incorporated into the cage using a process involving disassembly in 8 M urea solution followed by reassembly after gradual removal of urea.⁶³ The cage can accommodate 33 Dox molecules per apo-Fr without changing its overall structure and its size (12 nm) is considered to be optimal for an anticancer nano-medical agent to overcome the physiological barrier. Typical nanoparticles with sizes of about 100 nm are considered too large. The Dox loaded apo-Fr has excellent biocompatibility *in vivo*. The composite selectively delivers Dox molecules into tumor cells overexpressing TfR1 and does not require any additional functionalization to target cancer cells. The uptake of Dox-loaded ferritin in tumor cells was found to be 10-fold higher relative to free Dox and the composite has excellent antitumor activity compared to the clinically approved drug Doxil. In another study, the surface of the ferritin cage was modified with the RGD-4C (Cys-Asp-Cys-Arg-Gly-Asp-Cys-Phe-Cys) peptide which selectively binds to integrin $\alpha_v\beta_3$, a tumor angiogenesis gene which is overexpressed in tumor cells.⁶⁴ Dox molecules were pre-complexed with $\text{Cu}(\text{II})$ and loaded into the ferritin cage *via* passive diffusion with a loading capacity up to $\sim 73\%$ (w/w). Despite high loading, the ferritin cage structure was retained. The RGD modified ferritin cage loaded with Dox molecules is more effectively taken up by tumor cells due to the interaction between RGD-4C and integrins $\alpha_v\beta_3$ in U87MG cells. The Dox in the apo-Fr cage was released gradually and later identified after a long incubation time predominately in the nuclei where it interacts with DNA to induce cell death. The activity was found to be greater than that of free Dox. Similarly, an organic dye,

methylene blue, was incorporated into the ferritin cage *via* an assembly/disassembly route for developing light-activated anti-cancer agents.⁶⁶ In the presence of red light, the dye-encapsulated ferritin composite produces singlet oxygen which induces a cytotoxic effect in MCF-7 cells.

Besides protein cages, there are a number of synthetic cages such as metal-organic frameworks (MOFs), zeolites, and polymerosomes which are also available for controlling cellular functions.⁷⁵⁻⁷⁷ For example, engineered iron(III)-based metal-organic frameworks have been developed for drug delivery and imaging purposes.⁷⁸ Similarly, a polymeric nanoreactor encapsulates the enzyme penicillin acylase for local and controlled production of antibiotics.⁷⁹ The main advantage of the synthetic cages is that the cavity size can be easily tuned and also it can be functionalized for specific targeting ligands. However, the aqueous solubility, stability and biocompatibility remain an issue. In contrast, all such issues can be easily overcome by using the ferritin cage. The cage easily internalizes into cells. The preparation of the nanocomposites or metal functionalization using the ferritin cage can be achieved in a single step in buffer solution and does not require additional targeting or stabilizing ligands. Thus, the ferritin cage is a promising nano-platform for developing new biomaterials which are expected to provide the basis for new therapeutic and imaging applications.

3. Protein crystals

Protein crystals have been used in the determination of three-dimensional structures by X-ray crystallography since J. Kendrew first determined the three-dimensional protein structure of myoglobin in 1958.⁸⁰ In the last two decades, protein crystals have been generally regarded as porous materials because protein crystals have highly ordered arrangements of protein monomers and offer a wide range of pore sizes (20–100 Å), high porosity (50–80%), and large pore surface areas.^{5,81} However, because protein crystals tend to be very fragile in buffer solutions except for crystallization buffer, cross-linked enzyme crystals (CLECs) and cross-linked protein crystals (CLPCs) have been exploited to improve the mechanical stability of crystals for application as solid porous materials for separation, storage, and enzymatic reactions.^{81,102,103} Recently, these efforts have expanded the potential for using protein crystals as nanomaterials for functionalizing metal ions, metal complexes and metal nanoparticles as well as organic compounds (Table 2).⁵ In this section, we provide a survey of recent developments in functionalization of solvent channels of protein crystals for the construction of hybrid protein biomaterials.

3.1 Crystal engineering of hen egg white lysozyme

Among the engineered protein crystals, hen egg white lysozyme (HEWL) crystals are most frequently exploited as molecular templates for synthesis of metal nanoparticles and immobilization of metal complexes.¹⁰⁴ Because HEWL crystals are easily and abundantly obtained, we can investigate the properties and functions of the engineered crystals in detail as single crystals



Table 2 Recent functionalization of protein crystals

Protein	Composites	Methods	Functions	Ref.	
HEWL ^a	Au nanostructure	Soaking and chemical reduction (NaBH ₄)	Preparation of an Au nanostructure	82	
	Au NP	Soaking and autoreduction	Observation of the Au NP formation process	83	
	Au NP	Soaking and autoreduction	Catalyst: <i>p</i> -nitrophenol reduction	84	
	Au NP	Soaking and chemical reduction (NaOH)	Catalyst: <i>p</i> -nitrophenol reduction	85	
	Ag nanostructure	Soaking and photo reduction	Preparation of an Ag nanostructure	82	
	Ag NP	Soaking and chemical reduction (NaBH ₄)	Catalyst: <i>p</i> -nitrophenol reduction	86	
	CoPt NP	Soaking and chemical reduction (NaBH ₄)	Magnetism	87	
	Polypyrrole	Soaking and polymerization	Preparation of polypyrrole	88	
	Carbon dots	Soaking	Luminescence	89	
	CdS	Soaking	Fluorescence	90	
	Rh ³⁺	Soaking	Observation of the metal accumulation process	91	
		[Ru(benzene)Cl ₂] ₂	Soaking	Catalyst: transfer hydrogenation	92
		[Ru(CO) ₃ Cl ₂] ₂	Soaking	Extracellular matrix for CO releasing	93
Myoglobin	ZnP, Ru ₃ O	Heme substitution, soaking	Electron transfer system	94	
	Ru complexes, fluorescent dye	Chemical modification before crystallization	Porous modification	95	
Ferritin	Pd ion	Immobilization	Observation of the metal accumulation process	70	
	Zn, organic ligand	Co-crystallization	Synthesis of a porous crystalline framework	96	
	Dendrimer	Co-crystallization	Control crystal lattice structures	97	
	Phthalocyanine and pyrene tetrasulfonic acid	Co-crystallization	¹ O ₂ generation	98	
KDPGal ^b and Fkpa Protein	—	Fusion proteins by the genetic method	Porous crystal	99	
CCMV ^c	Avidin	Co-crystallization or soaking	Immobilization of functional molecules such as enzymes and Au NPs	100	
Plastocyanin	PEG	Chemical modification	Structure characterization	101	

^a HEWL = hen egg white lysozyme. ^b KDPGal = 2-keto-3-deoxy-6-phosphogalactonate. ^c CCMV = cowpea chlorotic mottle virus.

and in bulk. It has been reported that various metal ions and metal complexes are accumulated by coordination of amino acid residues on the surfaces of the solvent channels.^{91,105,106}

In addition, we can obtain different polymorphic structures of HEWL crystals by changing crystallization conditions. Such polymorphic structures include tetragonal, orthorhombic and monoclinic structures (T-, O-, and M-HEWL), which have different pore sizes, shapes, and arrangements of amino acid residues on the channels (Fig. 6).^{107–109} Therefore, HEWL crystals are one of the most promising crystals used as solid materials.

3.1.1. Preparation of inorganic nanoparticles. Mann *et al.* reported the first instance of using tetragonal HEWL crystals (*P*₄₃₂₁²) (T-HEWL crystals) in synthesis of metallic nanofilaments within the 1D solvent channels with a pore diameter of 1 to 2.2 nm. Glutaraldehyde cross-linked T-HEWL (CL-T-HEWL) crystals were employed for the formation of plasmonic Au and Ag nano-filaments within the channels (Fig. 7a–c).⁸² CL-T-HEWL crystals were soaked in a buffer solution containing NaBH₄. Au metallization was performed by soaking of dried CL-T-HEWL crystals containing borohydride in the solution including HAuCl₄·3H₂O. Ag nanorods were prepared by soaking Ag⁺ ions into the CL-T-HEWL crystals and then photoreduction of the Ag⁺ ion-containing CL-T-HEWL with UV radiation at a wavelength of 254 nm. TEM images of Au-T-HEWL crystals and Ag-T-HEWL crystals have electron dense stripes which were identified as metallic nanoparticles with average thicknesses of 1.6 nm for Au-T-HEWL and 1.5 nm for Ag-T-HEWL.

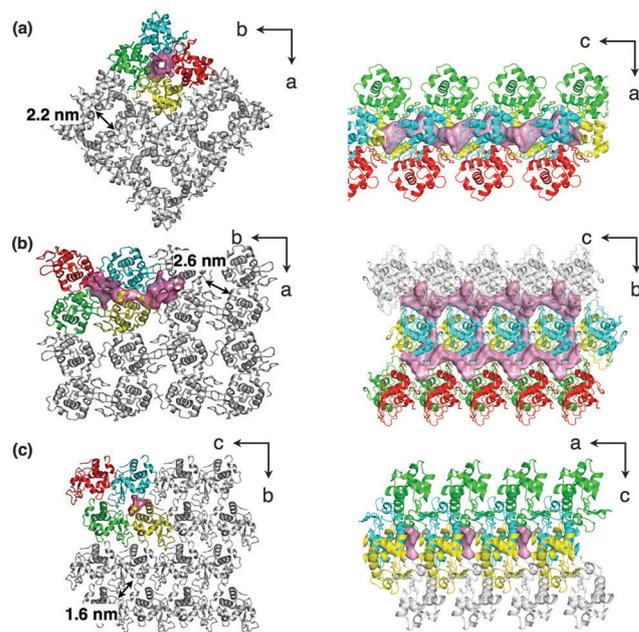
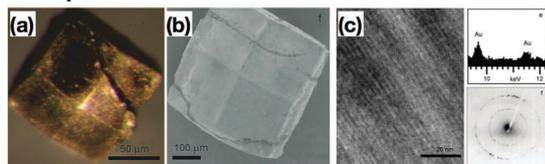


Fig. 6 Crystal lattice structures and the major solvent channels of (a) tetragonal (T)-HEWL, (b) orthorhombic (O)-HEWL, and (c) monoclinic (M)-HEWL taken from the PDB codes of 103L, 1BGI, and 5LYM, respectively.

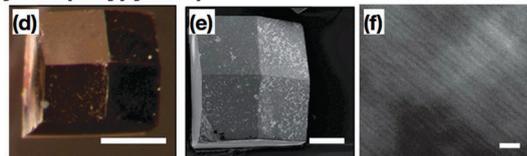
Synthesis of polypyrrole (PPy) in CL-T-HEWL crystals was also accomplished (Fig. 7d–f).⁸⁸ CL-T-HEWL crystals were soaked in the aqueous solution containing ammonium persulfate (APS) as



Au nanoparticles



Polymer (Polypyrrole)



Carbon dots (CD)



CD-CL-HEWL CD-CL-HEWL
oxCD-CL-HEWL oxCD-CL-HEWL
CL-HEWL CL-HEWL

Fig. 7 (a) Optical and (b) SEM micrographs of cross-linked lysozyme crystals containing chemically reduced Au nanoparticles. (c) TEM image of the thin fragments and corresponding EDX analysis and electron diffraction data. (d) Optical and (e) SEM micrographs of polypyrrole-cross-linked lysozyme crystals (PPy-CL-HEWL). (f) TEM image of the thin fragments of PPy-CL-HEWL. (g) Optical and (h) fluorescence micrographs of CL-HEWL (right), oxCD-CL-HEWL (middle), and CD-CL-HEWL (left). (i) TEM image of the thin fragments of CD-CL-HEWL. Reproduced from ref. 82, 88 and 89, with permission for (a–c), (d–f), and (g–i) from Wiley-VCH, the Royal Society of Chemistry and Wiley-VCH, respectively.

an oxidant for pyrrole. Then the APS-CL-T-HEWL crystals were exposed to pyrrole vapor for synthesis of PPy in the solvent channels of the crystals. TEM images of the PPy-CL-T-HEWL crystals showed continuous striations with a thickness of 1.8 nm, which they attributed to PPy domains because the striation images were not observed without polymerization of pyrrole. PPy-CL-T-HEWL crystals have low conductivity with a resistivity of $10^4 \Omega \text{ m}$, although CL-T-HEWL crystals have high resistivity ($>10^7 \Omega \text{ m}$).

Graphitic nanoparticles (carbon dots, CDs) were synthesized and confined within the solvent channels of the CL-T-HEWL crystals (Fig. 7g–i).⁸⁹ CL-T-HEWL crystals were soaked in an aqueous solution including citric acid and ethylene diamine and then exposed to microwave radiation to form oxygen-containing graphite nanoparticles (oxCD-CL-T-HEWL) by inducing pyrolysis of incorporated molecules. Carbon dots within the crystals (CD-CL-T-HEWL) were synthesized by incubation of the oxCD-CL-T-HEWL in an aqueous solution of sodium borohydride (NaBH_4). Fluorescence microscopy of CL-T-HEWL, oxCD-CL-T-HEWL and CD-CL-T-HEWL crystals indicated that the fluorescence intensities of the CD-CL-T-HEWL and oxCD-CL-T-HEWL crystals are dramatically increased and decreased, respectively, compared to the CL-T-HEWL crystal. It was found that fluorescence properties could be tuned by the addition of guest

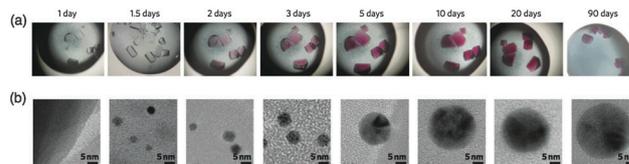


Fig. 8 Time-dependent growth of Au NPs in the single crystal of HEWL. (a) Optical images of single crystals of HEWL with Au(I) ions on different days of growth. (b) Corresponding TEM images. Reproduced from ref. 83 with permission for (a and b) from Nature publishing group.

molecules or an auxiliary chromophore at acidic or neutral pH, because pH-induced donor–acceptor coupled excitation energy transfer provides white or green light-emitting crystals.

Yi Lu *et al.* have reported time-dependent formation of gold nanoparticles (Au NPs) using a single crystal of T-HEWL (Fig. 8a and b).⁸³ The process of formation of Au NPs within the T-HEWL crystals was elucidated by auto-reduction of T-HEWL accumulating $\text{ClAuS}(\text{CH}_2\text{CH}_2\text{OH})_2$ (Au(I)) using transmission electron microscopy (TEM) and X-ray crystallography. Because the rate of auto-reduction of Au(I) is slow, it is possible to observe the process of formation of Au NPs. The time-dependent observation of Au NP formation could provide a means to elucidate biomineralization mechanisms and metal cluster formation mechanisms on the surfaces of proteins. Catalytic reactions for reduction of *p*-nitrophenol were also evaluated using Au NPs synthesized in the T-HEWL crystals with NaBH_4 .⁸⁴ Smaller Au NPs (2.2 nm) exhibited higher catalytic activities compared to the T-HEWL crystals with larger Au NPs ($>10 \text{ nm}$).

Fluorescent quantum dots were prepared in the single HEWL crystals by co-crystallization of lysozyme and Cd(II) ions in the presence of sodium sulfide.⁹⁰ The hybrid crystals were found to emit red fluorescence with a peak centered at $\sim 700 \text{ nm}$ under fluorescein isothiocyanate illumination. In the absence of sodium sulfide, the crystals containing Cd(II) ions are colorless and non-fluorescent, indicating that quantum dots of CdS are formed in the single crystal of lysozyme. In addition, when the CdS was prepared in aqueous solution in the presence of the lysozyme monomer, a low level of red fluorescence was observed, and red fluorescence emission was found to disappear after dissolution of the hybrid crystals containing CdS. These results show that only preparation of CdS inside the crystals exhibits enhancement of fluorescence due to the unique environments provided by the solvent channels of the crystals.

Ueno *et al.* synthesized magnetic CoPt NPs within the solvent channels of HEWL crystals with different sizes and compositions using three different crystal systems of HEWL (Fig. 6 and 9).⁸⁷ T-HEWL, O-HEWL and M-HEWL crystals were cross-linked with glutaraldehyde to maintain the crystal lattice during synthesis of CoPt-NPs. Cross-linked HEWL crystals (CL-O-HEWL, CL-T-HEWL, and CL-M-HEWL) were soaked in buffer solution containing Co(II) and Pt(II) ions. The CoPt NPs were synthesized by reduction of the CL-HEWL crystals containing Co and Pt ions with sodium borohydride. The average particle sizes of CoPt NPs synthesized in CL-HEWL crystals were



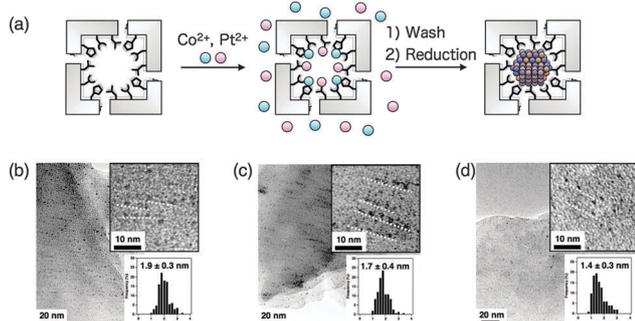


Fig. 9 (a) Schematic drawing of the preparation of CoPt-NPs in a solvent channel of a HEWL crystal. TEM images and particle size distribution of (b) CoPt-CL-O-HEWL, (c) CoPt-CL-T-HEWL, and (d) CoPt-CL-M-HEWL. The black dots are CoPt NPs and the white dotted lines (insets) provide a guide.

determined using TEM. Alignment of CoPt NPs in the solvent channels of the CL-HEWL crystals was observed (Fig. 9b–d). The composition of Co and Pt in CoPt NPs was analyzed by determining the X-ray fluorescence of CoPt NPs in CL-O-HEWL, CL-T-HEWL and CL-M-HEWL crystals. The results show that the ratios of Co and Pt atoms in CoPt NPs are 7.7:92.3, 3.8:96.2, and 6.3:93.7, respectively. The composition of CoPt NPs in the CL-HEWL would be affected by pre-organization of Co(II) and Pt(II) ions bound on the surface of the solvent channels of the HEWL crystals. The magnetic properties of CoPt-CL-HEWL crystals were examined using a superconducting quantum interference device (SQUID) magnetometer. CoPt-CL-O-HEWL has the highest coercivity value (4600 Oe) among the three CoPt-CL-HEWL crystals (with 1600 and 2900 Oe for CoPt-CL-T-HEWL and CoPt-CL-M-HEWL, respectively). The order of the coercivity of CoPt NPs within the crystals is generally proportional to the content of Co atoms of the CoPt NPs as determined in the previous report. These results indicate that the physical properties of inorganic metal nanoparticles prepared in the solvent channels can be tuned by the crystal lattice.

3.1.2. Functionalization of organometallic complexes. In another effort, organometallic complexes were accumulated in the solvent channels of the CL-HEWL crystal in an effort to develop catalytic reaction vessels (Fig. 10).⁹² CL-T-HEWL and CL-O-HEWL crystals were soaked in buffer solution including $[\text{Ru}(\text{benzene})\text{Cl}_2]_2$ complexes to prepare a Ru(benzene)-CL-HEWL composite. Catalytic activities of Ru(benzene)-CL-T-HEWL and Ru(benzene)-CL-O-HEWL crystals were investigated for transfer hydrogenation of acetophenone derivatives (Fig. 10). Ru(benzene)-CL-T-HEWL exhibits the highest activity for propiophenone with 53% conversion. Ru(benzene)-CL-O-HEWL shows higher enantioselectivity than Ru(benzene)-CL-T-HEWL for all products. The highest ee value is 36% ee S for isobutyrophenone. Interestingly, when only Ru complexes were employed for the catalytic reaction in buffer solution, it was found that the complexes could not catalyze transfer hydrogenation. In addition, the enantioselectivity and configuration of the hydrogenation products could be controlled by interactions of the substrate with the active sites of Ru complexes in the solvent channels, which differ among the crystal lattices (Fig. 10b–e).

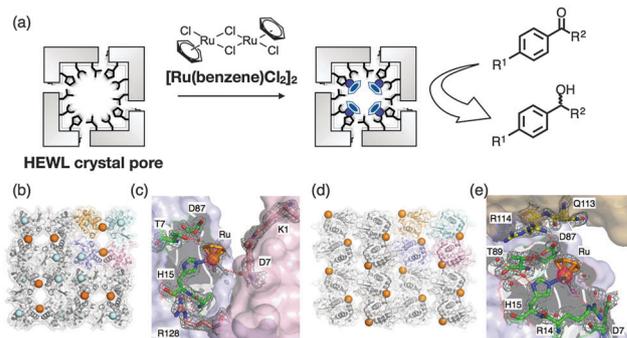


Fig. 10 (a) Schematic representation of the preparation of Ru(benzene)-CL-HEWL and catalytic transfer hydrogenation. Crystal structures in a lattice structure of (b) Ru(benzene)-CL-T-HEWL and (d) Ru(benzene)-CL-O-HEWL and active site structures of (c) Ru(benzene)-CL-T-HEWL and (e) Ru(benzene)-CL-O-HEWL.

HEWL crystals immobilized with organometallic Ru(CO) complexes can function as an extracellular matrix (ECM) for CO release.⁹³ The $[\text{Ru}(\text{CO})_3\text{Cl}_2]_2$ complex (CORM-2) was immobilized in the solvent channels of CL-T-HEWL crystals in an effort to store and release CO gas by coordination of amino acid residues such as His, Asp and Lys as observed from the X-ray crystal structure of Ru(CO)-CL-T-HEWL (Fig. 11). The number of Ru complexes in the crystals was found to be 10 per HEWL monomer. The IR spectrum of Ru(CO)-CL-T-HEWL exhibits four bands at 2055, 2025, 1981, and 1940 cm^{-1} . A pair of IR bands at 2055 and 1981 cm^{-1} was assigned to *cis*-Ru(CO)₂ complexes coordinated with His15. The other pair of bands at 2025 and 1940 cm^{-1} was expected for *cis*-Ru(CO)₂ complexes ligated to carboxylate of an Asp residue. The CO release phenomenon was examined in a myoglobin assay based on conversion of deoxy-myoglobin to carbonyl-myoglobin. The half life ($t_{1/2}$) value of CO release from Ru(CO)-CL-T-HEWL was found to be increased 10-fold longer than CORM-2. The amount of CO released from Ru(CO)-CL-T-HEWL was found to be 0.38 per Ru. The NF- κ B activity of Ru(CO)-CL-T-HEWL in the

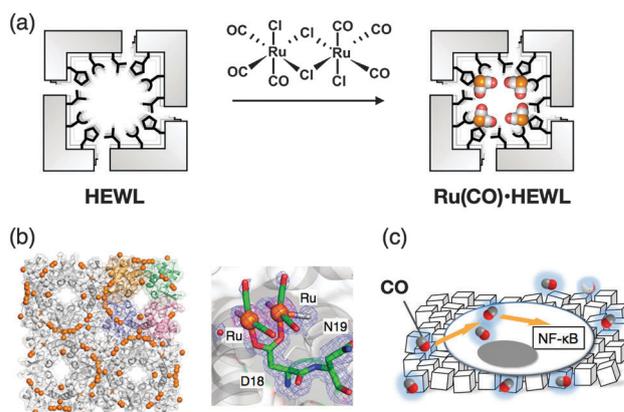


Fig. 11 (a) Schematic representation of the preparation of Ru(CO)-CL-HEWL. (b) Crystal structure in a lattice structure and coordination structure in the vicinity of Asp18 of Ru(CO)-CL-HEWL. (c) CO release in the vicinity of cells and activation of NF- κ B.



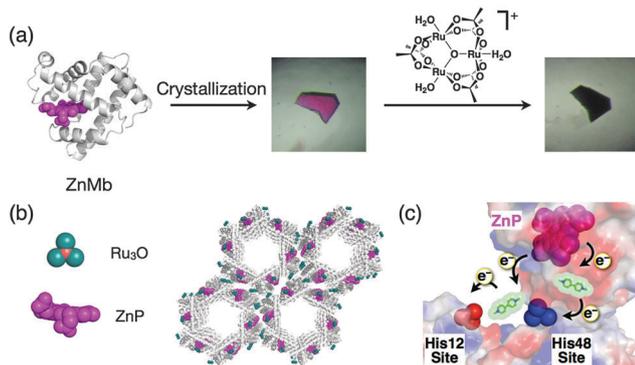


Fig. 12 (a) Construction of a photoinduced electron transfer system in the ZnP substituted Mb crystal. (b) Crystal structure of the Ru₃O soaked ZnMb crystal. (c) Schematic representation of methyl viologen mediated electron transfer between ZnP and Ru₃O in ZnMb crystals.

presence of TNF- α was found to be increased by 20% compared to the control.

3.2 Crystal engineering of myoglobin

Sperm whale myoglobin is crystallized with ammonium sulfate as a precipitant. The myoglobin crystal belongs to the space group *P6* and has hexagonal pores with a diameter of 40 Å. This pore can be employed as a reaction space and metal accommodation space within the crystals.⁹⁵ A Zn porphyrin (ZnP) and a Ru₃O cluster were found to be accumulated on the specific sites in the solvent channels of a single crystal of Mb by the replacement of heme and coordination of His48 and His12 to construct an artificial photo-induced electron transfer system (Fig. 12).⁹⁴ Methyl viologen (MV) soaked in the crystals was employed as a mediator between the ZnP and the Ru₃O cluster. This functionalized Mb crystal provides a longer half-life of the charge-separated state which is 2800-fold higher than that of the same system in organic solvent. This result suggests that the protein crystal can accumulate functional molecules at specific sites in the solvent channels to develop unique activity.

3.3 Crystal engineering of ferritin

Ferritin (Fr) is a spherical protein cage composed of 24 subunits with an interior space 8 nm in diameter. Apo-Fr has been utilized as a protein container for deposition of metal nanoparticles and for the incorporation of metal complexes.^{32,34} Recently, crystal engineering using ferritin has been developed to construct novel assembly structures. Ueno and co-workers elucidated the mechanism for accumulation of Pd ions on the interior surface of recombinant horse L-chain apo-ferritin (apo-rHLFr) using X-ray crystal structural analysis of a series of apo-rHLFr containing Pd ions (Fig. 13a).⁷⁰ Apo-rHLFr containing different amounts of Pd ions (50, 100, and 200 equivalences *vs.* apo-rHLFr) was crystallized. X-ray crystal structures of these Pd-apo-rHLFr composites show that Pd ions are deposited at specific binding sites located at the accumulation sites and 3-fold channels. At the accumulation center, Pd ions are selectively bound to Cys48 and His49 with several different binding modes at an early stage of Pd binding. The conformational

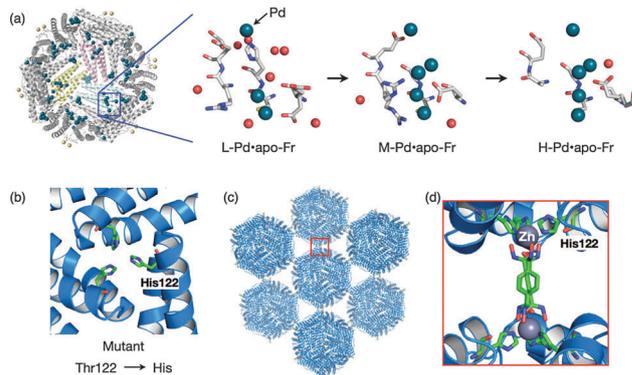


Fig. 13 (a) Accumulation process of Pd ions on the interior surface of apo-rHLFr. (b) Structure of the 3-fold channel of the ferritin mutant. (c) The bcc packing of the bdh²⁻ Zn-ferritin lattice mediated by coordination of bdh²⁻ with Zn bound at 3-fold channels. (d) Closeup view of the interface of two ferritin molecules, bdh²⁻ ligand bridge ferritin molecules. The structure was taken from PDB code 5CMR.

changes of His49, Glu45 and Arg52 stabilize and promote Pd(II) binding at the accumulation center. These interactions would promote the natural accumulation reactions such as biomineralization and metal cluster formation in protein scaffolds.

Tezcan and co-workers have established the construction of protein supramolecules from 1D protein nanotubes to 3D protein crystals by metal-mediated protein assemblies using engineered cytochrome *cb*₅₆₂ and ferritin.^{110,111} A 3D porous crystalline framework has been constructed recently by using the metal-directed protein assembly using ferritin, which was engineered at the 3-fold channel (Fig. 13b–d).⁹⁶ Thr122 in the exterior of the 3-fold axis channel was replaced with His to introduce a Zn coordination site. In addition, original cysteine residues of Cys90, Cys102, and Cys130 were replaced with either Glu or Ala to prevent Zn binding at these sites. 3D porous frameworks are formed by coordination of Zn²⁺ with the mutated ferritin in the presence of the bidentate ligand, benzene-1,4-dihydroxamic acid (H₂bdh) as well as metal organic frameworks (MOFs) which are composed of metal ions and organic ligands. The crystal structure of the framework has a different crystal lattice structure (*I432*) from the crystal produced in the presence of only ferritin and Zn ions (*F432*), and ferritin molecules were found to be connected by coordination of bidentate ligands to Zn ions (Fig. 13c and d). The self-assembling structure with metal coordination provides highly porous frameworks with a solvent content of 67%. This work indicates an opportunity to generate template proteins based on crystalline biomaterials for useful applications.

Kostiainen and co-workers have recently reported functionalization of a ferritin crystal by co-crystallization of dendrimers or organic dyes.^{97,98} The electrostatic self-assembly of apo-Fr and poly(amidoamine) (PAMAM) dendrimers provides two-component crystalline materials. The crystalline lattice structures of the co-crystals are affected by ionic strength and dendrimer generation. Two crystal symmetries (fcc and hcp) are allowed using apo-Fr and the dendrimer. The lattice structures also



depend on the size of the dendrimer and the ratio of the dendrimer and apo-Fr.⁷⁷ The organic dyes were also co-crystallized with apo-Fr. The supramolecular complex formed between cationic Zn phthalocyanine and anionic pyrene acts as molecular glue to crystallize apo-Fr *via* electrostatic interactions. The small-angle X-ray scattering (SAXS) analysis of the crystal shows the face-centered cubic (fcc) structure. The adsorption spectrum of the hybrid crystals indicates that the complex of phthalocyanine and pyrene is maintained in the crystals. The hybrid crystals can generate highly reactive ¹O₂ which is useful for various applications such as photodynamic therapy, water treatments, diagnostic arrays and as an oxidant in organic synthesis.⁹⁸

These results indicate that apo-Fr is useful for engineering crystals as well as *in vivo* applications due to its high stability. The engineered Fr crystals can be used to create functional solid materials because ferritin crystals can utilize both the interior of the cage and the solvent channels of the crystals.

3.4 Crystal engineering of other proteins

Artificial protein crystals have been designed by introducing and modifying functional moieties at the interface of proteins, such as fusion of two protein domains, which naturally form oligomeric proteins for the formation of novel protein crystals as reported in our recent review.⁵ Co-crystallization of two different proteins provides heterogeneous crystalline assemblies integrating different functions into nanostructures.¹⁰⁰ Protein-polymer conjugates were also crystallized and the protein-polymer interactions were investigated.¹⁰¹

Yeates and co-workers have reported novel protein caged assemblies of 750 kDa with a large interior space of about 130 Å by fusion of trimeric *Escherichia coli* 2-keto-3-deoxy-6-phosphogalactonate (KDPGal) aldolase and dimeric N-terminal domain *E. coli* FkpA protein (Fig. 14a–c).⁹⁹ The fusion protein is composed of KDPGal aldolase at the N-terminal domain, the N-terminal domain of FkpA at the C-terminal domain and a four residue alpha helical linker. The fusion protein designed by computational modeling was crystallized after a prolonged incubation time. The crystal structure showed that the atomic structure is in agreement with the modeled structure within a 1.2 Å root-mean-square deviation. In the crystal, large voids are formed with high solvent content (~82%) as well as porous organic molecules and metal organic frameworks. These specifically designed highly porous protein crystals can be used to immobilize large molecules such as proteins for X-ray crystal structure analysis.

Kostiainen *et al.* constructed virus crystals including different functional materials, such as proteins, dendrimers, and metal nanoparticles.¹⁰⁰ Cowpea chlorotic mottle virus (CCMV) was crystallized in the presence of avidin, to generate an electrostatically assembled composite with a non-close-packed body centered cubic structure (Fig. 14d). The multicomponent crystal was modified with biotin-conjugated fluorescein, horseradish peroxidase, and gold nanoparticles to incorporate these functional molecules into the crystals using avidin-biotin interactions. The multicomponent crystals were also prepared by co-crystallization of CCMV and pre-modified avidin with

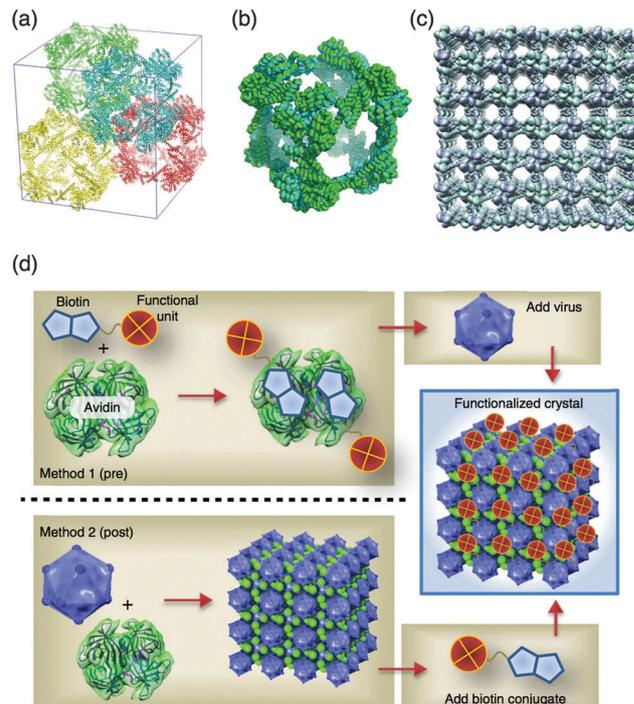


Fig. 14 (a–c) Crystal structure of the fusion protein of aldolase-three-helix cubic protein. (a) Four complete cages of the ATC-HL3 cage are packed within a unit cell. (b) The cube-shaped cage in the crystal. (c) The packing alignment of cages in the crystal produces a highly porous protein lattice. (d) Pre- and post-functionalization of CCMV-avidin crystals through biotin-avidin interaction. Reproduced from ref. 99 and 100 with permission from Nature publishing group.

biotinylated functional molecules. The functionalized crystals have 2 orders of magnitude higher enzymatic activity ($0.13\text{--}0.15\ \mu\text{M s}^{-1}$) than the corresponding free enzymes in supernatant solution which are obtained after washing crystals ($\sim 0.007\ \mu\text{M s}^{-1}$). Furthermore, only crystalline avidin-HRP CCMV crystals show high enzymatic activity compared to amorphous HRP without CCMV. These results show that active enzymes are accumulated into the crystals and that this method can be applied to several biotinylated molecules for applications in preparation of crystalline materials having catalytic, optical and magnetic functions.

Crowley *et al.* reported structural characterization of the β -sheet protein plastocyanin (Pc) conjugated with a single poly(ethylene glycol) (PEG) 5000 molecule by size exclusion chromatography (SEC), NMR spectroscopy, and X-ray crystallography.¹⁰¹ The crystal structure refined at 4.2 Å resolution has a highly porous structure with high solvent content (80%), in which the PEGylated proteins form double-helical assemblies, although the electron densities corresponding to the PEG chains are not observed due to the disorder. The volume of the PEG domain in the crystals was estimated to be within 10% of the calculated random coil PEG 5000. The NMR studies show that the PEG domains are not influenced by the proteins, suggesting that the PEG domains have minimal protein interactions.

These engineered protein crystals with novel crystal assembling structures have significant potential to provide large spaces within



crystals in the development of applications in solid functional materials to accumulate large molecules such as enzymes and proteins.

4. *In vivo* three dimensional protein crystals

As mentioned in the previous section, there is growing interest in functionalization of protein crystals. However, it remains challenging to establish large scale or more conventional systems for tailoring protein crystals with various functions. It is well known that certain proteins can be crystallized in living cells, such as peroxidase in peroxisomes, insulin in secretory granules, and polyhedrin crystals for generating functional solid-state catalysts, generating storage vessels, and encapsulating viruses, respectively.^{112–114} These protein crystals are spontaneously formed in living cells with high concentrations of the proteins in restricted locations. These crystals tend to be quite small compared to crystals prepared *in vitro* because growth of protein crystallization is limited to the size of living cells. Therefore, *in vivo* protein crystals have not been used for X-ray crystal structural analysis until recently due to radiation damage and difficulties in focusing on microcrystals.¹¹⁵ Enhanced techniques in micro-X-ray crystallography allow us to solve these problems by using microfocus X-ray beamlines and X-ray free-electron lasers.¹¹⁵ In 2007, Coulibaly *et al.* reported the first crystal structure of the *in vivo* protein crystal polyhedra, which forms in silkworms infected by cypoviruses (cytoplasmic polyhedrosis virus, CPV).¹¹⁶ Several *in vivo* protein crystals, including polyhedra and cathepsin B from *Trypanosoma brucei* (CatB), are produced in insect cells (Fig. 15).^{114,116,117} Structures of insecticidal Cry3A toxin in *Bacillus thuringiensis*, viral spindles, and coral fluorescent protein have been determined by microcrystal X-ray crystal structure analysis.^{114,118–120} In this section, recent progress in *in vivo* protein crystallization and *in vivo* crystal engineering toward the development of functional solid materials is described.

4.1 *In vivo* protein crystallization

The crystal structures of several *in vivo* protein crystals have recently been determined using synchrotron X-ray microbeam- and X-ray free-electron laser (XFEL)-based

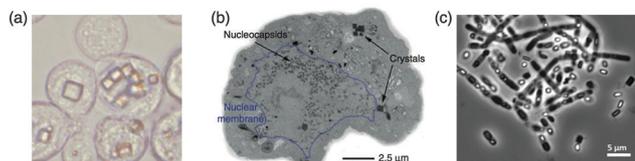


Fig. 15 *In vivo* three dimensional protein crystals. (a) Polyhedral crystals produced in Sf21 insect cells. (b) Scanning EM image of Sf9 insect cells infected with cathepsin B from *Trypanosoma brucei* (CatB) virus. (c) Phase contrast light micrograph of Bt cells. The dark rectangular shapes correspond to Cry3A toxin crystals. Reproduced from ref. 117 with permission for (b) from Nature publishing group. (c) was reproduced from ref. 118. Copyright (2014) National Academy of Science, USA.

crystallography techniques.^{114–116,118–120} Coulibaly *et al.* reported the crystal structures of polyhedral crystals (PhCs), which are crystals of polyhedrin monomers (PhMs) produced in insect cells infected by cypoviruses (CPVs) and nuclear polyhedrosis viruses (NPVs).^{116,121} During crystallization in the cell, polyhedra embed the virus particles. The viruses embedded in PhCs are protected against harsh conditions. The crystal structure of CPV PhCs shows that a trimer of PhMs tightly organized crystalline structures, in which the H1-helices are located at the N-terminus of PhMs, interact with neighboring PhMs. The dense pack of trimer structures forms narrow solvent channels in the crystals (Fig. 16a–c). The structures of wild-type and recombinant NPV PhCs were also determined by X-ray crystallography.^{121,122} The recombinant structure shows that molecular organization of NPV polyhedrin is different from CPV polyhedrin. The assembling structures of baculovirus PhCs are stabilized by disulfide bonds and domain-swapping of N-terminal regions (Fig. 16d–f). Therefore, the N-terminal domains play different roles in CPV and NPV PhCs for stabilization of the crystalline assembly.

Coulibaly *et al.* also reported a crystal structure of the spindle produced by insect poxviruses. Spindles are crystalline assemblies of fusolin protein that enhance the virulence of the viruses and the insecticidal activity of unrelated pathogens.¹¹⁹ To understand the mechanism of enhancement of the virulence and assembly of spindles for ultra-stable crystalline materials, the spindle structure was determined by X-ray crystallography. The crystal structure of spindle from *Melolontha melolontha* EV (MMEV) shows that the spindles are stabilized by cross-linking of the C-terminal molecular arm of fusolin as well as the H1 helix of CPV PhCs and disulfide bonds between fusolin dimers to form the 3D network through the entire group of crystalline assemblies of spindles.

X-ray free-electron lasers (XFELs) are also utilized for crystal structure analysis of *in vivo* protein crystals. The crystal structure of CatB grown in insect cells was determined using XFELs and

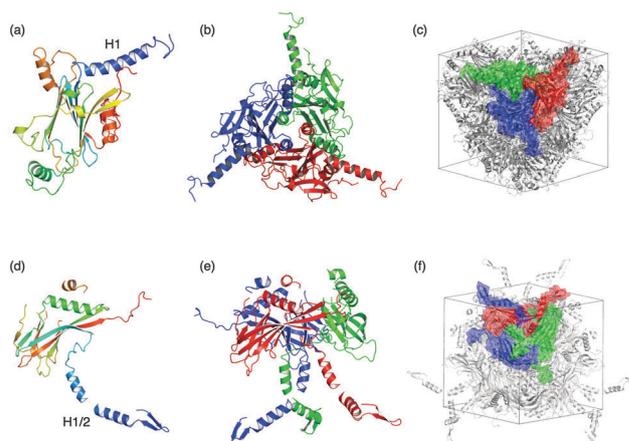


Fig. 16 Crystal structures of CPV polyhedra (a–c) and NPV polyhedra (d–f). Monomer (a and d) and trimer (b and e), and molecular assembling of the polyhedrin trimer in a unit cell (c and f) of CPV PhCs, and NPV PhCs taken from PDB ID: 2OH6, 3JVB, respectively.



refined to 2.1 Å.¹¹⁴ The native structure of CatB includes the pro-peptide and carbohydrate, which were not observed in the crystals of recrystallized CatB. This crystal structure analysis of *in vivo* protein crystals has potential for use in estimating the *in vivo* dynamic reactions of proteins, such as glycosylation, and phosphorylation, among others.

Very recently, crystal structure analysis of *in vivo* protein crystals within cells has been developed using microfocus beamlines with synchrotron and X-ray free-electron laser beam sources.^{118,120,123} Whole cells bearing CPV PhCs and engineered fluorescent protein variants were spread on a mesh loop, cooled, and irradiated at X-ray microfocus beamlines.^{120,123} The structure of CPV PhCs collected within cells indicates that there is no significant difference from isolated crystals. Sawaya and co-workers reported a crystal structure of Cry3A toxin produced in *Bacillus thuringiensis* (Bt) determined in an *in vivo* diffraction study using XFELs.¹¹⁸ The structure of Cry3A existing in living cells was determined to a resolution of 2.9 Å. The method of *in vivo* diffraction data collection does not require purification of the *in vivo* protein crystals and thus avoids destabilization of the crystals that can occur when the crystals are isolated from cells. In the future, *in vivo* chemical reactions could be analyzed by *in vivo* X-ray crystallography of *in vivo* protein crystals within living cells.

Miyawaki *et al.* have recently reported *in vivo* crystallization of a variant of coral fluorescent protein, Kikume Green-Red (KikGR), in HEK293 cells.¹²⁰ The KikGR variant was transfected into HEK293 and the crystallization process was observed using confocal microscopy. The time-dependent observation indicated that the crystals grew to micro-sized crystalline assemblies over the course of several minutes in living cells after 2–5 days of transfection (Fig. 17). TEM images and biochemical experiments indicate that the KikGR variant is crystallized within the cytoplasm and then the crystal is incorporated into the lysosome as a result of an autophagic process.

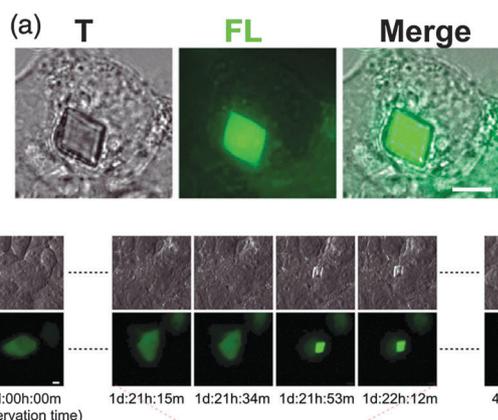


Fig. 17 (a) Transmission (T), fluorescence (FL) and merged images of the HEK293 cell bearing a crystal of a coral fluorescent protein, KikGR, variant. (b) Crystallization process of the KikGR variant revealed by long-term imaging. Differential interference contrast (DIC) and fluorescence (FL) images of HEK cells at the indicated times. Reproduced from ref. 120 with permission from Cell Press.

4.2 *In vivo* crystal engineering

The most prominent example of generation of *in vivo* protein crystals is the polyhedra produced in baculovirus-infected insect cells. CPV produces a protein known as polyhedrin in the replication cycle in insect cells to embed the viruses in spheres with a diameter of 75 nm in the crystals. The virus particles embedded in PhCs are resistant to dehydration, freezing and enzymatic degradation for several years because PhCs can tolerate harsh chemical treatments including incubation in concentrated urea, acid and detergents.¹¹⁶ Therefore, a PhC is a candidate for development as a crystalline solid material because the PhC is easily and abundantly crystallized in insect cells and has high stability.

Mori *et al.* reported encapsulation of exogenous proteins into recombinant PhCs instead of CPV by co-expression of the exogenous protein and PhM.^{124,125} The H1, N-terminal fragment of the PhM can function as a polyhedrin recognition signaling moiety which leads to the incorporation of exogenous proteins or enzymes because the H1-helix tightly interacts with neighboring PhMs as well as VP3 of CPV turret protein (Fig. 16a–c).^{124,125} Fusion proteins consisting of enhanced fluorescent protein (EGFP) fused with the H1-helix fragment of the PhM (H1-EGFP) or VP3-fragment of CPV (EGFP-VP3) were co-infected to prepare the composites of PhCs which include H1-EGFP or EGFP-VP3 (Fig. 18). Confocal laser scanning microscopy of the crystals indicates that the green fluorescence of EGFPs is distributed more close to the surface than to the core of the crystals. In addition, EGFP and *Discosoma* sp. red fluorescent protein (DsRed) are co-immobilized in one PhC by using the H1 and VP3 fragments. This technique was applied to immobilize various cytokines, such as FGF, LIF, and EGF in PhCs.^{126,127} These immobilized cytokines retain their biological activities, which include promotion of fibroblast growth and proliferation of mouse embryonic and induced pluripotent stem cells by slow release of the immobilized proteins.

Ueno *et al.* reported one-pot preparation of PhCs with protein kinase C (PKC) β II, a phosphorylation enzyme, by co-expression of PKC β II and PhMs, which is mutated for dissolution at optimum pH and retention of PKC activity after

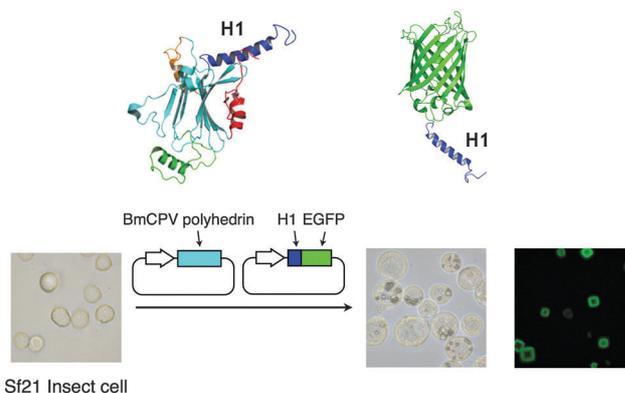


Fig. 18 Encapsulation of H1-EGFP into CPV PhC by co-infection and co-expression of CPV PhMs and H1-EGFP.



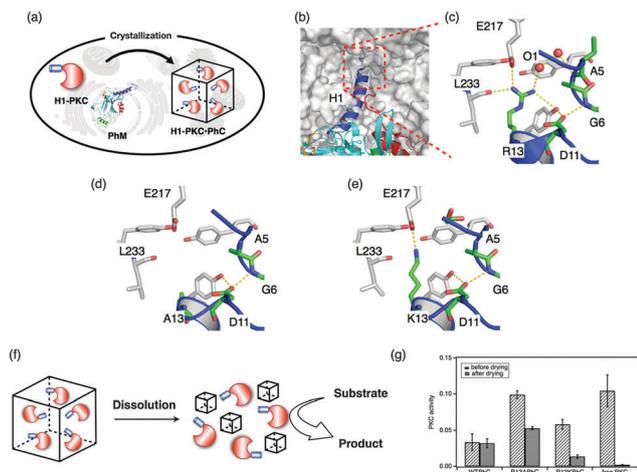


Fig. 19 Schematic representation of (a) *in vivo* preparation of H1-PKC-PhC. (b) The interaction of the H1 helix with neighboring polyhedrin molecules. Magnified structures in the vicinity position 13 of (c) WTPhC, (d) R13APhC, and (e) R13KPhC. Hydrogen bonds are indicated with yellow dotted lines. (f) Schematic representation of release of H1-PKC by dissolution and their enzymatic reaction. (g) PKC activity of H1-PKC-PhCs and recombinant free PKC β -II after air drying for 1 week at pH 8.5.

storage under drying. These modifications are expected to expand the application of the PhC as a solid material while the enzymes are retained at optimal pH (Fig. 19).¹²⁸ The crystal structure of WTPhCs indicates that the side chain of Arg13 at the tip of the H1-helix is expected to contribute to the robust character of the crystals by the formation of intermolecular hydrogen bonds. Two mutant crystals were designed, in which Arg13 is replaced with Ala and Lys, R13APhMs and R13KPhMs, respectively. H1-PKC-R13APhCs and H1-PKC-R13KPhCs both release large amounts of H1-PKC molecules at pH 8.5 upon dissolution of PhCs, although H1-PKC could not be released from WTPhCs. Phosphorylation reactions were examined to evaluate the enzymatic reactions. The stabilization of H1-PKC within the crystals was evaluated by determining enzymatic activity after incubation by air-drying for 1 week at 25 °C (Fig. 19g). These results indicate that H1-PKCs immobilized in the crystals retain the enzyme activity after incubation under harsh conditions, although free-PKC does not retain activity under the same conditions. These results show that exogenous enzymes with an H1 helix can be incorporated by co-expression into insect cells and the mutant crystals can release the H1-PKC at pH 8.5 by dissolution of the crystals. H1-PKC has higher tolerance to drying when incorporated into the crystals compared to free PKC.

It has been demonstrated that the surface of PhCs can be modified with carbohydrate (Fig. 20a).¹²⁹ Cys mutants, in which original Cys residues were replaced with Ala and one or two Cys residues, were introduced on the surface of the crystals. The introduced Cys residues were modified with propargyl maleimide and then the acetylene moieties were modified with Lewis X (LeX)-azide *via* the copper-catalyzed azide alkyne cycloaddition reaction. The reaction of LeX-modified PhCs with Alexa Fluor 488-conjugated antibody against stage-specific

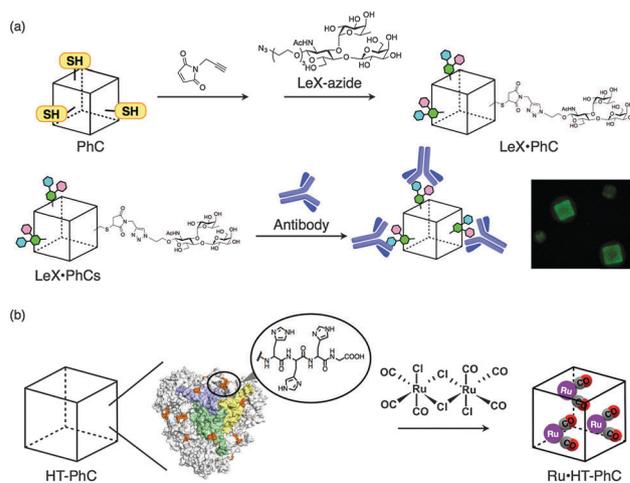


Fig. 20 (a) Schematic representation of surface modification of PhC with LeX moieties. The cystein residues were modified with propargyl maleimide then LeX-azide *via* thiol-maleimide chemistry and the copper-catalyzed azide alkyne cycloaddition, respectively. The LeX modified PhC was reacted with antibody. (b) Immobilization of Ru carbonyl complexes into PhC with a His-tag at the C-terminus.

embryonic antigen 1 (SSEA-1) was performed for 10 h at 4 °C. Confocal laser scanning microscopy of the composite shows that Cys-mutant PhCs react with the antibody more effectively than WTPhCs. The surface modification of protein crystals is expected to provide attractive materials for applications involving extracellular matrices, biosolid catalysis, and drug delivery systems.

A PhC also reacted with organometallic Ru carbonyl complexes in an effort to construct CO-releasing extracellular matrices (Fig. 20b).¹³⁰ Introduction of a hexahistidine tag (HT) to the C-terminus allows the crystal to accumulate Ru carbonyl complexes which are twice the size of WTPhCs, and release CO gas 3-fold more effectively than WTPhCs including Ru carbonyl complexes. The HT-PhC which includes Ru carbonyl complexes (Ru-HT-PhC) releases 6.2 ± 0.9 equiv. per PhM with a half-life ($t_{1/2}$) of 27.7 ± 1.6 min. The value is about 4-fold greater than that of the original CORM-2 (6.9 ± 2.8 min). The performance of a Ru-HTPhC as an extracellular scaffold was evaluated by measuring the activity of nuclear factor kappa B (NF- κ B) in living cells because CO gas is known to activate NF- κ B in the presence of tumor necrosis factor α (TNF- α). Ru-HTPhCs showed a 72% increase of NF- κ B activity compared to WTPhCs. The results indicate that the PhC can immobilize organometallic complexes to release the small molecules without degradation of the crystals. The hybrid crystals will be investigated for the development of solid functional materials for applications such as catalysis, metallodrugs, and bioimaging because organometallic complexes have various functions.

5. Conclusions and future perspectives

The design of protein assemblies to construct protein-based hybrid materials with metal ions, metal complexes, nanomaterials



and proteins has recently become a growing field with the aim of providing novel functions and mimicking natural functions in materials science and nanotechnology. This feature article summarizes recent development of strategies for functionalization of protein assemblies based on the ferritin cage and protein crystals for applications such as catalysis, magnetism, optics, imaging, and drug delivery both *in vivo* and *in vitro* systems. Protein assemblies provide confined environments for accumulation of functional molecules. Self-assembled protein cages containing metal complexes and inorganic materials have been recently utilized in cellular environments for delivery of CO as a biological gas and delivery of drugs due to their high biocompatibility and stability. Furthermore, protein cages have been employed as reaction vessels to incorporate multiple catalytic centers such as enzymes for cascade-type reactions in larger internal spaces functioning as natural micro-compartments. In the future, catalytic cascade reactions in cellular environments using protein assemblies incorporating metal complexes and enzymes will be investigated.

Protein crystals have also been recently developed not only in structural biology but also in materials science. Cross-linked protein crystals encapsulating metal ions, metal complexes and nanomaterials immobilized in solvent channels by coordination of amino acid residues and/or chemical modifications can function as solid biomaterials with catalytic, magnetic and optical properties. These efforts in protein crystal engineering have been expanded to *in vivo* protein crystallization. One of the advantages of using *in vivo* protein crystals as solid materials is that it is possible to prepare tailored protein crystals in large scale or conventional preparations because the crystals are spontaneously formed in living cells. Thus, design of confined chemical environments in protein cages and crystals has contributed to the development of new biohybrid materials.

Abbreviations

Hsp	Heat shock protein
Dps	DNA-binding protein for starved cells
CCMV	Cowpea chlorotic mottle virus
SOD	Superoxide dismutase
ROS	Reactive oxygen species
DCFH-DA	2,7-Dichlorofluorescein diacetate
TfR1	Transferrin receptor 1
TMB	3,3',5,5'-Tetramethylbenzidine
DAB	Di-aza-aminobenzene
QDs	Quantum dots
NIR	Near-infrared
MRI	Magnetic resonance imaging
RGD-4C	Cys-Asp-Cys-Arg-Gly-Asp-Cys-Phe-Cys peptide
PDT	Photodynamic therapy
CORM	Carbon monoxide releasing molecule
ATR	Attenuated total reflection
NF- κ B	Nuclear factor κ B
TNF- α	Tumor necrosis factor α
Dox	Doxorubicin

CLECs	Cross-linked enzyme crystals
CLPCs	Cross-linked protein crystals
HEWL	Hen egg white lysozyme
PPy	Polypyrrole
APS	Ammonium persulfate
CD	Carbon dot
NaBH ₄	Sodium borohydride
Au NPs	Gold nanoparticles
TEM	Transmission electron microscopy
SQUID	Superconducting quantum interference device
ECM	Extracellular matrix
ZnP	Zinc porphyrin
MV	Methyl viologen
H ₂ bdh	Benzene-1,4-dihydroxamic acid
MOF	Metal organic framework
PAMAM	Poly(amidoamine) dendrimer
SAXS	Small-angle X-ray scattering
Fcc	Face-centered cubic
KDPGal	2-Keto-3-deoxy-6-phosphogalactonate
CPV	Cytoplasmic polyhedrosis virus
CatB	Cathepsin B
XFEL	X-ray free-electron laser
PhC	Polyhedra crystal
PhM	Polyhedrin monomer
NPV	Nuclear polyhedrosis virus
Bt	<i>Bacillus thuringiensis</i>
KikGR	Kikume green-red
EGFP	Enhanced green fluorescent protein
DsRed	<i>Discosoma</i> sp. red fluorescent protein
PKC	Protein kinase C
SSEA-1	Stage-specific embryonic antigen 1

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