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Innate and engineered attributes of bacterial microcompartments for applications in bio-materials science

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Bacterial microcompartments (BMCs) are sophisticated all-protein bionanoreactors widely spread in bacterial phyla. BMCs facilitate diverse metabolic reactions, which assist bacterial survivability in normal (by fixing carbon dioxide) and energy dearth conditions. The past seven decades have uncovered numerous intrinsic features of BMCs, which have attracted researchers to tailor them for customised applications, including synthetic nanoreactors, scaffold nano-materials for catalysis or electron conduction, and delivery vehicles for drug molecules or RNA/DNA. In addition, BMCs provide a competitive advantage to pathogenic bacteria and this can pave a new path for antimicrobial drug design. In this review, we discuss different structural and functional aspects of BMCs. We also highlight the potential employment of BMCs for novel applications in bio-material science.

1. Introduction

A cell, the tiniest living unit, is a compartment of selected biomolecules in a complex metabolic environment. In the course of evolution, the cell itself has gradually developed membrane-enclosed sub-compartments within it to provide spatiotemporal regulation by assorting substrates and intermediates of interconnected metabolic processes. The physical barriers of all biological compartments are composed of either lipid or protein. While lipid membrane-bound compartments are present exclusively in eukaryotes, protein-based compartments exist in both prokaryotes and eukaryotes.[1] Ferritin and its homologs, bacterial microcompartments (BMCs), lumazine synthase, vault complex, and encapsulin are some examples of lipid-free proteinaceous compartments involved in various cell functions.[2–7] Ferritin is the first discovered protein compartment and is involved in the iron storage and homeostasis of iron.[8] It is present in both eukaryotic and prokaryotic cells. On the other hand, BMCs and encapsulins are two distinct lineages of all protein compartments present only in prokaryotes. These compartments are reported to segregate and orchestrate metabolic reactions inside them.[9,10] Another class of protein-based compartments is vault complexes, which are highly conserved among eukaryotes. These ribonucleoprotein complexes take part in nuclear-cytoplasmic transport, drug resistance, and

various cell signalling processes inside cells.[11] These protein-based compartments can serve as localized nano-reactors or nano-containers *in vivo* or *in vitro* and hence have attracted the attention of several researchers in the field. In past few years, they have been utilized to fabricate novel materials with a wide range of applications.[2,12–17] Fig. 1 chronologically summarizes the discovery and the first attempt towards the application of these naturally occurring protein compartments.

Among these protein compartments, bacterial microcompartments stand unique in terms of their shell architecture and diverse functions. They evolved transiently and conditionally in several bacterial and algal species to carry out certain conditional metabolic reactions.[18] In the 1950s, BMCs were first reported in the cytoplasm of cyanobacteria *Phormidium uncinatum* through Transmission Electron Microscopy (TEM).[19–22] In the beginning, these polyhedral bodies were thought of as protein inclusions and were later considered viruses due to their morphological similarity to phages. Follow-up studies revealed that they play an important role in enhancing CO_2 fixation during photosynthesis in cyanobacteria and some chemoautotrophic bacteria and were accordingly termed carboxysomes. In 1973, carboxysomes were first purified from the bacterium *Halothiobacillis neapolitanus*.[23,24] After 40 years of carboxysome discovery, homologs of carboxysomes were found to be present in other bacteria performing special catabolic reactions and were termed catabolosomes (or metabolosomes).[25,26] To date, BMC-related genes are clustered in 7000 loci in 45 bacterial phyla that can induce 68 BMC types and subtypes.[27] Studies followed by the isolation and purification of BMCs revealed that these structures are entirely composed of protein and these protein bodies (BMCs)

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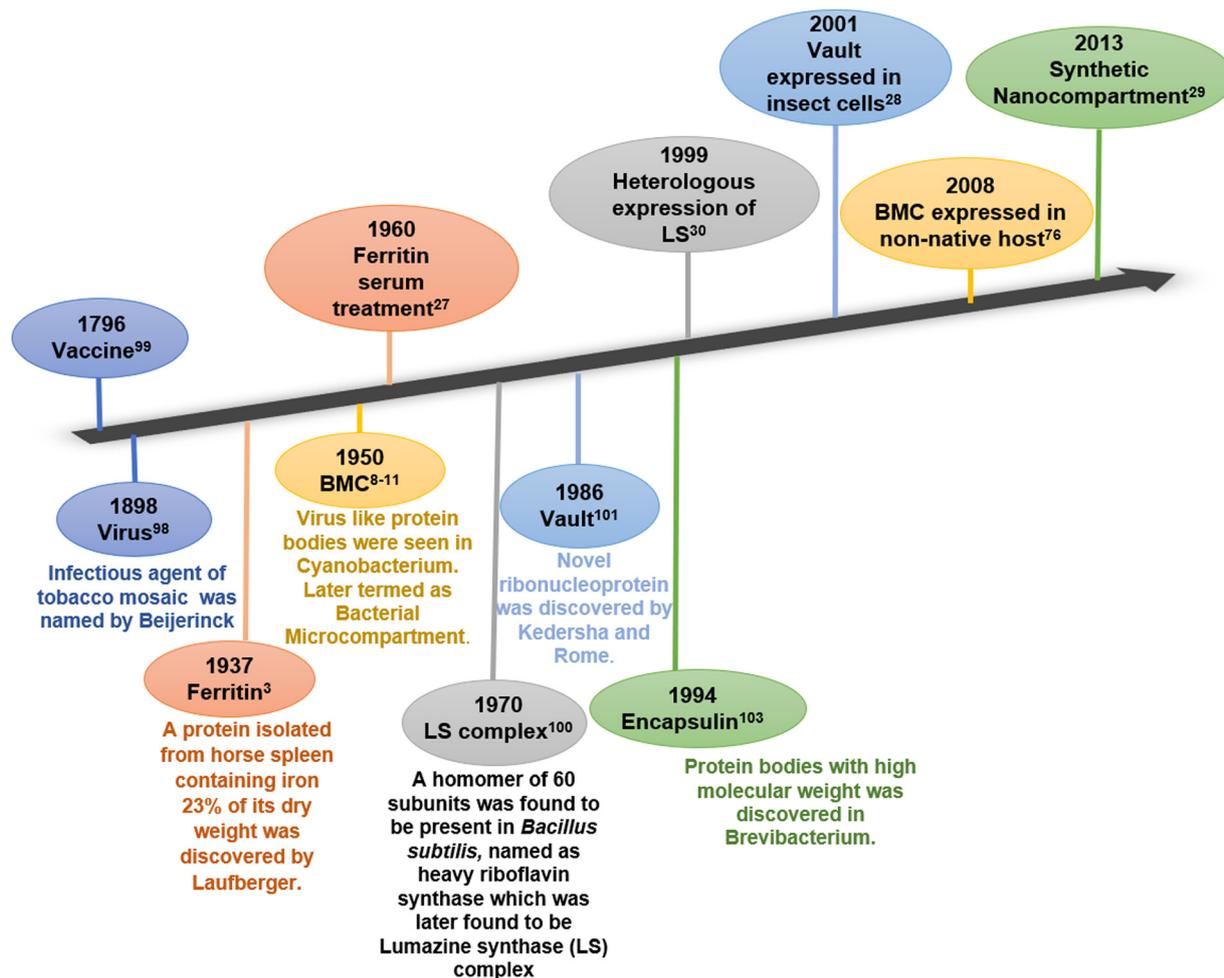*E-mail: sinhas@inst.ac.in*
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Fig. 1 Timeline of discovery (bottom) and first attempts towards applications (top) of various protein compartments.

have an enzymatic core within a protein shell.[28–30] Although the core enzyme cluster varies for different types of microcompartments, the protein subunits that make up the outer shell (shell proteins) are similar throughout several genres of BMCs.[31] The protein shell of all BMCs is made up of self-assembling protein subunits containing the two types of BMC domains.[32–35] These shell proteins have similar and superimposable structures and tile together to make up the heterogeneously self-assembled compartment envelope.[12,13] Entrapment of the metabolic machineries inside the small, confined semipermeable protein envelope of BMC increases the catalytic efficiency, orchestrates multi-enzyme pathways, and sequesters any toxic intermediates.[31] Typically, the genes that encode the BMCs are present in a single operon or clustered together.[36] These BMCs offer an open palette to design and fabricate customized biomaterials with desirable structure and function. Several efforts have already been employed to exploit the BMCs to construct synthetic bioreactors. In this review, we discuss in detail the natural and engineered properties of bacterial microcompartments towards the development of novel biomaterials for various applications. This review updates our current understanding of bacterial microcompartment applications and provides new insights into the future applications of these prokaryotic nano-organelles.

2. Innate features of BMC

2.1. Functional features of BMC

The main function of BMCs is to carry out different metabolic reactions in an efficient way inside bacterial cells, thus helping in their survival. According to these reactions, BMCs have two lineages – anabolic BMCs (carboxysomes) and catabolic BMCs (metabolosomes).

2.1.1. Fixing carbon dioxide in autotrophic bacteria – anabolic BMC. Carboxysomes are the first and only discovered anabolic BMCs to date. The core of carboxysomes consists of carbonic anhydrase and ribulose-1,5-bisphosphate carboxylase oxygenase (RuBisCO) for carbon fixation (Fig. 2). Based on the type of RuBisCO encapsulated, carboxysomes are divided into α-carboxysomes (Cso) and β-carboxysomes (Ccm). α-Carboxysomes are mainly distributed among chemoautotrophs and α-cyanobacteria and confine 1A RuBisCO, which is arranged layer-wise inside the shell with low-density lumen.[37] β-Carboxysomes present in β-cyanobacteria confine 1B RuBisCO, which is packed in a para-crystalline manner with high-density lumen.[38] Both types of carboxysomes are part of bacterial carbon dioxide concentrating mechanisms (CCM). In the CCM pathway, bicarbonate (HCO^{3-}) and inorganic CO_2 transporters in the bacterial cell

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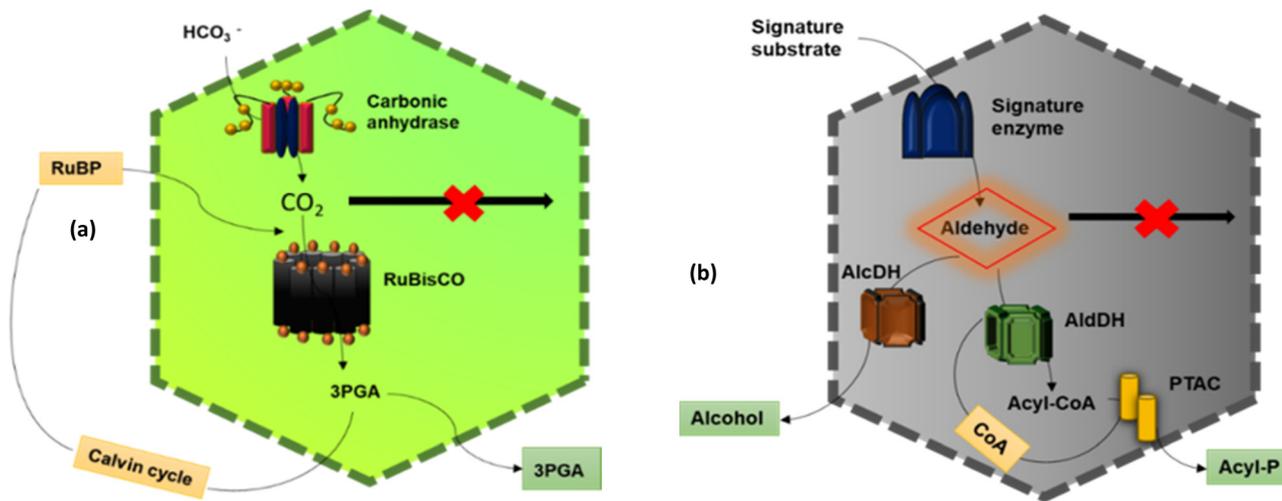*J. Mater. Chem. B*, 2023, **11**, 4842–4854 | **4843**

Fig. 2 Schematic diagram showing the action of a (a) carboxysome and (b) metabolosome.

membranes accumulate bicarbonate (HCO^{3-}) in the bacterial cytoplasm and from there, HCO^{3-} gets into the carboxysomes.[38,39] Inside the carboxysomes, carbonic anhydrase dehydrates bicarbonate to carbon dioxide. Due to the presence of the outer protein shell, carbon dioxide will be confined within the core, thus increasing the local concentration of CO_2 for RuBisCO to act on it, specifically eliminating the oxygenase activity.[40,41] Ribulose-1,5-bisphosphate (RuBP) carboxylation is catalyzed by RuBisCO forming 2 molecules of 3-phosphoglycerate (3PGA) that move out of the carboxysome to the cytoplasm from where some of the molecules are used in the Calvin Benson cycle and RuBP is regenerated.[38] Since carboxysomes are mainly distributed among oceanic photoautotrophs, they contribute 90% of the overall oxygen in Earth's atmosphere. Being a natural CO_2 concentrating reactor, it can be engineered for applications in the agricultural sector.

2.1.2. The utilization of various carbon sources – catabolic BMCs. Metabolosomes or catabolic BMCs are mainly involved in the degradation of different carbon sources by sequestration of their toxic intermediates. These microcompartments are expressed in bacteria under stress conditions, forcing the bacteria to survive on certain specific carbon sources like 1,2-propanediol, ethanolamine, ethanol, choline, fucose and rhamnose.[42–45] The metabolosomes are named according to the carbon source utilised. Pdu BMC (1,2-propanediol-utilizing BMC) and Eut BMC (ethanolamine-utilizing BMC) are some examples. The general mechanism for the utilization of these carbon sources inside these metabolosomes is shown in Fig. 3. A substrate-signature enzyme converts the substrate into a volatile and toxic aldehyde. It has been hypothesized that the BMCs have evolved to sequester this toxic volatile intermediate inside the protein cage and properly orchestrate it to downstream pathways.[46] The aldehyde product is then converted into alcohol and a cofactor derivative of carboxylic acid by alcohol dehydrogenase (AlcDH) and cofactor-dependent aldehyde dehydrogenase (AldDH) respectively. These cofactors are regenerated within the BMC by phosphotransacetylase (PTAC).[47,48] Based on the cofactor requirements of the enzymes, metabolosomes are again classified into vitamin B_{12}-dependent and vitamin
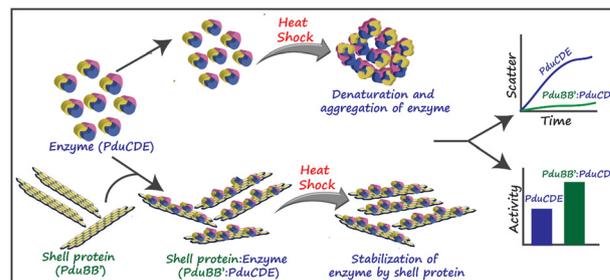

Fig. 3 The chaperone-like activity of the PduBB′ shell protein of *Salmonella enterica* on the enzyme, PduCDE, providing thermal stability to the enzyme. (Adapted from Kumar *et al.*[51]).

B_{12}-independent metabolosomes. Vitamin B_{12}-independent metabolosomes encapsulate enzymes called glycyl radical enzymes (GRE) and such microcompartments are called glycyl radical enzyme-containing microcompartments (GRMs).[49,50] One or more types of metabolosomes can be present in the same organism and the most widely studied are Pdu BMC and Eut BMC.

2.1.2.1. Propanediol Utilization Bacterial Microcompartment-Pdu BMC. PduBMC is one of the most studied metabolosomes. It is found mainly in *Salmonella* and other bacteria that contribute to the gut and soil microbiota.[52,53] The fermentation of fucose and rhamnose, common carbohydrates found in plant cell walls, bacterial capsules and glycoconjugates in eukaryotic cells, produces 1,2 propanediol. Pdu BMC catabolizes the 1,2 propanediol using vitamin B_{12}-dependent diol dehydratase. The reaction intermediate (propanaldehyde), being toxic and volatile, is prevented from moving out of the shell. Next, propanol dehydrogenase and propionaldehyde dehydrogenase convert propanaldehyde into propanol and propionyl CoA, respectively. Phosphotransacylase converts this propionyl-CoA into the propionyl-PO_4 ion; this reaction happens at the junction of the shell and cytoplasm, releasing the final product into the cytoplasm.[46,54]

2.1.2.2. Ethanolamine utilization bacterial microcompartment – Eut BMC. Eut BMC is involved in the degradation of the ethanolamine obtained from the phosphodiesterase-mediated degradation of phosphatidylethanolamine.[55] Here, the substrate-signature enzyme, ethanolamine ammonia lyase, converts ethanolamine into ammonia and acetaldehyde in the presence of vitamin B_{12} cofactor. Acetaldehyde as a volatile and lipophilic intermediate is sequestrated inside the shell and some of these molecules are converted into acetyl-CoA at first and then to acetyl phosphate, which is further converted to acetate.[56] Other molecules of acetaldehyde produce ethanol to maintain NADP/NAD$^+$ levels. Two different alcohol dehydrogenases are the key enzymes for this process and on completion of this metabolic circuit, one molecule of ATP is gained. A part of acetyl-CoA is released into the cytoplasm while another part is converted to acetal-P by phosphotransacetylase present at the border of the shell and cytoplasm.[57]

2.1.2.3. Ethanol utilization bacterial microcompartment – Etu BMC. This microcompartment was discovered in *Clostridium Kluyveri*, when it was grown on ethanol and acetate.[45] Before its discovery, researchers predicted the presence of a BMC involved in ethanol degradation from genome studies of *Clostridium Kluyveri*, in which the *etu* operon is located near the BMC loci.[45] The *etu* operon is considered to have of genes for two shell proteins, two ethanol dehydrogenases and three aldehyde dehydrogenases. These enzymes are involved in the conversion of ethanol generated from the fermentation of sugars or as a side metabolic product of Eut BMC, to acetyl-CoA.[58]

2.1.2.4. Glycyl radical enzyme containing microcompartments – Grm BMC. Grm BMCs represent a group of vitamin B_{12}-independent metabolosomes. They are involved in the degradation of propanediol, fucose and rhamnose, choline, *etc.*[49,50] The enzymes are commonly called glycyl radical enzymes. GRMs are subdivided into many groups, GRM1, GRM2, GRM3, GRM4, GRM5, GRM6, according to the substrate they metabolize. GRM1 and GRM2 are involved in choline metabolism. GRM3, 4 and 6 are similar to Pdu BMC and are involved in the degradation of 1,2-propanediol. GRM5 takes fuculose as the substrate.[27]

2.1.2.5. Other BMCs. Genome sequencing data of different bacteria done by many groups revealed the likelihood of the presence of more types of BMCs in the bacterial phyla.[59] Some of these are as follows. Sugar-phosphate utilization BMC (SPU BMC) is involved in the degradation of exogenous DNA and RNA in bacteria to produce energy. RMM BMC (*Rhodococcus* and *Mycobacterium* Microcompartemnt) is involved in the metabolism of amino acetone to propionyl-CoA. PVM BMC is involved in the metabolism of rhamnose and fucose, encapsulating a class-II aldolase enzyme.[44]

2.2. Structural features

Bacterial microcompartments can be viewed as polyhedral nano-reactors having a size range of 40–200 nm.[18] Although they are involved in diverse metabolic functions, they share some similarities in their structural features. This section presents a comparative analysis of the phenotypic and genotypic makeup of the different BMCs.

2.2.1. A family of self-assembling proteins with specific domains make up BMC–Shell/envelope proteins. The outer protein envelope, which makes up the polyhedral shell of the BMCs, is structurally composed of three types of self-assembling shell proteins, namely, hexamer (BMC-H), pseudo-hexamer/trimer (BMC-T) and pentamer (BMC-P).[32,33,60] BMC-H consists of a single domain, pfam00936 (called the BMC domain), and monomeric protein subunits, six of which assemble to form a hexamer structure. In the case of BMC-T, two pfam00936 domains form a tandem fusion and such subunits with two domains trimerize to form the overall hexagonal structure which structurally resembles BMC-H (hexamer). Hence, these trimeric proteins are also called 'pseudohexamers'.[61,62] During BMC-T assembly, they can interact side by side to form a single trimer (BMC-T^S) and/or they can interact face to face to form a double trimer (BMC-T^D). Both kinds of trimers are involved in the overall architecture of the bacterial microcompartment. Five monomers of the pfam03319 domain-containing subunits combine to form the BMC-P. While hexagonal BMC-T (*e.g.*, PduB, PduT EutL, Cso1D, CcmP) and BMC-H (*e.g.*, PduA, PduJ, PduU EutS, EutM, CcmK2, CcmO) form the facets of the polyhedra, pentagonal BMC-Ps (*e.g.*, PduN, EutN, CcmL) occupy the vertices and are twelve in number. Although pentamers are fewer in number, they are crucial for forming a closed compartment.[41,63] The deletion of a pentamer from the Pdu BMC resulted in the formation of tubular BMC, which was functional but less efficient, thus showing that the least abundant pentamer protein is important for maintaining a complete structural and functional bacterial microcompartment.[63]

The central axes of BMC-H and BMC-T form pores having average size ranges of 4 Å–7 Å and 12 Å–14 Å, respectively.[64] The size and the charge around the pore and its conformation decide the selectivity for the passage of the substrates, cofactors and products in and out of the BMCs.[64] For example, EutL (pseudohexamer) of Eut BMC has an average negative charge and is reported to open upon exposure to zinc ions.[65,66] Moreover, in the double trimer (BMC-T^D) one trimer faces the lumen and the other faces the cytoplasm. The pore of the cytoplasm-facing trimer is open and the other is obstructed by pore-surrounding amino acids, thus forming gated pores that open and close according to the substrate and amino acid interaction.[64,67] Thus, this semipermeable shell sequestrates the toxic intermediates of the reaction inside the compartment and protects the cell. Further, these pores can be engineered to tune their selectivity either by increasing or decreasing their size and altering the charge of amino acids around the pores. The permeability of different substances is gated based on the size and charges around each pore, which differs from bacteria to bacteria.

The shell proteins thus function as a semi-permeable physical barrier between the cytoplasm and the catalytic core. Some reports suggest that the shell proteins can protect the catalysts from thermal and pH stress. A recent study by Kumar *et al.* (Fig. 3) showed that the major shell protein of Pdu BMC shows

chaperone-like activity by conserving the structure and activity of its signature enzyme under thermal stress.[51]

2.2.2. Catalytic machineries inside the protein envelope – enzymatic core. The outer shell protein structures are similar in all types of BMCs but the constituents of the enzymatic core depend on their functions. Usually, enzymes that are compromised in the cytoplasmic environment and those that result in the formation of any toxic and volatile products are sequestrated within the BMC. For example, carboxysome-encompassing RuBisCO, which was evolved in the primitive environment when the amount of oxygen in the atmosphere started to increase. Oxygen, being a competitive inhibitor of CO_2 for RuBisCO, decreases its efficiency to bind to CO_2. When entrapped inside the compartment, RuBisCO is isolated from the cytoplasm environment containing oxygen, thus increasing its efficiency in binding to CO_2.[68] Metabolosomes like Pdu BMC and Eut BMC enclose aldehyde dehydrogenases, resulting in the production of toxic aldehydes, and are encapsulated along with an alcohol dehydrogenase and a phosphotransacetylase for conversion of the toxic intermediate to non-toxic products.[46] The compartment also provides a dedicated cofactor recycling system within it for the activity of encapsulated enzymes.[69]

2.2.3. Overall conserved sequence of the BMC operon – BMC loci. Since different bacterial microcompartments share similar structural attributes, the BMC operon contains some conserved regions that are homologous throughout different BMC in different bacterial phyla. All the proteins that make up a BMC are coded in a single operon called BMC loci.[36] On careful examination, it was observed that the operon had genes of shell proteins and the cargo enzymes positioned in a particular fashion. In a typical orientation, the operon starts with a promoter gene, followed by a regulator gene, which in most cases shows positive regulation of the operon. Next to the regulator, the shell/envelope protein genes are arranged so that the genes for the cargo enzyme are between two sets of shell protein genes. On induction of the respective operons with the corresponding substrates, transcription is initiated, followed by translation. The translated proteins then self-assemble to form the complete compartment. Two theories explain the self-assembly of bacterial microcompartments: the core-first assembly pathway and the concomitant assembly pathway. In the core-first assembly pathway, the enzymatic core is initially assembled and is then covered by shell proteins to form the complete BMC. This type of assembly is observed in β-carboxysomes. In β-carboxysomes, the cargo assembles first, forming a procarboxysome, onto which the shell proteins assemble and pinch off excess cargo to form mature carboxysomes. In the case of the concomitant assembly pathway, some shell proteins and some enzymes interact and form small self-assembled entities that finally self-assemble to form a complete BMC. This type of assembly is observed in α-carboxysomes and metabolosomes.[36]

2.2.4. A target sequence on cargo directs it to the BMC core – encapsulation peptide. All the enzymes that get entrapped inside microcompartments interact with the shell proteins for encapsulation and stabilization within the compartment.[70] This interaction is considered to be mediated by hydrophobic patches on the shell protein. The interaction between shell protein and core enzymes remained unknown for a very long time. Earlier comparative studies on diol dehydratase (the signature enzyme of PduBMC) and glycerol dehydratase (not associated with any BMC) revealed that one of the subunits of diol dehydratase has an N-terminal extension.[71] In 2005, it was demonstrated by Tobimartsu *et al.* that the N-terminal of the diol dehydratase is responsible for its lower solubility and the absence of the N terminal did not affect its catalytic activity. This led to further studies on the N-terminal sequence of different enzymes of Pdu BMC by different groups. In 2010, Fan *et al.* for the first time demonstrated that the N-terminal of PduP (aldehyde dehydrogenase) mediated its interaction with the PduA shell protein of Pdu BMC.[72,73] The same group demonstrated that deletion of the N-terminals of 35 amino acids of PduD reduced the encapsulation of PduCDE.[74] Further studies by Chowdhury *et al.* on Eut BMC revealed that the N-terminal sequence of EutC acts as an encapsulation peptide (EP). Later, several groups confirmed the involvement of the N-terminal or C-terminal targeting sequence in the encapsulation of enzymes inside BMCs.[75]

2.3. BMC provides competitive advantages to pathogenic bacteria

BMC enables bacteria to use alternative carbon sources other than glucose. Such alternative carbon sources like ethanolamine and 1,2-propanediol are constantly produced in the human gut through the metabolism of different food constitutents.[57,76] Enteric bacteria living in the animal gut can express different types of catabolic BMCs that utilize these substrates, thus providing a nutritional advantage over those bacteria that cannot express BMC.[77,78] Enterohaemorrhagic *E. coli*, for example, has a nutritional advantage over commensal *E. coli* because of ethanolamine metabolism when grown in bovine intestinal contents.[79] From these observations, researchers speculated that there was a connection between BMC and pathogenicity. It has been shown that the invading pathogen (*Salmonella enterica*) splits into two life cycle patterns; one directly attacks the gut epithelium causing inflammation, leading to anaerobic conditions and thus helping the other bacteria to colonize within the gut by functional BMC induction.[80] Studies on *S. typhimurium* suggest the upregulation of the *Eut* and *Pdu* operons during colonization of the bacteria in the chicken caecum.[81] Another study shows that deletion of the regulatory unit of *Pdu* and *Eut* operons reduces the proliferation of *S. typhimurium* in an animal model.[82] Although a direct connection between BMC and pathogenicity is yet to be found, it is evident that BMC provides growth advantages to bacteria irrespective of pathogenicity. Following these directions, drug molecules that can inhibit the assembly of BMC can be designed to restrict the growth of bacteria.

3. Engineering BMC for biomaterial applications

The confinement of enzymatic machinery inside the small volume of a protein shell commendably increases the catalytic

4846 | *J. Mater. Chem. B*, 2023, **11**, 4842–4854

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efficiency. Further, the protein shell also protects the encapsulated enzymatic cargo with high thermal and pH stability.[51] Another important property of bacterial microcompartments is that they can self-assemble heterogeneously, and each shell protein can self-assemble, homogenously forming different nanostructures based on the concentration in both *in vivo* and *in vitro* environments. All these features of BMCs, the high catalytic efficiency, multifunctionality, semi-permeability and self-assembling nature of the protein shell make BMCs superior to other protein compartments for exploitation as biosynthetic nano-reactors. Enticed by these special features, scientists have tailored BMCs for different bio-material applications.

The engineering of bacterial microcompartments was initially considered for use in expressing heterologous enzymes inside the shell to increase their catalytic efficiency. Later, inorganic catalysts like nanoparticles, and organic polymers like DNA were incorporated within or on the shell for industrial and biological applications. The self-assembly of the shell proteins has been used to attach desired reactive moieties nearby for electron conduction, charge transfer, *etc.* and the core with diverse functionality.

Any engineered BMCs or their components need to be supported by *in vitro* physical and biochemical characterization, following the expression and purification of these structures as a whole or as components. Further purification will be needed for the covalent attachment of any moieties to these proteins. To date, four different types of BMCs have been purified from different organisms by three main methods: cell lysis, followed by ultracentrifugation, sucrose gradient separation and affinity chromatography.[83] Purification of BMCs can be carried out from host organisms as well as heterologous expression systems.[83]

3.1. The fate of heterologous expressions of BMC and its components

An initial step in engineering bacterial microcompartments is the expression of BMC proteins in non-native hosts. This step is important as native BMC proteins will be expressed along with the engineered ones otherwise. For the first time, Parson *et al.* demonstrated the heterologous expression of Pdu BMC of *Citrobacter freundii* in *E. coli*, which was a fully functional and complete recombinant metabolosome.[84] Similarly, the α-carboxysome of *Halothiobacillus neapolitanus* was also successfully expressed in *E. coli*, which was observed to be fully intact and operative.[68] In 2018, β-carboxysomes of *Syncechococcus elongates* were expressed in *E. coli*, which was also functional.[85] These are some examples of the first few successful attempts at the heterologous expression of different bacterial microcompartments. Since the BMC operon can be expressed heterologously, engineering the same for synthetic compartments and structures was considered. Although the BMC operon codes for many proteins, all proteins are not important in forming the complete polyhedral structure of BMC. This was identified in the year 2010 when Parsons *et al.* expressed all the shell proteins[86] of the *Pdu* operon heterologously in *E. coli* and later deleted individual shell proteins to study the interaction and importance of different shell proteins for forming Pdu BMC. Following this, they were successful in expressing an empty minimal microcompartment *in vivo*. While conducting these experiments by deleting certain shell protein genes, researchers observed that with certain combinations of shell proteins, novel architectures that were different from those of the polyhedral organelles were formed. A combination of PduA and PduB formed axial filaments that showed motility when tagged with PduV (enzyme).[87] Cheng *et al.* showed that PduA, PduBB′, PduJ and PduN are only required to form a complete functional Pdu BMC. PduJ and PduN deletion resulted in the formation of an elongated BMC, while the deletion of PduA resulted in an enlarged BMC.[88] Choudhary *et al.* showed that the EutS shell protein alone can form a compartment-like structure by heterologously expressing the recombinant EutS of *Salmonella enterica* in *E. coli*.[89] In 2018, Hagen *et al.* formed an *in vitro* minimal microcompartment using a physical mixture of *Haliangium ochraceum* BMC-H, BMC-T and BMC-P in a ratio of 60 : 20 : 12, respectively. Another example of *in vivo* nanoarchitecture formation is the tubular Pdu BMC observed as a result of PduN deletion from the *Pdu* operon.[63] A recent report suggests the formation of pleiotropic structures including nanotubes and nanocones when six shell proteins of a GRM from *Rhodopseudomonas palustris* were expressed in *E. coli*.[90] The self-assembly of shell proteins into such diverse types of structures disclosed the wide plasticity of BMC in bio-material applications.

3.2. Repurposing BMC as a nanoreactor

Researchers have so far explored both anabolic and catabolic bacterial microcompartments for various biotechnology applications. BMC as a nanoreactor is prominent among all the other applications. Being such a well-orchestrated system, BMCs can carry out complex engineered pathways in the most efficient manner resulting in enhanced yield. The encapsulation of non-native cargo with encapsulation peptide (EP: described in the previous section) inside the BMC shell was the first step taken to fabricate the nanoreactor. Green Fluorescent Protein (GFP) fused with either aldehyde dehydrogenase or the first 34 amino acids from the N-terminal of the enzyme was targeted to the compartment. Although the efficiency of targeting was not high, this study demonstrated how cargo could be targeted to BMC.[83] In the same year, pyruvate decarboxylase and alcohol dehydrogenase fused with EP were targeted to Pdu BMC to produce a BMC that was efficient in producing ethanol from pyruvate[91] (Fig. 4). Wagner *et al.* showed that 3 different non-native enzymes, beta-galactosidase, esterase Est 5 and glycerol dehydrogenase, with 19 amino acids from PduP can be targeted to BMC without altering the enzyme efficiency.[92] When the BMC operon from *Citrobacter freundii* is co-expressed with polyphosphate kinase (PPK 1) fused with EP, the enzyme became encapsulated in BMC. PPK 1 loaded in BMC was able to sequester inorganic phosphate (Pi), a common water pollutant with more efficiency than free PPK 1. BMC can thus be repurposed to fabricate a reactor that can be utilized in wastewater treatment.[31] In another attempt, a hydrogen-producing nanoreactor was created using shell proteins of α-carboxysomes and directing hydrogen-producing machineries, [Fe–Fe] hydrogenase, ferredoxin (Fd) and ferredoxin:NADP$^+$ oxidoreductase

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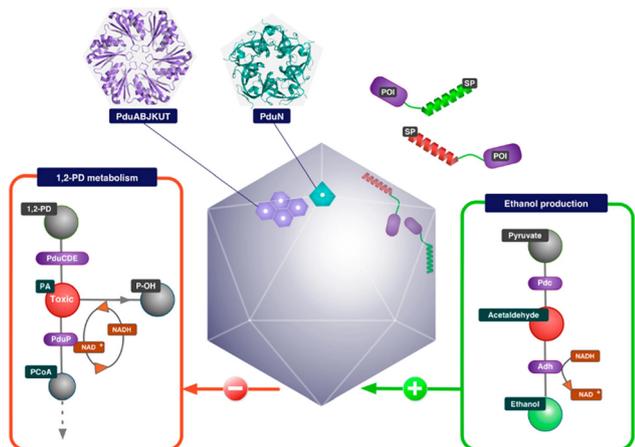*J. Mater. Chem. B*, 2023, **11**, 4842–4854 | **4847**

Fig. 4 Schematic illustration of an ethanol bioreactor, repurposing Pdu BMC. (Adapted from Wagner et al.[92]).

(FNR), into the protein shell. The developed reactor was not only producing hydrogen more efficiently than the free hydrogenase enzyme but also protecting the hydrogenase from oxygen.[93] To enhance CO_2 fixation by Rubisco in carboxysomes, the components of Rubisco activase (CbbO and CbbQ) were installed inside the carboxysome shell.[94] Prentice et al. invented recombinant bacteria expressing BMC with heterologous enzymes that were used for the accumulation of metabolites. In this case, lower molecular weight substrates were converted to polymeric or higher molecular weight products by the encapsulated enzymes. These polymeric metabolites were stored within the synthetic BMC.[95] In the context of developing nanoreactors by engineering BMC, these are some versatile paradigms for using EP to target non-native cargo into the BMC lumen. Just like EP, a hetero-dimeric coiled-coil system can also be used to target cargo in the BMC lumen, as described by Lee et al.[96] It was also reported that the SpyTag/SpyCatcher system was utilized to incorporate various fluorescent cargo inside the compartment.[97] Using this method, recent reports proclaimed the development of a synthetic formate utilizing microcompartments (sFUT).[98,99] This recombinant microcompartment was loaded with pyruvate formate lyase (PFL) and phosphate acetyl transferase and thus it was able to utilize formate and acetyl-phosphate to produce pyruvate. Zhang et al. covalently attached the SpyTag/SpyCatcher system to EutM, the hexameric sheet-forming shell protein of Eut BMC, for conducting augmented cascade reactions.[100] These studies revealed the calibre of BMC for developing a bio-reactor for various applications.

3.3. Delivery vehicle derived from BMC

One of the challenges addressed by many researchers is to develop a vehicle for delivering cargo molecules like drugs, DNA, RNA, metal nanoparticles, peptides or proteins inside cells. These cargo-loaded systems could have potential applications in cancer diagnosis and therapy.[101,102] In this context, very few attempts have been made with BMC. The possibility of using BMC as a cargo carrier was reported elsewhere.[103] The shell protein of BMC could be engineered to have cysteine at the desired position and with that, a drug molecule of specific function can be attached.[103] Recently, a protein shell (PS) was fabricated from a sheet-forming shell protein of Pdu BMC (Fig. 5). Here, anticancer drugs like doxorubicin and curcumin were loaded in the lumen of the protein shell and their release was checked.[104] Moreover, the non-native enzyme, Cyt C, was also able to be encapsulated inside the protein shell and this enzyme-loaded protein shell successfully converted pyrogallol to phloroglucinol.[104] In a recent report, DNA was targeted to the lumen of BMC for the first time. In this work, a DNA segment with the Lac repressor binding site was used and the Lac repressor was fused with an enhanced green fluorescent protein and Pdu BMC targeting peptide to target the DNA assembled inside the BMC lumen.[105] The use of BMCs has been also proposed in the field of phytonanotechnology using the nanostructures derived from the BMC shells for the controlled uptake and release of fertilizers, pesticides, or other agrochemicals that can interact directly to benefit the plant.[106] An advantage of BMCs as drug delivery vehicles is that they can shelter the cargo from thermal and pH stress.[51,107] However, the biocompatibility of BMC needs to be explored.

3.4. Developing protein-based hybrid materials

The symbiotic combination of biomolecules with inorganic nanoparticles or organic molecules produces hybrid materials, which are known to have better functionality and elaborate
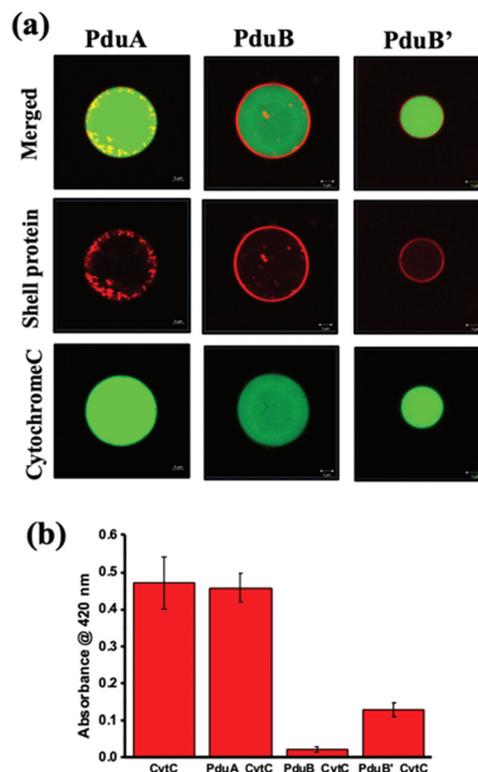

Fig. 5 (a) Confocal images of alexa 488-labelled cytochrome C loaded in a constructed protein shell. (b) Enzymatic conversion of pyrogallol to phloroglucinol by Cyt C inside the protein shell. (Adapted from Bari et al.).[104]

applications. To fabricate such materials, proteins or peptides have been used widely since their shape and size can be precisely controlled through genetic engineering. Whole Pdu BMC was used to reduce auric chloride to gold nanoparticles and after reduction, the gold nanoparticles were found to be present on the outer protein shell of BMC.[108] The developed hybrid system was able to perform standard gold nanoparticle-catalyzed reactions without losing the internal enzyme cascade activity of the BMC core. Another work from the same group demonstrated how the quaternary structure of the protein can affect the morphology and catalytic efficiency of nanocatalysts. In this work, they used a globular, non-self-assembling protein *i.e.*, BSA and a self-assembling and sheet-forming shell protein of Pdu BMC, *i.e.*, Pdu BB′, to develop a hybrid copper nano-flower (Cu NF). The Cu NF developed with BMC protein was found to have a different morphology and higher catalytic activity in comparison to BSA Cu NF.[109] In another study, Ccm (Cytochrome *c* maturation) system 1 (Ccm 1) was used to covalently attach heme with BMC-H to develop a long-distance electron transport system.[110] After the attachment of the heme moiety, shell proteins retained their property of self-assembling into hexameric structures (Fig. 6). As a proof of concept, the pore of a BMC shell protein was functionalized to bind inorganic redox counterparts such as copper or Fe–S clusters. The redox potential of these hybrid shell proteins was checked against the Standard Hydrogen Electrode (SHE). Inorganic counterparts bound to the concave side of the shell protein were found to have more positive formal potential in comparison to those variants having the same counterpart at the convex face.[110] Kaur *et al.* showed that a 2D sheet-forming shell protein, PduA (from Pdu BMC), when mutated to PduA [K26A] formed a 3D scaffold. Then, both 2D and 3D structures were used to synthesize a protein-gold nanoparticle hybrid material. In the case of a 3D scaffold, the size of the gold nanoparticle was found to be lower as this scaffold has a higher surface area that provides more nucleation sites for

nanoparticle formation. Moreover, gold nanoparticles synthesized in the presence of a 3D scaffold have greater efficiency in reducing *para*-nitrophenol.[111] These attempts at combining BMC shell proteins with organic or inorganic counterparts open many future directions to have novel hybrid material with diverse applications.

4. Conclusions and future outlook

The bacterial microcompartments are unique paradigms of biomolecular compartmentalization that can be exploited for myriads of applications in biology and biomaterial sciences (Fig. 7). To date, the literature has ample resources on the techniques that can be used for the genetic engineering of these nano-architectures leading to reactors with different shapes and properties. Further studies have also delineated the properties of the individual components. However, certain innate aspects of microcompartments still need to be explored and studied. The most relevant question is concerned with what determines the size and shape of these nano-organelles. The answer to this question will aid in the bio-synthetic production of microcompartment-like engineered nanoreactors. The next issue that needs attention is the mechanism of biogenesis of such a complex system. Although recent studies show that the biogenesis of different BMC paradigms occurs *via* liquid–liquid phase separation, more studies *in vivo* and *in vitro* need to be performed for conclusive outcomes.[112] Another aspect that remains to be deciphered is the role of BMCs in the pathogenesis of the micro-organism it is associated with. Such studies will help in the identification of alternative targets for microbial management.

Further, the components of BMCs can be exploited in permutations and combinations to form different protein architectures in 2- and 3-dimensions. These scaffolds will have high thermal and pH stability as indicated earlier and can
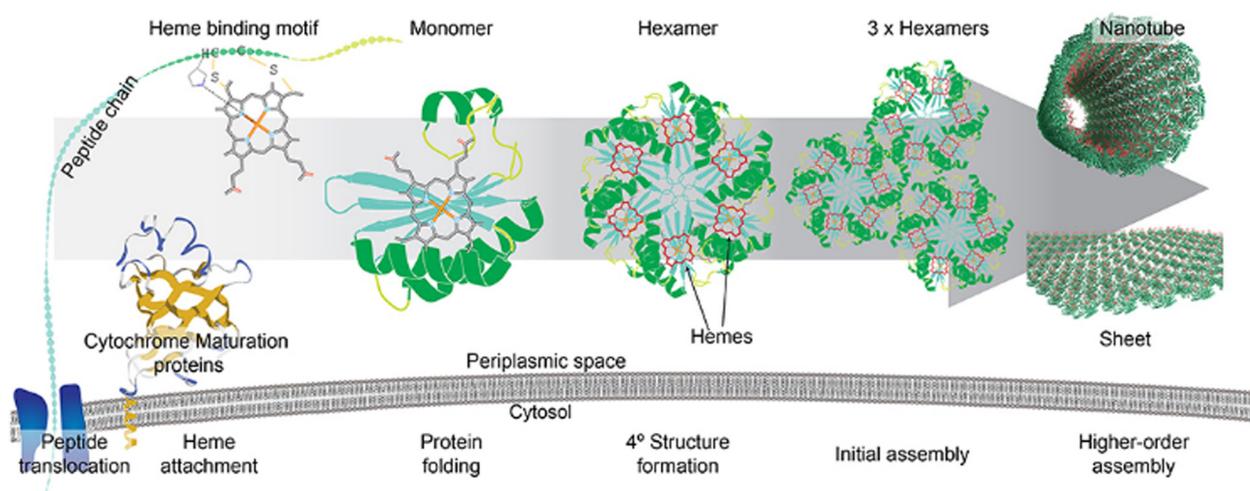

Fig. 6 The attachment of a heme moiety with a hexameric shell protein of BMC to construct a hybrid electron transporter system. (Adapted from Huang *et al.,* 2020).[110]

be used in catalysis, Raman spectroscopic probes, *etc.* The 3-dimensional structures such as protein vesicles derived from BMC components can be employed to carry bioactive molecules such as anticancer drugs, nucleic acids or peptides. This area needs the attention of researchers since self-assembling BMC shell proteins have remarkable potential for use as delivery vehicles.
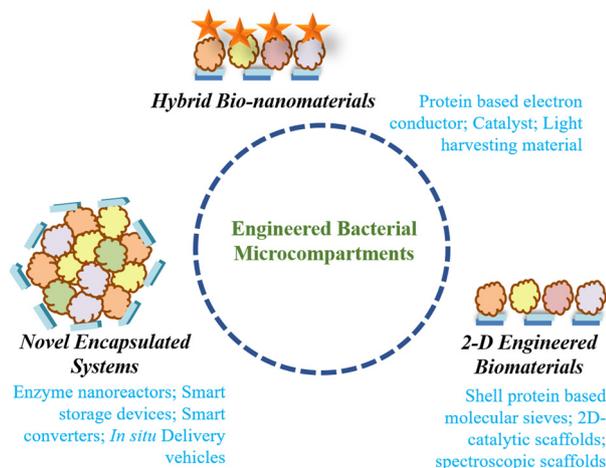

Fig. 7 Future pathways for BMC exploration.

Author contributions

SS conceived the theme of the review. SMR and AR performed the literature survey and wrote the manuscript. SS wrote and edited the manuscript.

Conflicts of interest

The authors declare no conflict of interest.

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