REVIEW

Molecular biomimetics: nanotechnology and bionanotechnology using genetically engineered peptides

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Nature provides inspiration for designing materials and systems that derive their functions from highly organized structures. Biological hard tissues are hybrid materials having inorganics within a complex organic matrix, the molecular scaffold controlling the inorganic structures. Biocomposites incorporate both biomacromolecules such as proteins, lipids and polysaccharides, and inorganic materials, such as hydroxyapatite, silica, magnetite and calcite. The ordered organization of hierarchical structures in organisms begins via the molecular recognition of inorganics by proteins that control interactions and is followed by the highly efficient self-assembly across scales. Following the molecular biological principle, proteins could also be used in controlling materials formation in practical engineering via self-assembled, hybrid, functional materials structures. In molecular biomimetics, material-specific peptides could be the key in the molecular engineering of biology-inspired materials. With the recent developments of nanoscale engineering in physical sciences and the advances in molecular biology, we now combine genetic tools with synthetic nanoscale constructs to create a novel methodology. We first genetically select and/or design peptides with specific binding to functional solids, tailor their binding and assembly characteristics, develop bifunctional peptide/protein genetic constructs with both material binding and biological activity, and use these as molecular synthesizers, erectors and assemblers. Here, we give an overview of solid-binding peptides as novel molecular agents coupling bio- and nanotechnology.

Keywords: bioinspiration; material-specific peptides; molecular recognition; biological materials evolution; binding and assembly; bionanotechnology

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1. Introduction

(a) Inspiration and lessons from biology

Nature provides inspiration for engineering structural and processing design criteria for the fabrication of practical materials to perform life’s functions (Calvert & Mann 1988; Sarikaya et al. 1990; Sarikaya 1994). During the last two decades, the realization that nanoscale inorganic materials have interesting physical characteristics based on their size (1–100 nm) has driven promises and expectations from nanotechnology with potential applications in both engineering and medical systems. Although there have been significant advances in the applications of nanotechnology, there have also been serious limitations mostly based on the problems associated with the assembly of nanoscale objects. These stem from the limitations in nanotechnological systems in controlling surface forces, inability to synthesize homologous sizes or shapes, and limitations in their higher scale, controlled organizations. In biological systems, on the other hand, inorganic materials are always in the form of nanometre-scale objects, which are self-assembled into ordered structures for full benefits of their function, that derive from their controlled size, morphology and organization into two- and three-dimensional constructions. Recently, this realization, therefore, brought biomimetics back into the forefront for renewed inspiration for solving nanotechnological problems (Sarikaya 1999; Ball 2001; Seeman & Belcher 2002). Biological materials are highly organized from the molecular to the nano-, micro- and macroscales, often in a hierarchical manner with intricate nanoarchitectures that ultimately make up a myriad of different functional elements, soft and hard tissues (Alberts et al. 2008). Hard tissues such as bones, dental tissues, spicules, shells and bacterial nanoparticles are examples that all have one or more protein-based organic components that control structural formation as well as become an integral part of the biological composites (Lowenstam & Weiner 1989; Sarikaya & Aksay 1995). These include slaffins and silicateins in silica-based structures, amelogenin in enamel and bone morphogenesis proteins or collagen in mammalian bone, calcite- or aragonite-forming proteins in mollusc shells and magnetite-forming proteins in magnetotactic bacteria (Berman et al. 1988; Cariolou & Morse 1988; Schultze et al. 1992; Paine & Snead 1996). The inorganic component could be of various types of materials (traditionally called ‘minerals’) with highly regular morphologies and three-dimensional organizations. These include piezoelectric aragonite platelets in nacre (figure 1a), precipitation-hardened single-crystal calcite with a complex architecture in sea urchin spines (figure 1b), optically transparent silica layers in sponge spicules (figure 1c) and superparamagnetic nanoparticles in magnetotactic bacteria (figure 1d).

The types of inorganics chosen by an organism have precursors or raw ingredients that are either in the soil, water or air that can relatively easily be accessed (Lowenstam & Weiner 1989; Sarikaya & Aksay 1995; Mann 1996). In addition to the intrinsic physical properties, the overall function and performance of the biological material is derived from the high degree of control that the organisms have over the formation of the structure of the material produced. The traditionally used term, ‘biomineralization’, therefore, is a misnomer, as the inorganics produced are not minerals but are materials with...
‘unique’ architectures with detailed micro- and nanostructures, including defect structures such as dislocations and mechanical or crystallographic twins, all specific to the organism that is producing them (Sarikaya 1994). For example, even in the case of mother of pearl, each of the organisms, e.g. pinctada, nautilus or abalone, producing it has different single-crystal aragonite platelets that are different from each other and each different from that of geological aragonite single crystal, in terms of the crystal itself, morphology and, more significantly, intrinsic physical properties, such as elastic modulus. From this point of view, these materials fabrication processes could be called biomaterialization to give the true meaning to the biological processes. The biological processing or fabrication (different from bioprocessing or biomimetic processing) is accomplished at ambient conditions of (near) room temperature, pH approximately 7.0 and in aqueous environments (Lowenstam & Weiner 1989; Coelenf & Antonietti 2008).

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As we see in figure 1a, nacre has a brick-and-mortar architecture consisting of layered segmented aragonitic (orthorhombic CaCO$_3$) tiles separated by an organic matrix. The organic is in the form of a 10 nm or thinner film that contains both proteins and polysaccharides, such as chitin. Either within the layer or on the surface of the organic film or within the particles themselves, the proteins possibly nucleate the inorganic, aragonite, establish its crystallography and control the growth. The resultant architecture, mother of pearl, is one the most durable hybrid composites with excellent specific toughness/strength combinations (Mayer & Sarikaya 2002). In figure 1b, sea urchin spines are single crystals of calcites (rhombohedral CaCO$_3$) with complex architectures. The spicule has high toughness and elastic modulus, unusual for a mineral calcite. Despite its single crystallinity, excellent mechanical property combinations in the spicule are probably due to the presence of nanoscale MgCO$_3$ precipitates, each associated with a strain field, toughening the otherwise brittle calcite matrix through microcrack closure (H. Fong & M. Sarikaya 2008, unpublished data). Both the formation of the complex architecture of the calcite and the presence of precipitates must, again, be due to the control that proteins have over these essential structural formations. As another example (figure 1c), the spicules of the sponge species *Rosella* are known to have excellent light collection (via the lens-shaped tip) and transmission (via the stem) properties with an interesting layered structure made up of non-crystalline silica (Sarikaya et al. 2001), all controlled by the silica-binding proteins known as silicatein (Morse 1999; Muller 2001). Finally, in magnetotactic bacteria (figure 1d), superparamagnetic single particles of magnetite (Fe$_3$O$_4$) form a string of particles aligned to sense the Earth’s magnetic field, aligning the bacteria and directing their motion via magnetotaxis (Frankel & Blakemore 1991). Each of the magnetite particles forms within a proteinaceous magnetosome membrane, a component of which directs the magnetite formation (Sakaguchi et al. 1993).

In each of the examples above, through materialization, the resultant hybrid composite structures, incorporating inorganic and proteinaceous components, are organized at the nanometre and higher dimensions, resulting in viable mechanical, magnetic and optical devices and each offers a unique design, not yet seen in man-made engineered systems. These functional biological systems are simultaneously self-organized, dynamic, complex, self-healing and multifunctional, and have characteristics difficult to achieve in purely synthetic systems even with the recently developed bottom-up processes that use molecules and nanocomponents.

Under genetic control, biological tissues are synthesized in aqueous environments in mild physiological conditions using biomacromolecules, primarily proteins but also carbohydrates and lipids. Proteins both collect and transport raw materials, and consistently and uniformly self- and co-assemble subunits into short- and long-range ordered nuclei and substrates (Sanchez et al. 2005; Tamerler & Sarikaya 2007). Whether in controlling tissue formation or being an integral part of the tissue in its biological functions and physical performance, proteins are an indispensable part of the biological structures and systems. A simple conclusion is that any future biomimetic system, whether for biotechnology or nanotechnology, should include protein(s) in its assembly and, perhaps, in its final hybrid structure (Sarikaya et al. 2003).

In traditional materials systems, the final product is a result of a balance of interactions, dictated by the kinetics and thermodynamics of the system, that are often achieved through ‘heat-and-beat’ approaches of traditional materials
science and engineering, which provide the energy for structural formations (Kingery 1976; Reed-Hill 1991). In biological systems, on the other hand, the same balance, and the energy, is achieved through evolutionary selection processes that result in the emergence of a specific molecular recognition using peptides and proteins (Pauling 1946). As we discuss below, and throughout the paper with examples, our approach is to engineer peptides with materials selectivity and use these as molecular building blocks in organizing functional materials systems in practical proof-of-principle demonstrations. Availability of new platforms will bring to the forefront new materials functionalities provided by the solid-binding peptides that will extend current technology via coupling nanoentities using the principles of biosorption beyond those provided by traditional chemisorption or physisorption.

(b) Molecular biomimetics pathways to nano- and bionanotechnology

Molecular biomimetics uses biology’s molecular ways in genetic selection or design of proteins and peptides that can control the synthesis of nanoscale objects and self-assembly of higher ordered multifunctional materials systems (Sarikaya et al. 2003). In the development of the molecular biomimetics protocols in nanotechnology, therefore, one uses solid-binding peptides and controls the formation, assembly and organization of functional nanoentities towards building useful technologies. To accomplish the overarching task, we integrate recent developments in molecular- and nanoscale engineering in physical sciences (nanoparticle formation, nano- and micropatterning such as dip-pen nanolithography and microcontact printing, and self and directed assemblies), and the advances in molecular biology, genetics and bioinformatics towards materials fabrication all at the molecular and nanometre scales (Sarikaya 1999; Sarikaya et al. 2003). Using closely controlled molecular, nano- and microstructures through molecular recognition, templating and self-assembly properties in biology, this field is evolving from the true marriage of physical and biological sciences towards providing practical application platforms (Niemeyer 2001; Sarikaya et al. 2004). The advantage of the new approach for nanotechnology is that inorganic surface-specific proteins could be used as couplers, growth initiators and modifiers, bracers and molecular erector sets, i.e. simply as building blocks for the self-assembly of materials with controlled organization and desired functions from the bottom up.

The realization of heterofunctional nanostructure materials and systems could be at three levels (Sarikaya et al. 2004), all occurring simultaneously with a closely knit feedback similar to the biological materials formation mechanisms (Albers et al. 2008). The first is that the inorganic-specific peptides are identified and peptide/protein templates are designed at the molecular level through directed evolution using the tools of molecular biology. This ensures the molecular scale and up processing for nanostructural control at the lowest practical dimensional scale possible. The second is that these peptide building blocks can be further engineered to tailor their recognition and assembly properties similar to biology’s way of successive cycles of mutation and generation can lead to progeny with improved features eventually for their usage as couplers or molecular erector sets to join synthetic entities, including nanoparticles, functional polymers or other nanoentities on to molecular
templates (molecular and nanoscale recognition). Finally, the third is that the biological molecules self- and co-assemble into ordered nanostructures. This ensures an energy-efficient robust assembly process for achieving complex nanostructures, and possibly hierarchical structures, similar to those found in biology (self-assembly; Sarikaya et al. 2004).

In this review, we provide an overview of molecular biomimetics approaches to achieve the premises of bionanotechnology with specific applications, mostly in medicine, and summarize their potentials and limitations. Here, we first summarize the protocols, adapted from molecular biology to materials science and engineering, for selecting polypeptides that recognize and bind to solids, and describe the protocols of combinatorial biology for identifying, characterizing and genetically engineering peptides for practical use. We emphasize cell surface and phage display approaches that are well adapted for the identification of solid material-specific peptides and explain ways to further tailor peptides using post-selection engineering and bioinformatics pathways. The protocols, established over years in this group, are presented in the quantitative binding characterization of the peptides using various spectroscopic techniques. We also briefly discuss possible mechanisms through which a given peptide might selectively bind to a material. Finally, we present extensive practical examples of current achievements in the usage of solid-binding polypeptides as building blocks to demonstrate their wide range of applications and, finally, discuss future prospects.

2. Genetic selection and directed evolution of solid-binding peptides

(a) Biocombinatorial selection of peptides

Genetically engineered peptides for inorganics (GEPIs) are selected through affinity-based biopanning protocol (Sarikaya et al. 2003). Biopanning steps consist of contacting the library with the material of interest, then washing out weak or non-binders and repeating the process to enrich for tight binders to select a subset of the original library exhibiting the ability to tightly interact with the desired surface. During the biopanning step, a minimum of three to five cycles of enrichment are usually performed. Generally in early rounds, low-affinity binders can be accessed if the selection is performed under mild conditions. In later rounds, as the conditions get harsher, tight binders are also recovered. Because the chimera is encoded within the phage genome or on a plasmid carried by the cell, the identity of the selected sequences (e.g. their amino acid compositions) can be deduced by DNA sequencing (figure 2).

We selected peptides for a variety of materials including noble metals (such as Au, Pt and Pd), metals (Ag and Ti), oxide and nitride semiconductors (e.g. Cu$_2$O, ITO, GaN, ZnO), minerals (such as mica, hydroxyapatite, calcite, aragonite, sapphire and graphite) or biocompatible substrates (such as silica, titania and alumina) that were selected by using either phage display (specifically filamentous phage strain M13) or cell surface display (specifically flagellar display; Sarikaya et al. 2004). There are also a number sequences selected for various materials by other groups (Gaskin et al. 2000; Feldheim & Eaton 2007; Guo et al. 2007). The ones selected via cell surface display include gold (Brown 1997) and zinc oxide (Kjærgaard et al. 2000), whereas phage display
selected ones are for their affinity towards gallium arsenide (Whaley et al. 2000), silica (Naik et al. 2002a; Eteshola et al. 2005), silver (Naik et al. 2002b), zinc sulphide (Lee et al. 2002a), calcite (Li et al. 2002), cadmium sulphide (Mao et al. 2003) and titanium oxide (Sano et al. 2005). Some biocombinatorially selected peptides have been used to assemble inorganic particles (Whaley et al. 2000; Lee et al. 2002a,b; Mao et al. 2003; Matthes et al. 2008) or to control nucleation of the compounds that they were selected for (Li et al. 2002; Naik et al. 2002a,b; Tullman et al. 2008).

When one is focusing on the material-specific peptide interactions, finding a consensus sequence might lead to a misleading result. This could be due to the high potential that a genetic bias in the selection by the organism may produce the same sequence without the diversity. As is well known, the wealth of genetic diversity leads to an assortment of sequences, which presumably reflects the
heterogeneity of the inorganic substrates at the atomic, topographic, chemical and crystallographic levels. Chemical diversity of the surfaces alone could produce a variety of sequences due to the different binding strategies that the peptide library could entail that are derived from the shape and lattice complementarities, electrostatic interactions, van der Waals interactions or various combinations of these mechanisms (Kulp et al. 2004; Evans et al. 2008; Seker et al. 2009). The ultimate robust usage of the inorganic-binding peptides for the fabrication and assembly of hybrid materials and systems requires fundamental studies towards better insights into peptide–solid molecular interactions and their incorporation into the design of desired material-specific peptides.

(b) Structural design concepts: mutation, multimerization, conformational constraints

Both the amino acid content (chemistry) as well as the sequence of the amino acids (molecular conformation) in a given selected set of peptides could affect their binding characteristics. We have recently demonstrated that the molecular constraints can be used to tune the architectural features and, consequently, the binding properties of the first generation of selected peptides. Specifically, we used a high-affinity 7-amino acid Pt-binding sequence, PTSTGQA, to build two different constructs: one is a Cys–Cys constrained ‘loop’ sequence (CPTSTGQAC) that mimics the domain used in the pIII tail sequence of the phage library construction, and the second is the linear form, a septapeptide, without the loop (Seker et al. 2007). By incorporating surface plasmon resonance (SPR, measuring binding) and circular dichroism (CD, determining molecular architecture), one is able to analyse the consequence of the loop constraint on peptide adsorption and kinetics and the conformation of peptides. These studies are related to each other with a comparative approach (as determined in figure 2).

One may also modify the binding activity of a given selected peptide by simply increasing the number of repeats of the original sequence. This multimerization could be accomplished using the simple tandem repeat, i.e. sequential attachment of the original sequence. We applied a multiple-repeat-based strategy on both phage display selected platinum and quartz binder (7 and 12 amino acid sequences each, respectively) and cell surface selected gold binders (14 amino acids each). One would expect that, as the number of repeats increased, there would be an increase in the binding activity of a given peptide. Surprisingly, however, not in all cases was the increase in the number of repeating peptides reflected in the enhancement of binding activity. In addition, material selectivity behaviour of each of the single peptides also changed when they were used in multiple-repeat forms. These results indicate that, rather than the amino acid content in a given material-binding sequence, it is the molecular conformation (secondary structure) that is more relevant, which dictates the solid-binding function. These preliminary results, therefore, show that there is a correlation between conformational instability (or adaptability) and binding ability (Seker et al. 2009). It is imperative that, in the next stage of multimerization studies, one could incorporate designed linkers between successive sequences to intentionally conform the overall multiple-repeat second-generation peptides for desired binding and other biological functions.
(c) Binding and assembly of peptides on solids

In the design and assembly of functional inorganic solids, it is essential to understand the nature of polypeptide recognition of and binding on to solid materials. Although considerable research has been directed in the literature towards understanding peptide binding to solids, it is not yet clear how proteins recognize an inorganic surface and how it could be manipulated to enhance or reduce this binding activity. This problem is similar to protein–protein recognition in biology (Pauling 1946); in the current hybrid systems, the problem reduces to one of peptide–solid interface. Here, the peptide is relatively small, perhaps approximately 10 amino acids long (1 kDa), and the inorganic solid is relatively flat but with atomic and molecular features with mostly crystallographic lattice organization. The specificity of a protein for a surface may originate from both chemical (e.g. H-bonding, polarity and charge effects) and physical (conformation, size and morphology) recognition mechanisms (Izrailev et al. 1997; Evans 2003; Dai et al. 2004; Evans et al. 2008). Recent studies have also demonstrated that the peptide overall molecular architecture (i.e. constraint versus linear) plays a key role in the solid recognition (Hnilova et al. 2008). For a given system, these mechanisms may be all significant, but with varying degrees depending on the peptide sequence, chemistry and topology of the solid surface, and the conditions of the solvent (water). Therefore, each, to a certain degree, would contribute towards a collective behaviour. Similar to the molecular recognition in biomacromolecular systems, the major contribution, however, comes from amino acid sequences that lead to a specific molecular conformation on the surface of the solid, and to a lesser extent on composition and overall amino acid content of the peptide, as demonstrated in the example below (see §2e).

(d) Peptide binding to solids and kinetics

Among the experimental approaches to rapidly monitor protein adsorption and binding on inorganics is fluorescence microscopy (FM), which has now become a routine tool as a first step in the qualitative evaluation of these sequences with respect to their affinity and selectivity (figure 3). FM imaging is an essential part of the screening protocol in our laboratory. However, this type of characterization does not provide quantitative information of polypeptide adsorption or detailed binding kinetics or mechanism(s). Another frequently used technique in molecular biology binding assays is ELISA, an immunofluorescence labelling detection using monoclonal antibody conjugated with secondary antibody fragments (Brown 1992; Whaley et al. 2000; Naik et al. 2002a; Dai et al. 2004; Sarikaya et al. 2004). Although time consuming and statistically less significant, scanning probe microscopy (SPM) protocols could also be used, which require the integration of sample preparation, self-assembly, tip design, observation conditions, data analysis and interpretations of specific polypeptides binding on to inorganic surfaces (Whitesides et al. 1991). Both atomic force microscopy (AFM) and scanning tunnelling microscopy (STM) techniques have been used to acquire static information of peptide binding to solids. The quantitative data towards determining kinetic parameters of binding could, however, be obtained using more established techniques such as quartz crystal microbalance (QCM; Murray & Deshaires 2000; Bailey et al. 2002) and SPR spectroscopy (Czenderna & Lu 1984; Homola et al. 1999).
Both QCM and SPR (figure 2) have been used to quantitatively analyse peptide adsorption kinetics under various protein concentrations, solution properties, such as pH and salinity, and solid surface conditions (Sarikaya et al. 2004; Sano et al. 2005; Tamerler et al. 2006b; Seker et al. 2007; Hnilova et al. 2008). Recently, conventional spectroscopy techniques, such as X-ray photoelectron spectroscopy and time-of-flight secondary ion mass spectroscopy techniques, have also been shown to provide the fingerprint of peptide adsorption on to surfaces (Coen et al. 2001; Suzuki et al. 2007). Although difficult to carry out, the application of solid and liquid state NMR could provide quantitative information of molecular conformations of peptides, essential information towards the understanding of the mechanism of polypeptide binding on to solids (Evans 2003). Finally, molecular modelling that studies interface interactions between a peptide and a solid will lead to rapid evaluations of various types of hybrid interfaces. These studies, e.g. molecular dynamics, that make use of computational chemistry, biology and physics, are still in their infancy, but are expected to provide protocols in the near future through the implementation of model experimental systems coupled with theoretical approaches (Evans et al. 2008).

A detailed understanding of the peptide recognition and assembly processes will inevitably lead to better insights into the design of peptides for tailored binding. A better knowledge of the mechanisms of the quantitative adsorption may become possible through high-resolution surface microscopy (e.g. AFM and STM), molecular spectroscopy and surface diffraction studies as well (such as small angle X-ray diffraction). Many of these techniques, with their advantages and pitfalls, have been discussed extensively in the literature; in this review, we will discuss one technique, SPR, which provides the most practical information.
on binding kinetics and materials selectivity of peptides for solids and, therefore, is frequently used in our research in the identification of the most promising peptides that are in frequent use today for practical implementations (§3).

(e) Peptide adsorption via molecular architectural control

Most studies on the adsorption behaviour of combinatorially selected inorganic-binding peptides on to solids have focused mainly on their amino acid compositions (Naik et al. 2002b; Mao et al. 2003). Only recently have some studies addressed the peptide structural constraints on the adsorption behaviour and affinity to solids (Tamerler et al. 2006a; Makrodimitris et al. 2007; Seker et al. 2007; Gungormus et al. 2008; M. Gungormus, D. Khatayevich, C. So, C. Tamerler & M. Sarikaya 2008, unpublished data). It is well known in protein engineering that the protein molecular architecture affects its function (Alberts et al. 2008). In this example, we hypothesized that the structure–function relationship also persists in peptide binding to inorganic materials (figure 4). To assess the hypothesis, we used two gold-binding peptides that were originally selected in a cyclic form, i.e. constraint architecture, and compared their adsorption and conformational behaviours to those of their linear, free, forms using, respectively, SPR and CD spectroscopy and computational modelling. We used two gold-binding sequences that were originally selected using the FliTrx cell surface approach (Hnilova et al. 2008). These two peptides, AuBP1 (WAGAKRLVLRRE) and AuBP2 (WALRRSIRRQSY), were synthesized...
using a solid-state technique in an open dodecapeptide version, called linear (l) as well as in constraint form, i.e. through an 18-aa Cys–Cys constrained loops, called cyclic (c), to mimic the original FliTrx displayed peptide conformations. We first carried out the CD spectroscopy to assess the molecular conformations and found that the cyclic versions of AuBPs have mainly random coil structures; however, the linear versions of AuBPs also have some degree of polyproline type II (PPII) rigid structures in addition to the random coil structures \cite{Hnilova et al. 2008}. The percentage of PPII structure in \textit{l}-AuBP2 is greater than that in \textit{l}-AuBP1, and, thus, the structural differences between the \textit{l} and \textit{c}-versions of AuBP2 are much bigger than the structural differences between the \textit{l} and \textit{c}-versions of AuBP1.

The SPR analysis showed that both the linear and cyclic forms of AuBPs have high affinities to gold (e.g. $\Delta G_{ads} = -8.7 \text{ kcal mol}^{-1}$). We also found that both the linear and cyclic forms of AuBPs have random coil and PPII structures, which cooperatively promote unfolded, conformationally labile peptides that may enhance their adaptability to interfacial features that exist on gold surfaces. One would expect differences in the binding characteristics between the cyclic and linear forms as the structure may change. In fact, we found that AuBP2 has an order of magnitude higher affinity in the cyclic version than the linear one (figure 4). This difference is consistent with the observation of significant structural change in the molecular conformations of the cyclic and linear versions of AuBP2 in solution. On the other hand, the binding affinities of AuBP1 in the cyclic and linear forms are quite similar. In this case, the molecular structures of this peptide in the two architectures are similar, as we show both experimentally (CD) and via modelling. On the basis of all the evidence, we show that the sequence of the amino acids in a given peptide and its molecular conformation may be the key determinants that facilitate peptide-selective binding on solid materials \cite{Hnilova et al. 2008}.

3. Implementations of solid-binding peptides in bionanotechnology

Once a bank of fully characterized solid-binding peptides becomes available, then it could be used as a ‘molecular toolbox’ for a wide range of applications from solid synthesis to molecular and nanoscale assemblies. Here, the peptide is not only useful in linking one nanomaterial to another, but a GEPI could also be used for genetically fusing it on to another functional protein and the system used as a bifunctional molecular construct, where the peptide would be the ligand. Alternatively, a GEPI could be fused, chemically, on to a synthetic polymer, to create multifunctional hybrid polymeric structures. Below, we will demonstrate a few uses of various GEPIs in generating new functional materials systems to understand their potential usage as molecular building blocks.

(a) GEPI-assisted synthesis of nanoinorganics

Given that these genetically engineered peptides recognize and bind to minerals, there may also be an inherent capability within the sequences to influence the morphology of these minerals as well, a prospect that has not yet been fully explored in great detail. Once this is achieved, peptide-based molecular scaffolds developed may have great potential for applications in tissue
regeneration. An example from our recent work on biomineralization using hydroxyapatite (HA)-binding peptides (Gungormus et al. 2008; M. Gungormus, D. Khatayevich, C. So, C. Tamerler & M. Sarikaya 2008, unpublished data) is shown in figure 5a,b. We demonstrated that the biocombinatorially selected HA-binding peptides could offer a route for regulating calcium phosphate-based nanocrystal formation within a biomedical context. Specifically, a successful generation of cysteine-constrained M13 bacteriophage heptapeptide library was screened against HA powder. Using the library, we selected 49 sequences and two were identified for further investigation. One of these peptides exhibited the highest binding affinity (HABP1), and the other, a much lower binding affinity (HABP2) to HA, for subsequent calcium phosphate formation and biophysical characterization studies. Here, we were interested in learning whether HA-binding polypeptide sequences could also regulate calcium phosphate formation in vitro, and likewise, determine the contributions of primary sequence and secondary structural properties that are associated with HA affinity as well as calcium phosphate formation capability. We found that both peptides affected calcium phosphate formation, with the former exhibiting a higher inhibitory activity over the latter, inducing a desired morphology on the formed calcium phosphate mineral (figure 5a). The resulting nanoparticles are plate shaped, several tens of nanometres in length and only a few nanometres in thickness. These particles resemble hydroxyapatite particles in dentine in human tooth (Fong et al. 2000). These results

Figure 5. Peptide-assisted biomaterialization using GEPIs. (a,b) Hydroxyapatite synthesis in the presence of biocombinatorially selected HABP1 compared to a control containing no peptide: (a) control; (b) with HABP1. (c,d) Au nanoparticle synthesis in the presence of AuBP1 with respect to a control prepared by a non-specific peptide: (c) control; (d) with AuBP1.
reveal a possibility of peptides in controlling particle morphology that is the major difference in differentiating dental hard tissues (dentine, cementum and enamel) as well as bone architectures. Peptide-controlled morphogenesis of Hap nanoparticles could be used in regulating materialization in hard-tissue regeneration or filler design for tissue restoration.

Another example is in the morphology control of gold particles using gold-binding peptides (figure 5c,d). Gold nanoparticles with 12 nm diameter monosize can be formed at ambient conditions using the well-known Faraday’s technique by reducing AuCl₃ with sodium citrate (or other reducing agents; Turkevich et al. 1951). In the presence of peptide, reducing the gold concentration and lowering temperature allow particle formation at a slower rate, giving the protein time to interact with surfaces during the growth and providing conditions to examine the effect of gold binding during colloidal gold formation. We conducted a search for

Figure 6. Targeted co-assembly of molecular functional entities via GEPIs, i.e. nanoparticle (QD) and fluorescein molecule. (a) Biotinylated QBP1 and (b) fluorescein-conjugated QBP1. (c) A schematic description of the assembly process. Assembly of nanoparticle (QD) functionalized with streptavidin targeting biotinylated QBP on a microcontact-printed micropatterned silicon substrate (containing native silicon oxide): (i) preparing the stamp; (ii) incubation with QBP-bio; (iii) stamping on the surface of silicon wafer; (iv) incubation with SA-QD and immobilization; (v) incubation of QBP1-fluorescein; (vi) assembly of QBP1-fluorescein on exposed silica regions; and (vii) final product incorporating targeted-assembled QD and fluorescein using QBP1 as the targeting molecule. Fluorescence microscopy images (using the appropriate filter) (d) after step (iv) and (e) after step (vi). (f) Image of the overlay of (c) and (d), corresponding to step (vii).
mutants that modulated the architecture, i.e. particle versus thin film, of gold crystallites (Hnilova et al. 2008). The selection of mutants was based on the change of colour of the gold colloid (from pale yellow to a red colloid), which was related to altered rate of crystallization. Forty gold mutants were tested this way, and the sequence analysis showed that two separate mutants that accelerated the crystal growth also changed the particle shape from cubo-octahedral (the usual shape of the gold particles under equilibrium growth conditions) to flat, thin films (figure 5c,d). This new observation is interesting from the point of enzymatic effect of protein on crystal growth rather than traditionally assumed templating effect. The polypeptides, in spite of being slightly basic, may have caused the formation of gold crystals similar to those formed in acidic conditions. This suggests that the role of the polypeptides in gold crystallization is to act as an acid, a common mechanism in enzyme function, and the protocol could be used to regulate the shape of metal nanoparticles for photonic and electronic applications.

As demonstrated with the examples above, biocombinatorially selected peptides can have enzymatic effects in the synthesis, morphogenesis and fabrication of inorganic nanomaterials. Similar to biological systems, it may be expected that the solid-binding peptides may have further potential for size, crystallography and mineral selectivity, with potential usage in a variety of practical applications, from filler material in papers to paints, as well as specialized coatings (Sarikaya et al. 2004).

Figure 7. Potential application areas of GEPIs in the molecular biomimetics field, which include molecular probing, separation, nanotechnology and nanomedicine, with potential of growing in to new areas (dotted hexagons).
Directed and mediated assembly of functional nanoentities

Protein microarray technologies, used in proteomics and clinical assays, require efficient patterning of biomolecules on selected substrates (Gristina 1987; Blawas & Reichert 1998; Chicurel & Dalma-Weiszhausz 2002; Cutler 2003; Min & Mrksich 2004; Cretich et al. 2006), which is possible provided that the proteins are spatially immobilized on solid substrates via various lithography techniques, e.g. soft lithography (Xia & Whitesides 1998), dip-pen lithography (Lee et al. 2002a, b) and photolithography (Revzin et al. 2001). Recently, protein immobilization has become a key issue in bionanotechnology since immobilization provides physical support to the molecule, resulting in improved stability and activity and, furthermore, helps to separate proteins from solution, rendering them reusable (Castner & Ratner 2002; Bornscheuer 2003). The approaches for biomolecule immobilization on glass or metal (e.g. gold) substrates generally require surface functionalization by self-assembled monolayers (SAMs) of bifunctional molecules, such as amino-terminated aminoalkyl-alkoxysilanes for silica and carboxyl-terminated alkanethiols for gold substrates (Mrksich & Whitesides 1996; Ostuni et al. 1999). Despite their widespread usage, these traditionally available linkers have certain limitations, such as causing random orientation of the protein on solid surfaces and requiring multistep chemical reactions and, furthermore, the assembled monolayers can be unstable during immobilization (Fujiwara et al. 2006; Park et al. 2006). To overcome these limitations, it is preferable to have molecules as direct linkers to the solid substrate of interest, which not only have all the desired features of the conventional chemically prepared SAMs but also have specificity to a given solid substrate and assemble on to it efficiently. In addition, the molecule used as the linker could be amenable to genetic manipulation for selecting the best linker site to the displayed protein or nanoentity without causing any effect in reducing the binding activity. Solid-binding peptides can provide the multifunctionality as a preferred linker with high structural stability incorporating a target molecule aligned consistently to carry out a desired function (Sarikaya et al. 2003).

Here we demonstrate the solid-binding peptide as a molecular assembler for two different nanoentities, quantum dots (QDs) and fluorescent molecules, and sequentially assemble them on a micropatterned surface using the material specificity of the GEPI (Kacar et al. 2009). In this case, directed immobilization of the QDs is followed by the GEPI-mediated assembly of the fluorescent molecule using the microcontact printing and self-assembly procedures schematically illustrated in figure 6c. The directed immobilization of SA-QD on a QBP1-biopatterned surface is shown in figure 6d as red stripes, imaged with a fluorescent microscope using a QD605 filter, revealing red fluorescent contrast. Here, the dark stripes represent the regions originally unoccupied, exposing the bare quartz surface. Next, following the procedure in figure 6c, the assembly of the fluorescent molecule, i.e. fluorescein, is mediated using the QBP1-F molecular conjugate. The assembled conjugate molecules are imaged in green, as shown in figure 6e, using a FITC filter. At this step, the QBP1-F molecular conjugate diffuses towards the regions of the substrate previously unoccupied, after the initial directed immobilization of QDs (figure 6e). Both images in figure 6d, e were recorded from the same area of the sample, showing regular

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alternating lines of red and green stripes, corresponding to the directed-assembled QDs and mediated-assembled fluorescein molecules, respectively. This result demonstrates that the QBPI is active as an efficient molecular linker as well as a versatile PDMS ink. Furthermore, we demonstrate here the co-assembly of two diverse nanoentities without the involvement of complex surface modification, often involved in the traditional silane-based procedures (Fujiiwara et al. 2006). The patterning protocols developed here would be useful as microscale platforms for a wide range of applications from generating photonic lattices to co-assembling multi-enzyme or multi-protein assays.

4. Future prospects of solid-binding peptides as molecular building blocks in bionanotechnology

The joining of biology with materials requires an ability to design, engineer and control interfaces at the materials/bio intersections as these sites are significant in the implementation of nanotechnology, developments of new materials and protocols in molecular engineering, and realization of bionanotechnology (figure 7). Biology controls all interfaces between molecular materials, tissues and organs using peptides and proteins which are also the agents of molecular communication. In a sense, proteins are the workhorses in biology carrying out the chemical, physical and biological functions of organisms. Similar to biology, in engineering and technological systems, we can genetically select peptides with an ability to bind to inorganic materials to create a new fundamental building block to couple bio and synthetic entities. As we describe here, GEPIs have short amino acid sequences with material-selective binding and self-assembling properties. Once selected using combinatorial mutagenesis, GEPIs can be further tailored to enhance/modify their binding ability and multifunctionality. The multifunctionality could be introduced either using two or more material-binding peptides to create novel ways of making dissimilar materials thermodynamically compatible, or by genetically fusing a functional protein, e.g. enzyme or antibody, to develop heterofunctional molecular constructs.

Solid-binding peptides coupled with solid substrates form a new generation of novel hybrid materials systems (Sarikaya et al. 2003). Genetic control of the coupling and the resulting function of the hybrid material are new approaches with potential to overcome limitations encountered in the progress of a wide range of applications in which traditionally synthetic linkers, such as either thiol or silane, have been used. The attachment of biomolecules, in particular proteins, on to solid supports is fundamental in the development of advanced biosensors, bioreactors, affinity chromatographic separation materials and many diagnostics such as those used in cancer therapeutics (Blawas & Reichert 1998; He et al. 2006; Behrens & Behrens 2008). Protein adsorption and macromolecular interactions at solid surfaces play key roles in the performance of implants and hard-tissue regeneration (Gottlieb et al. 2008; Ma 2008). Proteins adsorbed specifically on to probe substrates are used to build protein microarrays suitable for modern proteomics (Cutler 2003; Cretich et al. 2006). Enzyme immobilization on substrates (e.g. nanoparticles in a colloid) will greatly enhance the usage of industrial enzymes (Kasemo 2002). Designing bifunctional peptides (e.g. attached to a probe) coupled to nanoparticles, e.g. QDs or fluorescent...
molecules, will provide new avenues for multicomponent biosensor design (Li et al. 2004; Rusmini et al. 2007). The same (nanoparticle/GEPI-probe) platform, where the probe is an antibody and the nanoparticle is a therapeutic or imaging entity, will provide a new molecular platform for cancer probing (Weissleder 2006; Tamerler & Sarikaya 2007). The examples given above illustrate only some of the achievable goals using these new classes of functional molecular linkers. All these and a wide variety of other applications form the core of biological materials science and engineering (Sarikaya et al. 2003) which can be designed and genetically engineered (figure 7). Based on its recognition and self-assembly characteristics, the role of a GEPI in these hybrid structures would be to provide the essential molecular linkage between the inorganic components, and, at the same time, be an integral component of the overall structure providing to it functional (e.g. mechanical) durability. Owing to the intrinsic properties mimicked after natural proteins, in the coming years and decades, we are likely to see engineered inorganic-binding polypeptides used more and in a wide range of applications from particle synthesis and assembly with genetically controlled physical and chemical characteristics in materials science to probing for biological targets in biology and medicine (Weissleder 2006; Sengupta & Sasisekharan 2007; Tamerler & Sarikaya 2008).

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