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# Ultrafast, cytocompatible mineralization of calcium phosphate in the formation of stratified nanoshells of artificial spores†

 Duc Tai Nguyen,<sup>a</sup> Sang Yeong Han,<sup>a</sup> Hyunwoo Choi,<sup>a</sup> Nayoung Kim,<sup>a</sup>  
Gulaim A. Seisenbaeva,<sup>id</sup> Vadim G. Kessler<sup>id</sup> and Insung S. Choi<sup>id</sup>\*<sup>a</sup>

**Spatially controlled confinement of catalytic enzymes within nanoshells holds substantial potential for applications in bioreactors, synthetic cells, and artificial spores. The utilization of amorphous calcium phosphate (CaP) precursors enables the extremely rapid (<5 seconds) construction of thick (~400 nm) CaP nanoshells, stratified with distinct enzymes, on various tannic acid-primed substrates. *Saccharomyces cerevisiae* cells are nanoencapsulated with enzyme-embedded, multilayered CaP nanoshells in a cytocompatible manner, providing an advanced chemical tool for interfacing living cells with functional entities in a spatially controlled configuration.**

Sporulation is an adaptive-survival tactic of many bacteria and fungi characterized by the formation of a robust, protective physicochemical barrier around the cell in response to nutrient deprivation or other adverse environmental conditions.<sup>1,2</sup> Inspired by biological sporulation, durable cell-in-nanoshell structures, referred to as artificial spores, have been developed by chemically interfacing individual living cells with external materials in a cytocompatible fashion.<sup>3–6</sup> Artificial spores not only provide cytoprotection against harsh environmental stresses but also allow for the chemical manipulation of cellular metabolic activities, empowering the nanoencapsulated cells with exogenous functions.<sup>7,8</sup>

Artificial spores can be constructed with various cytocompatible materials, encompassing both organic and inorganic species, to ensure cell viability during the process of single-cell nanoencapsulation (SCNE).<sup>9–11</sup> Inspired by the silica (SiO<sub>2</sub>) and calcium carbonate (CaCO<sub>3</sub>) exoskeletons found in certain unicellular organisms, such as diatoms and coccolithophores, bioinspired mineralization has been employed to fabricate artificial nanoshells composed of SiO<sub>2</sub>,<sup>12–15</sup> CaCO<sub>3</sub>,<sup>16,17</sup> titanium dioxide (TiO<sub>2</sub>),<sup>18,19</sup> or calcium phosphate (CaP).<sup>20</sup> The mild reaction

conditions for and the robustness of mineral nanoshells make bioinspired mineralization a powerful approach for the protection and functionalization of living cells at the single-cell level.

Although artificial-spore approaches with Ca minerals have been demonstrated, their applications and methodological advancements remain limited, largely due to the complexity, as well as the heterogeneity and low efficiency, of thin-film formation from Ca minerals. In nature, cell-mediated mineralization actively controls the nucleation sites, crystal growth, crystallinity, and morphology of biominerals.<sup>21</sup> However, in bioinspired mineralization, it is extremely challenging to replicate the controlled, closed, and isolated microsystems of living cells due to the stochastic nature of nucleation, particularly in the case of polymorphic CaP. Therefore, the open systems for chemical biomineralization have thus far strictly required the optimization of various factors, including solution pH, temperature, ionic strength, influx rate of metal ions, nucleation sites, and crystal growth duration, to achieve successful Ca-mineral SCNE.

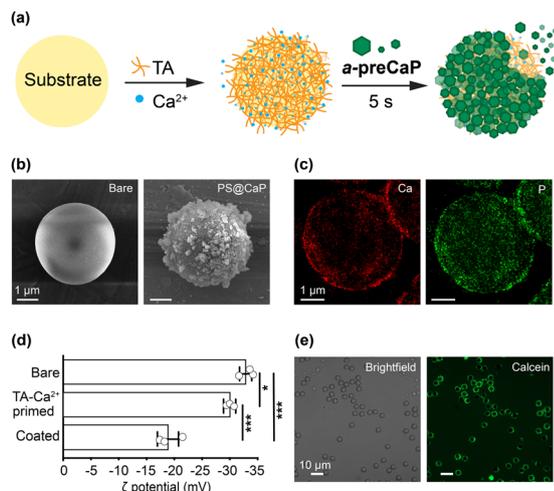
Typical Ca-mineral processes proceed through the mixing of a Ca<sup>2+</sup> solution with a HPO<sub>4</sub><sup>2–</sup> or PO<sub>4</sub><sup>3–</sup> solution at controlled pH and reaction times, which can extend to tens of minutes or hours for the formation and growth of amorphous Ca minerals on the cell surface, always carrying a high risk of unwanted nucleation in solution. For localized heterogeneous nucleation, the substrates have been pretreated with Ca<sup>2+</sup>-adsorbing polymers, such as poly(acrylic acid)<sup>22</sup> and polyvinylpyrrolidone,<sup>17</sup> to induce surface seeding. In addition, capping agents (*e.g.*, triethylamine<sup>23</sup> and polyphenols<sup>24</sup>) or zwitterion polymers<sup>25</sup> are often required to prevent the overgrowth of nanoparticles and their transformation to hydroxyapatite. Recent advances include the use of zeolitic imidazolate framework-8, which promotes exclusive surface growth of CaP through ion exchange between the surface Zn<sup>2+</sup> and Ca<sup>2+</sup> in solution.<sup>24</sup> However, the stochasticity, randomness, and inflexibility of current biomineralization protocols still limit their applications in cell-surface engineering, particularly when the incorporation of drugs, genetic materials, and enzymes is beneficial in advanced cell therapies. Hence, it is of the utmost importance to develop a rapid,

<sup>a</sup> Department of Chemistry, KAIST, Daejeon 34141, Republic of Korea.  
E-mail: ischoi@kaist.ac.kr

<sup>b</sup> Department of Molecular Sciences, SLU, Uppsala 75007, Sweden

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**Fig. 1** (a) Schematic illustration of the **a-preCaP** protocol. (b) FE-SEM images of bare PS and PS@CaP. (c) TEM-EDX Ca and P mapping of PS@CaP. (d) Zeta ( $\zeta$ ) potentials of bare PS, TA- $\text{Ca}^{2+}$ -primed PS, and PS@CaP. Data are represented as mean  $\pm$  SD (unpaired, two-sided Student's *t*-test;  $n = 3$ ; \*\*\* $p < 0.001$ , \* $p < 0.05$ ). (e) CLSM images of PS@CaP stained with calcein.

highly efficient, reproducible, and versatile methodology for bio-inspired CaP or other mineralization. In this work, we remarkably shorten the CaP mineralization process by simply interfacing an amorphous CaP precursor (**a-preCaP**) with tannic acid (TA)-primed surfaces (Fig. 1a). The formation of CaP-mineral shells around individual living cells is completed within 5 s through rapid heterogeneous nucleation of **a-preCaP**. In addition, the large adsorption surface area of the **a-preCaP** clusters enables the hierarchical incorporation of external functional biomolecules, such as enzymes, into the multilayered CaP shells, providing the catalytic empowerment of living cells in SCNE.<sup>8,26</sup>

The mechanisms on CaP formation in organisms remain uncertain. Biological CaP is often amorphous or low crystalline, and its deposition is suggested to proceed through an amorphous precursor phase, consisting of CaP clusters, rather than the ion association pathway in classical nucleation theory.<sup>27,28</sup> CaP minerals have been shown to evolve from pre-nucleation clusters of multiple  $\text{Ca}^{2+}$  and  $\text{PO}_4^{3-}$  ions into larger post-nucleation clusters and loosely packed polymeric assemblies before crystallization.<sup>29,30</sup> These nanometric structures are metastable and are often regarded as the building blocks of CaP precipitation. We utilized the concept of the amorphous precursor phase to initiate and accelerate the localized heterogeneous deposition of CaP minerals in SCNE. Specifically, we produced **a-preCaP** by mixing 4.8 mM of  $\text{CaCl}_2$  and 3.2 mM of  $\text{K}_3\text{PO}_4$  in water. Transmission electron microscopy (TEM) images showed an interconnected network of CaP assemblies, each approximately 10–20 nm in size (Fig. S1a, ESI<sup>†</sup>). Dynamic light scattering (DLS) analysis further indicated the existence of small CaP clusters (average diameter: 4.4 nm) and a large network of CaP assemblies (average diameter: 1000 nm) (Fig. S1b, ESI<sup>†</sup>). **a-preCaP** appeared amorphous or ill-defined, as evidenced in the selected area electron diffraction (SAED) pattern (Fig. S1c, ESI<sup>†</sup>). We observed that CaP minerals were produced in the solution immediately after the addition of a small amount of

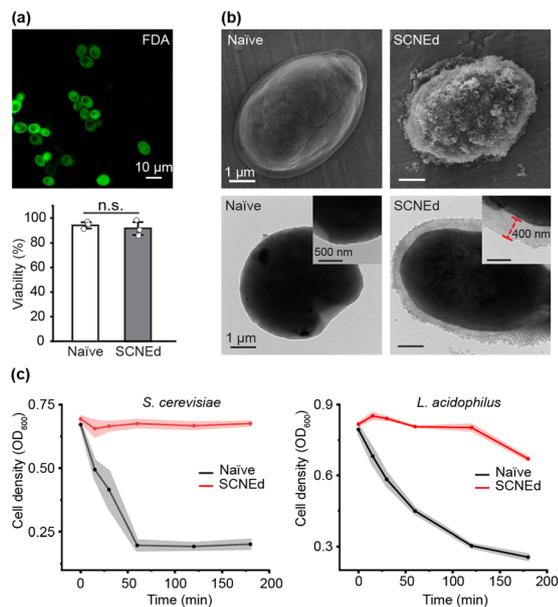
TA to the **a-preCaP** solution (Fig. S2a, ESI<sup>†</sup>). The **a-preCaP** solution (2 mL) became turbid upon the addition of 100  $\mu\text{L}$  of TA (6 mM). We thought that TA effectively reduced the nucleation energy barrier, thereby facilitating CaP nucleation.<sup>31</sup>

Following the initial assessment, we investigated the CaP shell formation with abiotic polystyrene (PS) microparticles (sulfonated, diameter: 3.97  $\mu\text{m}$ ) as a model. In brief, the PS particles were primed with a mixture of TA (4 mM) and  $\text{Ca}^{2+}$  (8 mM) and then subjected to gentle shaking with the **a-preCaP** solution for 5 s, resulting in the formation of PS@CaP. Field-emission scanning electron microscopy (FE-SEM) images showed a dramatic change in surface morphology, from smooth to particulate (Fig. 1b). Elemental mapping with energy-dispersive X-ray (EDX) analysis of PS@CaP revealed the Ca and P peaks, confirming the formation of the CaP shells (Fig. 1c). In addition, zeta ( $\zeta$ ) potential changes from  $-32.8$  to  $-30.7$  mV after TA- $\text{Ca}^{2+}$  priming, and to  $-18.9$  mV, after CaP deposition, further supported the shell formation (Fig. 1d). The X-ray diffraction (XRD) analysis indicated that the CaP shells were amorphous (Fig. S2b, ESI<sup>†</sup>). The CaP shells were visualized in the confocal laser-scanning microscope (CLSM) images, after staining the shells with calcein, a green-fluorescent  $\text{Ca}^{2+}$  indicator (Fig. 1e). The analyses underscored the rapid and intimate interactions between **a-preCaP** in solution and TA on the surface, leading to the ultrafast formation of CaP shells.

**a-preCaP** was also found to strongly interact with proteins. For example, preincubation of the **a-preCaP** solution with bovine serum albumin-Alexa Fluor 647 (BSA-Alexa 647, final concentration: 10  $\mu\text{g mL}^{-1}$ ) inhibited the shell formation on the TA- $\text{Ca}^{2+}$ -primed PS surface, with predominant homogeneous nucleation (Fig. S3a, ESI<sup>†</sup>). However, lowering the BSA-Alexa 647 concentration to 5  $\mu\text{g mL}^{-1}$  led to heterogeneous nucleation within a short time, suggesting the potential for incorporating proteins into the CaP shell. UV-vis analysis indicated that about 60% of BSA-Alexa 647 was incorporated into the CaP shell within 5 s (Fig. S3b, ESI<sup>†</sup>). The immobilization efficiency nearly doubled compared with the post-adsorption of BSA-Alexa 647 onto PS@CaP (efficiency: 30%), presumably due to the high surface area of **a-preCaP**.

Our ultrafast protocol for CaP shell formation was successfully applied to living *Saccharomyces cerevisiae*, creating the SCNEd cell, *S.cerevisiae*@CaP. The excellent cytocompatibility of our protocol was confirmed by the fluorescein diacetate (FDA) assay, which showed no noticeable decrease in viability after SCNE (viability, relative to naïve cells: 97.5%) (Fig. 2a). The CaP shell formation was supported by the changes from  $-24.6$  mV in  $\zeta$  potential to  $-20.1$  mV after TA- $\text{Ca}^{2+}$  priming and to  $-7.4$  mV after CaP shell formation (Fig. S4, ESI<sup>†</sup>), as well as by the CLSM analysis, alongside FE-SEM and TEM analyses. TEM images showed the complete CaP shells, about 400 nm thick, after just 5 s of gentle shaking, highlighting the ultrafast CaP-shell formation through the **a-preCaP** approach (Fig. 2b). As a comparison, we also formed *S.cerevisiae*@CaP by the conventional method,<sup>20</sup> where  $\text{Ca}^{2+}$ -TA-primed *S. cerevisiae* was incubated in 800  $\mu\text{L}$  of a  $\text{Ca}^{2+}$  solution (6 mM), and 200  $\mu\text{L}$  of a  $\text{PO}_4^{3-}$  solution (16 mM) was added dropwise to the cell suspension at 20  $\mu\text{L min}^{-1}$  (final concentrations: 4.8 mM of  $\text{CaCl}_2$  and 3.2 mM of  $\text{K}_3\text{PO}_4$ ), resulting in much thinner ( $\sim 100$  nm), uneven, and incomplete shells (Fig. S5, ESI<sup>†</sup>).

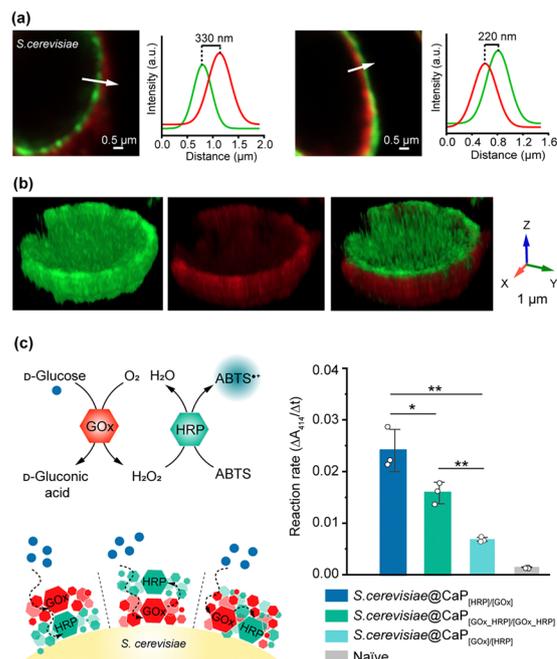




**Fig. 2** (a) Live/dead assay: CLSM image and viability of *S. cerevisiae* before and after SCNE. FDA (green): live and PI (red): dead. Data are represented as mean  $\pm$  SD (unpaired, two-sided Student's *t*-test; *n* = 3; n.s.: not significant). (b) (top) FE-SEM and (bottom) TEM images of naive and SCNEd *S. cerevisiae*@CaP. (c) Time-dependent OD<sub>600</sub> changes of naive (black) and SCNEd cells (red) under lytic enzyme attack. (left) *S. cerevisiae* and (right) *L. acidophilus*.

Furthermore, we observed no noticeable heterogeneity in shell formation, whether between batches or within a batch (Fig. S6, ESI<sup>†</sup>). Our ultrafast, cyto-compatible approach can be simply expanded to the SCNE of other cell types, such as probiotic *Lactobacillus acidophilus* (Fig. S7, ESI<sup>†</sup>). The thick and durable CaP mineral shells noticeably protected the SCNEd cells from attacks by lytic enzymes (Fig. 2c). The cell density (OD<sub>600</sub>) of SCNEd *S. cerevisiae* or *L. acidophilus* was monitored alongside their naive counterparts in solutions containing lyticase or lysozyme, respectively. SCNEd cells maintained high cell densities (*S. cerevisiae*: 97.5%, *L. acidophilus*: 82%), while only 30% of naive cells remained after 3 h of reaction, indicating the enhanced resistance of SCNEd cells conferred by the thick CaP shells.

As a proof-of-concept for hierarchical protein embedment in CaP shells, two different BSA proteins, BSA-Alexa 488 (green) and BSA-Alexa 647 (red), were separately compartmentalized within the double-layered CaP shells on *S. cerevisiae*. The shell-forming process was repeated with BSA-adsorbed  $\alpha$ -preCaP, producing *S. cerevisiae*@CaP<sub>[layer 1]/[layer 2]</sub>. CLSM images showed distinct green and red layers surrounding *S. cerevisiae* in *S. cerevisiae*@CaP<sub>[BSA-Alexa 488]/[BSA-Alexa 647]</sub> and *S. cerevisiae*@CaP<sub>[BSA-Alexa 647]/[BSA-Alexa 488]</sub> (Fig. 3a). Z-stack CLSM images of *S. cerevisiae*@CaP<sub>[BSA-Alexa 488]/[BSA-Alexa 647]</sub> captured the spatial arrangement of red and green layers, confirming the orderly compartmentalization of BSA-Alexa 488 and BSA-Alexa 647 within the double-layered CaP shells (Fig. 3b). Line profiles of the shells revealed a 200–300 nm separation between the green and red fluorescence peaks, further validating the compartmentalization of the proteins. UV-vis analysis determined incorporation efficiencies of 91.2% and 81.5% for the first



**Fig. 3** (a) CLSM images and line profiles of (left) *S. cerevisiae*@CaP<sub>[BSA-Alexa 488]/[BSA-Alexa 647]</sub> and (right) *S. cerevisiae*@CaP<sub>[BSA-Alexa 647]/[BSA-Alexa 488]</sub>. (b) Reconstructed 3D z-stacked CLSM images of *S. cerevisiae*@CaP<sub>[BSA-Alexa 488]/[BSA-Alexa 647]</sub>. Green: BSA-Alexa 488; red: BSA-Alexa 647. (c) (left) Schematic for cascade reactions of GOx and HRP. (right) Graph of reaction rates for different combination and order of GOx and HRP in *S. cerevisiae*@CaP shells. Data are represented as mean  $\pm$  SD (unpaired, two-sided Student's *t*-test; *n* = 3; \*\**p* < 0.01, \**p* < 0.05).

and second layers, respectively. It is to note that the hierarchical compartmentalization of proteins within mineral-based nanoshells remains elusive. In addition to proteins, DNA was immobilized within the CaP shells, as evidenced by the red fluorescence of propidium iodine in CLSM images (Fig. S8, ESI<sup>†</sup>). Moreover, our protocol enabled the formation of multi-layered shells, as demonstrated by the triple-layered CaP shell in *S. cerevisiae*@CaP<sub>[BSA-Alexa 647]/[BSA-Alexa 488]/[DNA-Hoechst]</sub> (Fig. S9, ESI<sup>†</sup>). These results underscored the potential of our system for embedding various functional entities within multilayered CaP shells in a hierarchical manner, benefitting the development of cascade reaction systems.<sup>8,32–34</sup>

Two enzymes—glucose oxidase (GOx) and horseradish peroxidase (HRP)—were compartmentalized within double-layered CaP shells on *S. cerevisiae* cells, with the order and combination of enzymes varied in each layer. In this system, GOx catalyzes the glucose oxidation, generating hydrogen peroxide (H<sub>2</sub>O<sub>2</sub>) as a byproduct. HRP subsequently utilizes H<sub>2</sub>O<sub>2</sub> to oxidize 2,2'-azino-bis(3-ethylbenzothiazoline-6-sulfonic acid) (ABTS) as a probe into ABTS<sup>•+</sup>, producing a characteristic dark green radical ( $\lambda_{\max}$  = 414 nm). As a negative control, naive cells showed no significant change in ABTS<sup>•+</sup> absorbance over 4 h, confirming the lack of enzymatic activity (Fig. S10a, ESI<sup>†</sup>). In contrast, SCNEd cells functionalized with GOx and HRP demonstrated robust cascade reactions, evidenced by a marked increase in ABTS<sup>•+</sup> absorbance (Fig. S10b, ESI<sup>†</sup>). Notably, *S. cerevisiae*@CaP<sub>[HRP]/[GOx]</sub> achieved a 3.53-fold higher reaction rate compared with *S. cerevisiae*@CaP<sub>[GOx]/[HRP]</sub>,



highlighting the critical role of spatial enzyme arrangement within CaP multilayers for efficient substrate channeling (Fig. 3c).<sup>8,26</sup> Furthermore, *S.cerevisiae*@CaP<sub>[HRP]/[GOx]</sub> showed a 1.52-fold rate increase compared with *S.cerevisiae*@CaP<sub>[GOx\_HRP]/[GOx\_HRP]</sub>, where both enzymes were co-mixed in each layer. These results showed the potential of hierarchical enzyme compartmentalization within CaP shells to enhance the catalytic efficiency of multicompartmental systems by regulating the directional trafficking of substances and substrates. We also stored naïve and *S.cerevisiae*@CaP<sub>[GOx\_HRP]</sub> cells at 4 °C for 7 days without any added nutrients, showing no noticeable decrease in cell viability compared with naïve cells, although the reaction rates decreased linearly over time, presumably due to leaching (Fig. S11, ESI†).

In summary, the **a-preCaP** strategy enables facile and ultra-fast CaP-based nanoshell formation on various substrates, generating thick (~400 nm) CaP nanoshells around tannic acid-primed substrates in under 5 s. Beyond cytocompatibility, **a-preCaP** could adsorb biomolecules, enabling spatially controlled compartmentalization of target molecules within the CaP nanoshells. We have demonstrated that the combination and arrangement of GOx and HRP within the double-layered shells significantly affect the enzymatic cascade performance, emphasizing the critical role of spatial organization in multi-enzyme systems. The multi-compartment, multifunctional artificial spores hold promise for advanced cell therapies, cell factories, cell microrobots, and targeted delivery systems.

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## Data availability

The data supporting this article have been included as part of the ESI.†

## Conflicts of interest

There are no conflicts to declare.

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