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High-temperature ternary Cu–Si–Al alloy as a core–shell microencapsulated phase change material: fabrication *via* dry synthesis method and its thermal stability mechanism†

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In the quest for efficient high-temperature thermal energy storage systems (TES) and power-to-heat-to-power systems (PHP), this study focuses on the development of Cu–12.8Si–20Al/Al₂O₃ core–shell microencapsulated phase change materials (MEPCMs). The Cu–12.8Si–20Al alloy, with melting point range of 738–758 °C was selected as the core PCM. Two subsequent physical methods were performed to optimize the MEPCMs: (1) uniformly coating the core with shell nanoparticles *via* a dry synthesis mechanical impact technique; (2) conducting heat oxidation in an O₂ atmosphere to foster a robust shell structure. To ascertain the optimal structure for the MEPCM, we investigated three shell variants: α-Al₂O₃, AlOOH, and a mixture of both. Significantly, the α-Al₂O₃ nanoparticles manifested a dual-layered shell, defined by an internally sintered α-Al₂O₃ nanoparticles layer and an overlying sub-nanoparticles layer. This construction enhanced the MEPCMs' thermal resilience: allowing them to withstand over 600 cycles of endothermic and exothermic phases, as well as affirming their endurance under extensive 100 h air exposure at 900 °C. The synergy between α-Al₂O₃ and AlOOH in the mixed shell revealed a pivotal role of AlOOH, which served as an adept sintering agent to enhance the MEPCM's thermal stability. In conclusion, the Cu–Si–Al/Al₂O₃ MEPCM was successfully produced as a promising candidate in high-temperature latent heat storage applications.

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Introduction

The production capacity of variable renewable energy (VRE), such as wind and concentrated solar power (CSP), continues to grow each year due to the reduction in equipment costs.^{1,2} However, VRE presents challenges in flexibly meeting power demands. Power demand must align with the amount of power supplied for a stable power supply. As such, power-to-heat-to-power (PHP) systems that store energy in inexpensive thermal energy storage (TES) materials and release it according to

energy demand are attracting attention.^{3–6} TES materials and the heat storage temperature in a PHP system are determined by the operating temperature of its heat cycle with which it will be integrated. From the perspective of the Carnot efficiency, a higher operating temperature is desirable. Therefore, high-temperature TES has recently become necessary, with proposals like the Brayton cycle for CSP operating at over 700 °C.⁷ Presently, sensible heat storage (SHS) materials, one of the TES materials, are utilized due to their low cost.^{8,9} However, SHS materials have a low energy storage density of 25 kW m⁻³,³ leading to problems such as the need for large-scale storage facilities and corresponding limitations in their application.

Latent heat storage (LHS) materials, which store latent heat accompanying phase changes of materials, are attracting attention due to their high energy storage density.³ Molten salts and alloy series materials that undergo a phase change in the high temperature range have been researched.¹⁰ Molten salts are economical (≥110 US\$ per tonne),¹¹ which has led to extensive material development¹² and application research.¹³ However, the low thermal conductivity¹⁴ and the associated inflexibility have been problematic. On the other hand, alloy-based LHS

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materials, having high thermal conductivity, high latent heat capacity, and stability against heat, are gaining attention.¹⁵ For instance, the thermal conductivities of representative high-temperature LHS materials aluminum (melting point: 660–661 °C) and copper (melting point: 1077–1083 °C) are 204 W m⁻¹ K⁻¹ and 350 W m⁻¹ K⁻¹, respectively.¹⁵ Rea *et al.* have successfully demonstrated a prototype TES system using Al–Si LHS materials.¹⁶ However, alloy-based LHS materials have a high corrosiveness,¹⁰ leading to challenges such as storage container damage, leakage, and safety concerns. Moreover, LHS materials transitioning between liquid and solid phases posed issues in handling.

Microencapsulated phase change materials (MEPCMs), a type of anti-corrosive packing,¹⁰ is considered promising due to benefits like increased heat transfer surface area of the PCM and reduced volume change associated with phase transition.^{17,18} The MEPCMs are expected to prevent corrosion and improve PCM manageability. MEPCM production methods are classified into physical, physico-chemical, and chemical.¹⁹ For high-temperature MEPCMs, physico-chemical methods are primarily utilized. Previous research has proposed several methods for preparing high-temperature MEPCMs, including sol-gel^{20,21} and sacrificial layers.²² Our research group has developed a highly durable, high-temperature MEPCMs using two-steps: (1) creating an ALOOH precursor layer on PCM microspheres using hydroxide precipitation in hot water and (2) a subsequent heat-oxidation treatment to yield a stable Al₂O₃ shell.^{23,24} These hydrolysis and heat oxidation methods were also applied to develop Al–10mass%Zn MEPCMs (melting point: 640 °C).²⁵ However, MEPCMs employing Cu-based alloy PCMs with melting points exceeding 700 °C have not been developed.

While the need for cost-effective and straightforward fabrication of MEPCMs is evident, conventional high-temperature fabrication methods, like the sol-gel approach, come with challenges. These often demand reaction control and safety management due to wet processes, which are replete with complex chemical reactions. Consequently, there has been significant interest in encapsulation method that avoids the complexities of these chemical reactions.^{26,27} Among these, spray drying, which involves spraying shell material containing solvent onto the PCM surface, has gained traction.²⁸ However, it has limitations for high-temperature MEPCMs (above 500 °C) because of restrictions in shell material types.²⁶ Furthermore, several physical methods grapple with issues such as incomplete coating of the PCM particle surface, potential leaks, degradation in performance, and diminished mechanical strength.²⁶

Given these challenges, our study takes a novel turn, introducing a Cu-based MEPCM fabricated through the combination of high-speed impact blending (HIB) and a subsequent heat oxidation treatment. The principles of HIB method are rooted in the dynamics of mechanical impact processes.²⁹ It prepares complexes of both organic and inorganic materials without relying on solvents. This method, involving dry and impact blending, depends on the dynamic interactions between particles. These interactions achieve a balance between adhesion – where particles stick; and abrasion – where

particles might be worn down or refined. Although HIB method has been used in electrode particle coating^{30,31} and particle modifications,^{32,33} its utilization in the fabrication of MEPCMs remains underexplored. Notably, there is no existing documentation on MEPCMs creation using HIB, and the mechanisms for surface coating by HIB are not fully understood. For our Cu-based alloy MEPCM, we devised a twofold fabrication process: (1) HIB treatment to achieve a uniform coat on the PCM surfaces with shell nanoparticles (NPs), either ALOOH or α -Al₂O₃, and (2) a heat-oxidation process to sinter these shell NPs, forming a stable α -Al₂O₃ shell. This dry synthesis method, which combines HIB and heat oxidation treatment, not only offers an alternative to the pitfalls of wet processes but also sets the stage for scalable and economical solutions. The fundamental physics behind this approach grants the adaptability to alter the MEPCM shell structure by modifying the composition of the shell NPs. Additionally, by optimizing the PCM composition, the method holds promise for MEPCMs suitable for a wide temperature spectrum, from the lower to the upper extremes.

Experimental

Preparation of the MEPCMs

Microspherical particles of Cu–12.8mass%Si–20mass%Al (Cu–12.8Si–20Al) alloy, prepared through spinning disk atomization (Hikari Material Industry Co. Ltd), were used as the PCM raw material. The raw material's diameter, melting temperature, and LH were 20–45 μ m, 738–758 °C (Fig. 2a and Fig. S1, ESI[†]), and 254 J g⁻¹, respectively. ALOOH NPs powder was purchased from TAIMEI Chemicals Co., Ltd (type: C06, 50%, diameter: 0.7 μ m). α -Al₂O₃ NPs powder was purchased from Kojundo Chemical Laboratory Co. Ltd (purity: 99.99%, diameter: 0.3 μ m).

Fig. 1 illustrates a schematic diagram of MEPCM preparation using a hybridization system. Core particles (Cu–12.8Si–20Al) were mixed with shell NPs (α -Al₂O₃ and ALOOH) in an 8 : 2 volume ratio using a ball mill (140 rpm, 10 min). The resulting mixture (30 g) underwent a HIB treatment (rotation speed: 50 m s⁻¹, processing time: 3 min, atmosphere: Ar) to attach the shell NPs onto the core PCM surfaces by using a HIB machine (Model: NHS-0, Nara Machinery Co. Ltd, Japan). The HIB process was carried out in five batches for each parameter set, with the fifth batch as the sample (hybridized-PCM). Subsequently, the hybridized-PCMs were subjected to heat oxidation treatment in an oxygen atmosphere (99.5% purity, flowing at 200 mL min⁻¹). The temperature was ramped from room temperature to 1000 °C at a rate of 10 °C min⁻¹ and sustained at 1000 °C for three hours. Table 1 summarizes the naming of the MEPCMs pre and post heat oxidation. The HIB treatment PCM is named hybridized PCM. The PCM encapsulated by subsequent heat oxidation treatment is named MEPCM.

Characterization

The morphologies and surface structures of the synthesized MEPCMs were examined using a field-emission scanning electron microscopy (FE-SEM) (JSM-7001FA, JEOL Ltd, Japan).





Fig. 1 Schematic diagram of the MEPCMs making using the HIB method.

Table 1 The MEPCM composition and corresponding designation pre and post heat oxidation

Shell particle composition	Initial name (before heat oxidation)	Final name (after heat oxidation)
α -Al ₂ O ₃	Hybridized-PCM-alumina	MEPCM-alumina
α -Al ₂ O ₃ -45 vol%AIOOH	Hybridized-PCM-alumina-AIOOH	MEPCM-alumina-AIOOH
AIOOH	Hybridized-PCM-AIOOH	MEPCM-AIOOH

Following the heat-oxidation treatment, the elemental distribution was evaluated by inspecting the cross-sections of the MEPCMs. These cross-sectioned specimens were polished using an ion beam cross-section polisher (IB-09010CP, JEOL Ltd, Japan) before being characterized by energy-dispersive X-ray spectroscopy (EDS) (JSM-7001FA, JEOL Ltd, Japan). The phase composition was determined using powder X-ray diffraction (XRD) equipped with one-dimensional silicon strip sensor (MiniFlex600, Cu K α , Rigaku, Japan). The prominent XRD peaks were identified corresponding to α -Al₂O₃ (JCPDS card

No. 01-075-6776), AIOOH (JCPDS card No. 01-073-6509), Si (JCPDS card No. 00-005-0565), Al₄Cu₉ (JCPDS card No. 00-024-0003), and Cu₂O (JCPDS card No. 01-077-7719). The melting point and LHS capacity of the MEPCMs were quantified using a differential scanning calorimetry (DSC) analyzer (DSC-823, Mettler Toledo, USA). The cyclic durability over 600 cycles was assessed through thermal cyclic tests in a custom-built quartz furnace capable of lateral movement between two points. The high-temperature stability of MEPCMs was conducted by exposing them to an air environment at 900 °C for 100 h in a muffle furnace.



Fig. 2 SEM images of the MEPCMs before and after the treatment. (a) Raw material, (b) hybridized-PCM-alumina, (c) hybridized-PCM-alumina-AIOOH, (d) hybridized-PCM-AIOOH (e) MEPCM-alumina, (f) MEPCM-alumina-AIOOH, and (g) MEPCM-AIOOH. MEPCMs in (e)–(g) were derived by heat oxidation treatment of the corresponding HIB-treated MEPCMs from (b)–(d) at 1000 °C for three hours.



Results & discussion

Observation of MEPCMs

Fig. 2 displays the SEM images of MEPCMs, synthesized using a combination of HIB and heat-oxidation treatment at 1000 °C for three hours. All the hybridized-PCMs consistently exhibit spherical shapes (Fig. 2b–d). Following the HIB treatment, the presence of shell NPs (α -Al₂O₃, ALOOH) on the PCM core particle is apparent (Fig. 2b–d). Specifically, the α -Al₂O₃ shell NPs adheres to the particle's surfaces in a spherical form (Fig. 2b), while the ALOOH shell NPs adheres in a plate-like manner (Fig. 2d). Fig. 2c depicts a combination of rounded α -Al₂O₃ NPs and flat ALOOH NPs attached to the core PCM particle's surfaces.

Following the heat oxidation treatment at 1000 °C for three hours, all samples preserved their spherical shape, even when processed above the melting point of the Cu-12.8Si-20Al alloy (Fig. 2e–g). No leakage of liquid PCM or damage to the shell was detected, indicating the heat oxidation treatment successfully sintered the shell NPs, resulting in the formation of dense shells. The MEPCM-alumina is uniformly encapsulated by α -Al₂O₃ shell NPs (Fig. 2e), with minimal variations from its state prior to the heat oxidation (Fig. 2b). Fig. 2g illustrates the MEPCM-ALOOH encapsulated by a dense, seamlessly sintered surfaces. Given that ALOOH dehydrates to form Al₂O₃ around 500 °C,³⁴ this sintered layer is the result of ALOOH dehydration. Lastly, the surfaces of MEPCM-alumina-ALOOH are dominated by spherical α -Al₂O₃ shell NPs (Fig. 2f).

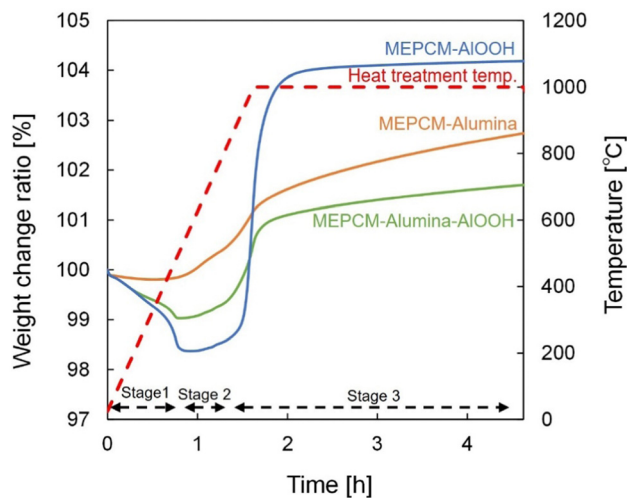
Heat oxidation of hybridized-PCM

Hybridized-PCMs formed a stable and dense α -Al₂O₃ shell by sintering the shell nanoparticles through heat oxidation treatment. Drawing from the findings of Nomura *et al.*, when an Al-based PCM coated with ALOOH undergoes heat oxidation around 1000 °C, a dense and robust Al₂O₃ shell formed on the PCM surfaces. This formation is primarily due to the dehydration of ALOOH coupled with the partial oxidation of Al.²³ Similarly, for hybridized-PCM that was encapsulated with ALOOH/Al₂O₃ shell nanoparticles, subjecting them to a heat oxidation treatment at 1000 °C instigates the dehydration of ALOOH and the favorable oxidation of specific Al quantities within the PCM (Fig. S2, ESI†). These synergistic reactions facilitate the sintering of shell nanoparticles, leading to the formation of a stable α -Al₂O₃ shell.

Fig. 3 delineates the thermogravimetric (TG) curves of hybridized-PCMs during the heat-oxidation. Apparently, the TG curves of hybridized-PCMs, when coated with ALOOH nanoparticles (Fig. 3), align closely with the trends of MEPCMs having Al-based alloy PCM surfaces, as coated by the hydroxide precipitation treatment.^{24,35,36} Here, the heat oxidation process can be classified into three distinct stages:³⁵

(i) Stage 1 – room temperature to 580 °C: this is where dehydration of ALOOH takes place,³⁴ resulting in discernable mass reduction. This decrease becomes more pronounced with an increasing concentration of ALOOH NPs.

(ii) Stage 2 – spanning 580 °C to 738 °C (pre-melting point of PCM): oxidation manifests on the solid-state PCM core



Sample name	Stage 1 [mass%]	Stage 2 [mass%]	Stage 3 [mass%]
MEPCM-Alumina	-0.05	0.34	2.40
MEPCM-Alumina-AIOOH	-0.95	0.18	2.44
MEPCM-AIOOH	-1.62	0.11	5.67

Fig. 3 TG curves of hybridized-PCM during heat-oxidation treatment.

surfaces, resulting in the development of a thin oxide film over the PCM core particle.

(iii) Stage 3 – beyond 738 °C (post-melting point of PCM): here, a significant gain is observed, attributable to the acceleration oxidation of Al. Simultaneously, the swift volumetric expansion, might induce tiny cracks in the oxide film. In the PCM transitions to a liquid state, it swiftly permeates these cracks, interacting with oxygen, eventually giving rise to a stable oxide.²⁴

The corresponding yielded composition post-heat oxidation treatment is further corroborated by the XRD pattern result. Fig. 4 exhibits the comparative analysis of the XRD patterns between hybridized-PCMs and developed MEPCMs. Both Al₄Cu₉ and Si in PCM were detected across all samples. Notably, ALOOH was discernible in both hybridized-PCM-ALOOH and hybridized-PCM-alumina-ALOOH. In contrast, in the XRD patterns for MEPCM-ALOOH and MEPCM-alumina-ALOOH, ALOOH peaks were absent, while clear α -Al₂O₃ peaks were present. Such observations can be linked to the transformation of the ALOOH shell nanoparticles into α -Al₂O₃, a conversion triggered during the 500 °C heat-oxidation treatment (Fig. 3). Further emphasizing this transformation, the intensity of the α -Al₂O₃ peak in MEPCM-alumina, MEPCM-ALOOH, and MEPCM-alumina-ALOOH was markedly higher compared to their hybridized-PCM counterparts. Additionally, the absence of copper oxide in hybridized-PCMs, followed by its detection post heat-oxidation, implies the oxidation of a portion of the PCM particles during the treatment.

Thermal energy storage and release performance

As illustrated in Fig. 5, the LHS capacity of the MEPCMs increases with the decreasing addition of ALOOH. The melting temperature of the synthesized MEPCMs was approximately





Fig. 4 XRD patterns of the MEPCMs after heat-oxidation treatment.

775–780 °C, whereas the solidification temperature ranged from 730–737 °C. The solidification temperature of all MEPCMs was about 45 °C lower than the melting temperature. Such supercooling phenomenon has been reported for other high-temperature MEPCMs.^{35,37} During solidification, both MEPCM-alumina and MEPCM-alumina-AlOOH exhibited a minor peak around 750 °C. Notably, this peak was absent in the MEPCM-AlOOH. Both MEPCM-alumina and MEPCM-alumina-AlOOH presented similar DSC curves. However, the DSC curve for MEPCM-AlOOH was distinctly different. The concentration of Al in the PCM of MEPCM-AlOOH likely diminished,

transitioning from 20% down to approximately 15% due to the significant oxidation of Al in the PCM during the heat oxidation (Fig. 3). Such oxidation elevated the onset temperature of melting for MEPCM-AlOOH (Fig. S1, ESI†). From these observations, it becomes clear that the composition of the shell NPs plays a pivotal role in determining the LHS characteristics of the MEPCMs.

Thermal stability and cyclic durability

To assess the thermal stability of the MEPCMs, these materials were subjected to extensive exposure in air at 900 °C. The structural response and changes in LHS capacity over this period are presented in Fig. 6a and b. Notably, both the MEPCM-alumina and MEPCM-alumina-AlOOH samples exhibited robust structural integrity, retaining their shape even after the extensive 100 h test period. This ability to withstand high-temperature environments speaks to the strength of the binding forces within these materials, as well as the resilience of their microstructures against thermal degradation.

In stark contrast, the MEPCM-AlOOH displayed shell damage around 40 h. Such damage can be attributed to weaker intermolecular forces or potential imperfections in its microstructure, rendering it more susceptible to thermal stresses. Quantifying the LHS capacity post the 100 h thermal test, values recorded were 198 J g⁻¹ for MEPCM-alumina, 197 J g⁻¹ for MEPCM-alumina-AlOOH, and 159 J g⁻¹ for MEPCM-AlOOH (Fig. 6b). When juxtaposed with their initial capacities, these represent reductions to 89.6, 93.8, and 78.3%, respectively. The data asserted that MEPCM-alumina and MEPCM-alumina-AlOOH exhibit superior thermal stability when exposed to high-temperature air.

Progressive heating and cooling cycles were, then, conducted in order to confirm both the LHS capacity and cyclic durability of the MEPCMs. During this cyclic test, the MEPCMs



Fig. 5 DSC curve of various AlOOH addition MEPCMs after heat oxidation treatment. (T_m : melting temperature of MEPCM, T_s : solidification temperature of MEPCM, ΔH : latent heat storage capacity).





Fig. 6 (a) SEM images and (b) LHS capacity of heat-oxidation-treated samples, following thermal endurance test for 40 h and 100 h at 900 °C in air. Samples include MEPCM-alumina, MEPCM-alumina-AIOOH, and MEPCM-AIOOH. Samples with visible shell damage are marked with red circles.

were heated to 865 °C and cooled to 600 °C over 600 cycles. The heating and cooling average rates were approximately 58 °C min⁻¹ and 28 °C min⁻¹, respectively. Fig. 7b and c depict the SEM images of the MEPCMs after 100 and 600 cycles, respectively. Notably, Fig. 7b reveals structural compromises

in the MEPCM-AIOOH after just 100 cycles: ruptured shells and evident PCM leakage. Such degradation can be traced back to the inherent stresses introduced by volumetric changes – expansion and contraction – during the phase transitions of the PCM particles.³⁸ However, in a testament to their



Fig. 7 SEM images of (a) heat-oxidation-treated samples after (b) 100 and (c) 600 thermal cycles. Samples include MEPCM-alumina, MEPCM-alumina-AIOOH, and MEPCM-AIOOH. Samples with visible shell damage are marked with red circles.



superior structural stability, both MEPCM-alumina and MEPCM-alumina-AOOH samples, as seen in Fig. 7c, exhibited remarkable resilience. Even after an exhaustive 600 cycles, they retained their original morphology without any discernible cracks or indications of PCM leakage.

Fig. 8 presents the heating (a) – cooling (b) DSC curves of MEPCM-alumina, MEPCM-alumina-AOOH, and MEPCM-AOOH after 600 cycles of repeated melting and solidification. In Fig. 8b, the solidification temperature remains unchanged after cyclic testing, at approximately 731–738 °C. In contrast, the LHS of melting after 600 cyclic testing is 208 J g⁻¹ for MEPCM-alumina, 203 J g⁻¹ for MEPCM-alumina-AOOH and 156 J g⁻¹ for MEPCM-AOOH, decreasing to 91.2, 98.5, and 81.3% of those before cyclic testing, respectively (Fig. 8c). Thus, MEPCM-alumina and MEPCM-alumina-AOOH maintain high LHS capacity, nearly identical to the as-prepared MEPCMs. To elucidate the compositional changes and oxidation of Cu–Si–Al PCM after 600 cycles, XRD analysis was undertaken (Fig. S3, ESI†). The findings suggest that the majority of the PCM in MEPCM-alumina and MEPCM-alumina-AOOH remained intact, resisting oxidation throughout the cyclic testing. In contrast, the peak intensity of α -Al₂O₃ in MEPCM-AOOH after 600 cyclic tests was higher than that in MEPCM-AOOH before the cyclic tests. This suggests that the decrease in the LHS capacity of MEPCM-AOOH was due to shell breakage (Fig. 7) and subsequent oxidation of the PCM.

Further insights into the MEPCM shell cross-section of the as-prepared MEPCMs were sought to understand the encapsulation mechanism and associated mechanical properties. The observation analyses revealed negligible structural changes pre and post 600 cyclic tests (Fig. 9 and Fig. S4–S6, ESI†). Of these, the α -Al₂O₃ shells of both MEPCM-alumina and MEPCM-alumina-AOOH were discerned to have a dual-layered composition: an inner layer comprised of tiny sintered NPs, indicative

of originating from α -Al₂O₃ NPs, and an outer layer consisting of aggregated NPs. The sintered shell NPs layer was about 300–400 nm thick, consisting of tiny sintered NPs around 150 nm in size, specifically α -Al₂O₃ NPs. The thickness of the shell NPs layer was about 1.2 μ m. Voids were discerned within the shell NPs layer of MEPCM-alumina, while in MEPCM-alumina-AOOH, the α -Al₂O₃ NPs acted as a binder, promoting inter-particle connectivity during its transformation into α -Al₂O₃ through the sintering process.

Within the shell layer of MEPCM-alumina-AOOH, the α -Al₂O₃ shell NPs seemed to have undergone a mild fusion (Fig. S6, ESI†). This mild fusion of the α -Al₂O₃ NPs indicates that the addition of AOOH to MEPCM-alumina-AOOH acted as a binding agent. It aided in linking adjacent α -Al₂O₃ NPs as they transformed into α -Al₂O₃ during the sintering process initiated by the heat oxidation treatment. The dual-layered shell structures of MEPCM-alumina and MEPCM-alumina-AOOH exhibited high durability, retaining their integrity even after 600 cyclic tests (Fig. S4, ESI†). This shell structure effectively protected against PCM corrosion and oxidation and retained 98.5% (MEPCM-alumina-AOOH) or 91.2% (MEPCM-AOOH) of LHS capacity after 600 cyclic tests.

In contrast, the MEPCM-AOOH shell (Fig. 9 and Fig. S7, ESI†) was composed of alumina-sintered bodies, forming a dense structure where individual and sintered shell NPs could not be observed. The thickness of the alumina-sintered layer was approximately 0.2–1 μ m, and differences in thickness and density were seen depending on the location. Some areas showed the presence of voids or fragile areas prone to cracking within the sintered body. Compared to MEPCM-alumina and MEPCM-alumina-AOOH, MEPCM-AOOH had lower cyclic durability. It is thought that the sintered shell particle layers of MEPCM-alumina and MEPCM-alumina-AOOH, the sudden expansion pressure associated with the melting of PCM, were



Fig. 8 LHS capacity of as-prepared MEPCM-alumina, MEPCM-alumina-AOOH, and MEPCM-AOOH samples after 100 cycles, 300 cycles, and 600 cyclic tests. (a) and (b) DSC curves for melting and solidification, respectively. (c) LHS capacity of all samples. (T_m : melting temperature of MEPCM; T_s : solidification temperature of MEPCM; ΔH : latent heat storage capacity).





Fig. 9 Cross-sectional SEM images of the MEPCMs before and after 600 cyclic tests. Samples include MEPCM-alumina, MEPCM-alumina-AIOOH, and MEPCM-AIOOH. Orange lines indicate the interface between the PCM core and its shell.

distributed by many sintered NPs, contributing to their high durability. On the other hand, since the MEPCM-AIOOH shell is formed from a large sintered body,²⁴ the breakage was likely caused by the failure to distribute the rapid expansion pressure caused by the melting of the PCM, which was received by the vulnerable areas. The inhomogeneity of the MEPCM-AIOOH sintered shell may have contributed to its low durability.

MEPCM-alumina-AIOOH demonstrated higher cyclic durability than MEPCM-alumina (Fig. 8). In the shell NPs layer of MEPCM-alumina-AIOOH, the added AIOOH acted as a sintering agent, causing the α - Al_2O_3 NPs to fuse. This fusion made the

shell particle layer denser and is postulated to have provided more excellent mechanical durability.

Nomura *et al.* reported that Al-Si/ Al_2O_3 core-shell MEPCMs, prepared by hydrolysis and heat oxidation, exhibit shell formation through two fundamental mechanisms: (1) crack caused by expansion pressure resulting from melting of the PCM core and crack repair by PCM oxidation and (2) oxidation of the PCM surfaces due to oxygen diffusion.³⁶ Similarly, the shell of Cu-Si-Al/ Al_2O_3 core-shell MEPCMs, prepared *via* HIB treatment and heat oxidation, is thought to form through a comparable mechanism. Fig. 10 presents the expected

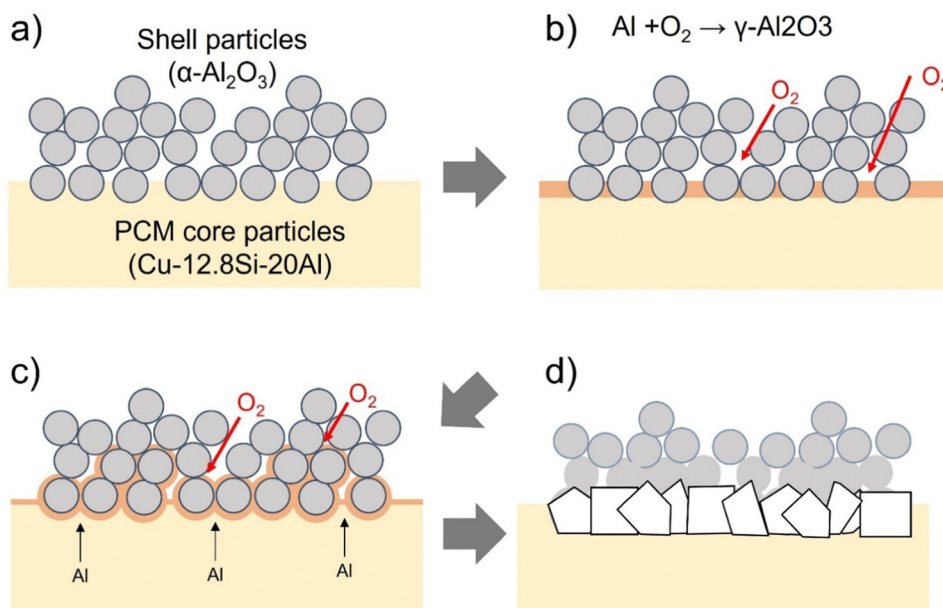


Fig. 10 Schematic diagram of shell formation by heat oxidation treatment of MEPCM-alumina. (a) After HIB treatment, (b) oxidation of PCM core particles and formation of oxide film, (c) diffusion of PCM due to melting of PCM, and (d) sintered α - Al_2O_3 shell.



schematic diagram illustrating the shell formation process during the heat oxidation treatment of MEPCM-alumina. The TG curve of MEPCM-alumina shows a significant weight shift between 538–1000 °C (Fig. 3). When the heat oxidation temperature is below the melting point (580–738 °C), the PCM core particles (Cu–12.8Si–20Al) undergo oxidation and form a γ -Al₂O₃ oxide film on their surface. If the heat oxidation temperature exceeds the melting point (738–1000 °C), two phenomena occur: (1) the initiation of cracks and subsequent PCM leakage and (2) crack repair through PCM oxidation. These processes are repeated, leading to the formation of an Al₂O₃ shell.³⁶ Near 950 °C, the generated γ -Al₂O₃ transitions to α -Al₂O₃ formation, followed by the sintering of α -Al₂O₃ shell NPs, which results in the formation of a sintered shell NPs layer.

Conclusions

In conclusion, this study has successfully developed a Cu–12.8mass%Si–20mass%Al@Al₂O₃ MEPCM with excellent thermal cyclic durability and thermal stability by HIB treatment followed by heat oxidation treatment. The effect of shell NPs composition on the shell structure and durability of MEPCMs was investigated. The primary results are as follows.

(1) The HIB treatment successfully enabled a uniform coating of the PCM surfaces with shell NPs. After the HIB process, heat oxidation caused the formation of shell NPs sintering and the merging of shell NPs, leading to the construction of a stable α -Al₂O₃ shell.

(2) MEPCM-alumina and MEPCM-alumina-AIOOH, utilizing α -Al₂O₃ NPs as a shell, formed a dual-layered shell comprised of a sintered shell NPs layer and a shell-NPs layer. These demonstrated impressive durability over 600 cycles and superior thermal stability in high-temperature air. MEPCM-AIOOH utilizing AIOOH as seamless and dense α -Al₂O₃ shell was formed.

(3) Incorporating AIOOH NPs into the shell composition enhanced the thermal durability of the shell. During the heat oxidation, AIOOH acted as a binding agent, facilitating the formation of a denser α -Al₂O₃ shell.

The MEPCM fabrication method that employs HIB treatment offers potential for broader application in the microencapsulation of diverse PCMs, as it is not reliant on specific PCMs or shell NPs. Furthermore, the combined strategy of HIB and heat oxidation treatment holds benefits for mass production and cost efficiency due to its physical process.

Author contributions

Masahiro Aoki: data curation, investigation, writing. Melbert Jeem: data curation, investigation, writing – review & editing. Yuto Shimizu: investigation. Takahiro Kawaguchi: investigation. Minako Kondo: investigation. Tomokazu Nakamura: investigation. Chihiro Fushimi: investigation. Takahiro Nomura: investigation, funding acquisition, project administration, supervision, writing, writing – review & editing.

Data availability

The data supporting this study are available from the corresponding author upon reasonable request.

Abbreviations

ΔH	Latent heat storage capacity
CSP	Concentrated solar power
DSC	Differential scanning calorimeter
EDS	Energy-dispersive X-ray spectroscopy
HIB	High-speed impact blending
LH	Latent heat
LHS	Latent heat storage
MEPCM	Microencapsulated phase change material
SEM	Scanning electron microscopy
SHS	Sensible heat storage
TES	Thermal energy storage
TG	Thermogravimetric analysis
T_m	Melting temperature
T_s	Solidification temperature
PHP	Power to heat to power
VRE	Variable renewable energy
NPs	Nanoparticles

Conflicts of interest

The authors declare no conflict of interest.

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References

- Renewable energy – powering a safer future, <https://www.un.org/en/climatechange/raising-ambition/renewable-energy>, (accessed February 21, 2023).
- M. K. DeValeria, E. E. Michaelides and D. N. Michaelides, *Energy*, 2020, **190**, 116440.
- S. Wu, C. Zhou, E. Doroodchi, R. Nellore and B. Moghtaderi, *Energy Convers. Manage.*, 2018, **168**, 421–453.
- Y. He, S. Guo, J. Zhou, F. Wu, J. Huang and H. Pei, *Energy Convers. Manage.*, 2021, **229**, 113779.
- T. Uchino, T. Yasui and C. Fushimi, *Energy Convers. Manage.*, 2021, **243**, 114366.
- S. Liu, H. Bai, P. Jiang, Q. Xu and M. Taghavi, *J. Energy Storage*, 2022, **50**, 104577.
- L. Sang, X. Lv, Y. Wang, J. Huang and Y. Wu, *J. Energy Storage*, 2023, **61**, 106724.



- 8 M. Tawalbeh, H. A. Khan, A. Al-Othman, F. Almomani and S. Ajith, *Int. J. Thermofluids*, 2023, **18**, 100326.
- 9 J. Liu, Z. Chang, L. Wang, J. Xu, R. Kuang and Z. Wu, *ACS Omega*, 2020, **5**, 19236–19246.
- 10 M. Liu, X. Zhang, J. Ji and H. Yan, *J. Energy Storage*, 2023, **63**, 107005.
- 11 C. Li, Q. Li and R. Ge, *Energy Rep.*, 2022, **8**, 12740–12764.
- 12 Y. T. Wu, Y. Li, N. Ren, R. P. Zhi and C. F. Ma, *Sol. Energy Mater. Sol. Cells*, 2018, **176**, 181–189.
- 13 M. Opolot, C. Zhao, M. Liu, S. Mancin, F. Bruno and K. Hooman, *Renewable Sustainable Energy Rev.*, 2022, **160**, 112293.
- 14 B. Xu, P. Li and C. Chan, *Appl. Energy*, 2015, **160**, 286–307.
- 15 S. C. Costa and M. Kenisarin, *Renewable Sustainable Energy Rev.*, 2022, **154**, 111812.
- 16 J. E. Rea, C. J. Oshman, A. Singh, J. Alleman, G. Buchholz, P. A. Parilla, J. M. Adameczyk, H. N. Fujishin, B. R. Ortiz, T. Braden, E. Bensen, R. T. Bell, N. P. Siegel, D. S. Ginley and E. S. Toberer, *Energy Convers. Manage.*, 2019, **199**, 111992.
- 17 R. Al-Shannaq, J. Kurdi, S. Al-Muhtaseb and M. Farid, *Sol. Energy*, 2016, **129**, 54–64.
- 18 G. Peng, G. Dou, Y. Hu, Y. Sun and Z. Chen, *Adv. Polym. Technol.*, 2020, **2020**, 9490873.
- 19 A. Jamekhorshid, S. M. Sadrameli and M. Farid, *Renewable Sustainable Energy Rev.*, 2014, **31**, 531–542.
- 20 X. Wu, M. Fan, S. Cui, G. Tan and X. Shen, *Sol. Energy Mater. Sol. Cells*, 2018, **178**, 280–288.
- 21 M. D. Romero-Sanchez, R.-R. Piticescu, A. M. Motoc, F. Aran-Ais and A. I. Tudor, *Manuf. Rev.*, 2018, **5**, 8.
- 22 B. Zhao, Y. Guo, C. Wang, L. Zeng, K. Gao, N. Sheng, E. Gariboldi and C. Zhu, *Chem. Eng. J.*, 2023, **457**, 141352.
- 23 T. Nomura, N. Sheng, C. Zhu, G. Saito, D. Hanzaki, T. Hiraki and T. Akiyama, *Appl. Energy*, 2017, **188**, 9–18.
- 24 N. Sheng, C. Zhu, H. Sakai, Y. Hasegawa, T. Akiyama and T. Nomura, *Sol. Energy Mater. Sol. Cells*, 2019, **200**, 109925.
- 25 T. Kawaguchi, H. Sakai, N. Sheng, A. Kurniawan and T. Nomura, *Appl. Energy*, 2020, **276**, 115487.
- 26 Y. Huang, A. Stonehouse and C. Abeykoon, *Int. J. Heat Mass Transfer*, 2023, **200**, 123458.
- 27 G. Leng, G. Qiao, Z. Jiang, G. Xu, Y. Qin, C. Chang and Y. Ding, *Appl. Energy*, 2018, **217**, 212–220.
- 28 G. Alva, Y. Lin, L. Liu and G. Fang, *Energy Build.*, 2017, **144**, 276–294.
- 29 W. J. Tang, Z. Y. Fu, J. Y. Zhang, W. M. Wang, H. Wang, Y. C. Wang and Q. J. Zhang, *Powder Technol.*, 2006, **167**, 117–123.
- 30 T. Kawaguchi, H. Nakamura and S. Watano, *Powder Technol.*, 2017, **305**, 241–249.
- 31 T. Kawaguchi, H. Nakamura and S. Watano, *Powder Technol.*, 2018, **323**, 581–587.
- 32 J. Yang, A. Sliva, A. Banerjee, R. N. Dave and R. Pfeffer, *Powder Technol.*, 2005, **158**, 21–33.
- 33 R. Pfeffer, R. N. Dave, D. Wei and M. Ramlakhan, *Powder Technol.*, 2001, **117**, 40–67.
- 34 Boehmite Powder/Products, <https://www.taimei-chem.co.jp/product/15.html>, (accessed September 6, 2023).
- 35 Y. Shimizu, T. Kawaguchi, H. Sakai, K. Dong, A. Kurniawan and T. Nomura, *Sol. Energy Mater. Sol. Cells*, 2022, **246**, 111874.
- 36 T. Nomura, C. Zhu, N. Sheng, G. Saito and T. Akiyama, *Sci. Rep.*, 2015, **5**, 9117.
- 37 T. Kawaguchi, J. Yoolerd, H. Sakai, Y. Shimizu, A. Kurniawan and T. Nomura, *Sol. Energy Mater. Sol. Cells*, 2022, **237**, 111540.
- 38 T. Nomura, J. Yoolerd, N. Sheng, H. Sakai, Y. Hasegawa, M. Haga, G. Saito and T. Akiyama, *Sol. Energy Mater. Sol. Cells*, 2018, **187**, 255–262.

