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Crystallization by local cooling of supercooled sodium acetate trihydrate composites for long-term heat storage

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Highlights

- Crystallization of supercooled SAT composites started between -9°C and -24°C
- Sodium acetate-water mixtures released heat in two stages during crystallization
- Higher, more uniform crystallization temperatures in samples with rusty steel profiles
- Carboxymethyl cellulose and polymeric additives improved heat release during crystallization
- Proof of applicability of CO₂ evaporation and Peltier element devices to initiate crystallization

Abstract

Sodium acetate trihydrate (SAT) can be used for long-term heat storage in buildings by utilizing its ability to supercool stably to ambient temperatures while preserving its heat of fusion. Additives are used to stabilize the SAT and avoid phase separation. A reliable method for initializing the solidification of supercooled SAT composites is needed to operate a heat storage unit based on supercooled SAT. The crystallization temperatures of SAT composites during cooling were therefore investigated and experiments carried out using methods applying local cooling in a small part of prototype heat storage units to initialize crystallization. To find the crystallization temperatures of SAT composites, supercooled samples were cooled down in a freezer. The influence of rusty metal parts submerged in melted SAT composite samples and various periods at rest in a supercooled state were investigated with regard to supercooling stability and crystallization temperature. Carboxymethyl cellulose, extra water, liquid polymer HD 310, metal-based graphite flakes, and silicone oil were applied as additives to form the different SAT compositions. Samples with 60 g SAT in glass jars were subject to repeated heating and cooling cycles. It was found that samples containing steel profiles crystallized in the range of -9 to -15°C , while SA-water mixtures without steel profiles cooled down to -24°C before crystallizing. Furthermore, samples with carboxymethyl cellulose and liquid polymer HD 310 showed a greater temperature rise during crystallization, which is in accordance with previous findings on heat contents. SA-water mixtures showed a second minor temperature peak at temperatures below -20°C , when the sample was cooled down again after the first temperature rise. Devices were developed to initialize the crystallization of supercooled SAT composites in prototype heat storage units using rapid local cooling of the SAT composite. We successfully initialized crystallization by evaporating pressurized liquid carbon dioxide in a small chamber on one side of the PCM container and by using Peltier elements. Our experiments showed that the controlled initialization of crystallization by cooling is feasible for all the SAT composites investigated.

Keywords: Sodium acetate trihydrate, stable supercooling, long-term heat storage, CO₂ evaporation, Peltier elements.

Nomenclature

Dimensional variables

T	temperature (°C)
m	mass (kg)
t	time (min)

Greek letters

ΔT	temperature difference (K)
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Subscripts

C	cooling
CR	crystallization
Ref	reference
Steel plate	steel plate attached to a Peltier element
Heat sink	heat sink attached to a Peltier element
PCM	phase change material
Surface	the surface of the PCM container
CO ₂	carbon dioxide evaporation chamber

Abbreviations

CMC	carboxymethyl cellulose
SA	sodium acetate
SAT	sodium acetate trihydrate
CO ₂	carbon dioxide
PCM	phase change material
wt.	weight

1. Introduction

1.1. Long-term heat storage with sodium acetate trihydrate

Due to the mismatch between solar energy resources and the demand of thermal household services, long-term storage of heat is essential for solar heating systems that aim to cover the heat demand with solar fractions in the range of 80–100%. The concept of a novel compact seasonal heat store based on supercooling of Sodium Acetate Trihydrate (SAT) has therefore been defined in previous research activities at the Technical University of Denmark [1].

SAT, chemical formula $\text{CH}_3\text{COONa} \cdot 3\text{H}_2\text{O}$, is a salt hydrate with a melting point of 58 °C. It has been identified as a potential heat storage material due to its relatively high heat of fusion (264 kJ kg⁻¹ [2]; 237–243 kJ kg⁻¹ [3];) and its relatively high specific heat capacities in both liquid and solid phases [4]. The density of solid SAT in a closed sample that has solidified from supercooled state has been determined to be 1.24–1.28 kg L⁻¹ [5]. SAT is attractive for domestic heat storage applications because of its high volumetric heat storage capacity (~500 kJ L⁻¹ from 20 °C in solid state to 80 °C in liquid state), the low price and its non-toxic properties. Reviews on thermal energy storage with PCM [6], [7] therefore list SAT among the salt hydrates identified for domestic applications.

Supercooling is a phenomenon in which a substance in liquid state cools below its solidifying temperature without solidifying. SAT can be used for long-term heat storage because of its ability to supercool stably to ambient temperatures while preserving its heat of fusion [2], [8], [9]. Once heat is in demand, the solidification of the supercooled solution can be triggered and the preserved heat of fusion will be released as it crystallizes. This has been demonstrated by Dannemand et al. with both flat [10] and cylindrical [11] prototype heat storage units. Flat units have also been used to demonstrate a segmented heat storage in a solar heating system for domestic heat supply [12]. Deng et al. [13] have investigated the performance of a hot water tank that makes use of the stable supercooling of SAT in the mantle.

1.2. Sodium acetate trihydrate composites

For utilization of SAT in domestic heat stores, volumes of several cubic meters are needed for solar heating systems with yearly solar fractions close to 1 [1]. For economic reasons chemically pure SAT cannot be used. Instead, industrial SAT products were successfully applied in previous research [10], [12], [14].

Additives are needed to stabilize the SAT in its liquid phase. The solubility of sodium acetate (SA) in the corresponding crystal water at 58 °C (the melting point of SAT) is not high enough to dissolve all anhydrous salt (experimentally measured by Green [15]). Moreover, as an incongruently melting salt hydrate, SAT suffers from phase separation over repeated heating and cooling cycles and over long periods in supercooled state. Phase separation is caused by the density difference between SA and salt water. When SA molecules settle down to the bottom of a SAT volume and a leaner salt-water solution remains in the upper part of the volume, full reformation of SAT crystals can no longer be achieved [16]. This effect reduces the material's heat storage capacity.

To overcome phase separation, Furbo and Svendsen [17] suggested adding extra water to the salt hydrate. However, tests with prototype storage units [10], [11] showed a decrease in heat capacity after a number of storage cycles had been applied. Thickening agents were therefore used to increase the viscosity of the liquid solution and avoid SA to settle at the bottom of storage units. Carboxymethyl cellulose (CMC) and Xanthan rubber were investigated by Cabeza et al. [3] and Dannemand et al. [15] for this purpose. Liquid polymer solutions have also been identified as able to

reduce the phase separation effect by increasing the solubility of SAT in the liquid solution [19]. A detailed study on the stability and the heat content of supercooled SAT with various additives was conducted by Kong et al. [14]. Composites containing CMC and the liquid polymer solution HD 310 were found to provide the highest heat of fusion.

Graphite flakes for thermal conductivity enhancement have also been investigated [18]. Oils (paraffin, silicone) that do not mix with SAT have been added to increase heat transfer by filling insulating cavities formed during the solidification and contraction of the SAT composition.

1.3. Crystallization of sodium acetate trihydrate

To use SAT as a heat storage material, we need greater understanding of the factors that influence supercooling stability, nucleation mechanisms and crystal growth.

1.3.1. Supercooling stability

The crystallization process from pure SA aqueous solution to SAT requires a large degree of supercooling; earlier research found a crystallization temperature of $-15\text{ }^{\circ}\text{C}$ [20]. The behaviour of SA aqueous solutions has been experimentally investigated and can be explained using phase diagrams [4], [15].

The difference between the actual concentration and the solubility concentration of a solution at a given temperature is called the supersaturation. When an SA aqueous solution supercools, it becomes supersaturated without forming crystals. Supersaturation is the driving force for solution crystallization processes, so a supercooled SA aqueous solution stays in a meta-stable state until nucleation takes place (start of the crystallization process).

In most latent heat storage applications, this phenomenon is unwanted [6], [21]. Measures have even been developed to deliver nucleation centres in order to reduce the supercooling of SAT [22], [23]. Nevertheless, for long-term heat storage, the utilization of supercooled salt hydrates has been studied since the late 1920s [24].

Additives were used to overcome phase separation in liquid SAT and to increase the thermal conductivity of solid SAT. Since pure SAT is too expensive for heat storage applications, impure low-cost SAT products were used to realize large heat storage units in a cost-effective way. And it may well be that the available information on the crystallization properties of SAT, such as phase diagrams, does not apply to SAT composites.

1.3.2. Nucleation

Nucleation is the birth of new crystal nuclei. Mullin [25] defined two kinds of nucleation: either from solution (primary nucleation) or in the presence of existing crystals (secondary nucleation).

For primary nucleation, Mullin defined the formation of crystal nuclides from a homogenous fluid as homogeneous nucleation. This is not a common event, because supercooled solutions are probably seeded without our knowledge by the presence of atmospheric dust during material preparation. Because the rate of nucleation of a solution can be affected considerably by the presence of such impurities, this so-called heterogeneous nucleation leads to “spontaneous” nucleation events, the mechanisms of which are not fully understood [26].

A supersaturated solution nucleates much more readily when crystals of the solute are already present or added [25]. Seed crystal injection into a supercooled SAT solution is a secondary nucleation method, which has proved reliable even where there is less supercooling [4], [12].

1.3.3. Crystal growth

As soon as stable nuclei have been formed in a supersaturated solution, they begin to grow into crystals of visible size. Dietz et al. [27] and Munakata and Nagata [28] have measured the crystal growth rate of supercooled SA-aqueous solutions in tubes. The growth rates were found to be solely a function of the solution concentration and the degree of supercooling. Measured and calculated crystal growth rates have been evaluated by Ma et al. [29], but we do not yet have parameters for the degree of supercooling or models for crystal growth of melted SAT composite material.

1.4. Mechanisms for the controlled initialization of crystallization

Heat storage applications that make use of the stable supercooling of SAT need reliable mechanisms for starting crystallization. Several possibilities were tested in laboratory-scale experiments, fewer in setups with volumes of 100 L or more:

- Adding seed crystals is a common method of initializing the crystallization in SAT in laboratory-scale setups [4], [20], [27]. Crystallization with seed crystals works as long as the crystals do not melt and become inactive [30]. Seed crystal injection in large storage units has been demonstrated by some of the present authors [12].
- Triggering crystallization using ultrasonic waves has been reported as successfully tested [31], but other researchers [32] have stated that it failed.
- Local cooling of SAT in liquid phase down to its crystallization temperature (primary, heterogeneous nucleation). We demonstrated how the cooling power resulting from the endothermic process of evaporating carbon dioxide (CO₂) could supply a temperature low enough to initiate nucleation in supercooled SAT composites in prototype storage units. Previous studies have shown that SAT will spontaneously crystallize between $-15\text{ }^{\circ}\text{C}$ and $-25\text{ }^{\circ}\text{C}$ [20]. The phase diagram of CO₂ shows a triple point at 5.1 bars at $-57\text{ }^{\circ}\text{C}$. At atmospheric pressure, the phase change is directly from solid to gas at $-78\text{ }^{\circ}\text{C}$. This indicates that the temperatures resulting from evaporating CO₂ may be low enough to reach the temperature where SAT will crystallize. Dannemand et al. have previously tested prototype heat storage units utilizing the stable supercooling of SAT composites. In their laboratory investigations, they initiated nucleation of SAT in storage units by evaporating CO₂ in a small chamber on one side of the PCM unit [1], [10], [33]. The method of using evaporation of CO₂ is also used in plumbing to freeze water in pipes to make temporary plugs of ice to enable repairs without draining the entire system.
- Local cooling of SAT in liquid phase using Peltier elements. A setup with a double-element Peltier device reached $-30\text{ }^{\circ}\text{C}$ and may be able to trigger crystallization [34].

Controlled seed crystal injection requires a rather complex device [12]. If the sealing membranes of the triggering units fail, there could be a risk of unwanted spontaneous solidification in supercooled storage material. In contrast, local cooling of SAT composites using CO₂ evaporation or Peltier elements can be done without any moving parts in the supercooled SAT and without penetrating the chamber containing the PCM and is therefore considered a potentially more durable and cost-efficient solution for closed PCM containers.

1.5. *Scope*

Local rapid cooling of a small part of a PCM storage unit has the potential to be used as a way to initiate the crystallization of supercooled SAT composites. It is therefore crucial to know the temperature at which relevant SAT composites crystallize during cooling. Apart from the additives in the SAT, the vessel containing the SAT composite may work as an impurity, affecting both the behaviour and the crystallization temperature of the SAT. So we investigated and determined the crystallization temperature of various promising SAT composites and of SAT in contact with steel. We also observed whether the length of time that the supercooled SAT composites were kept at 20 °C had any influence of the crystallization temperature. SAT composites with extra water, CMC, liquid polymeric solution, graphite flakes, and silicone oil were used for the experiments. The results of such investigations have not previously been reported in the literature.

Furthermore, this article includes the measurements and evaluations of two different methods of applying local rapid cooling of a part of the SAT composite volume in supercooled PCM prototype heat storage units: cooling by evaporation of pressurized liquid CO₂ and cooling by Peltier elements. Repeated experiments indicate that a reliable initialization of crystallization is possible. Measurements showed when temperatures were achieved that were sufficiently low for application with the various SAT composites. Detailed practical experience of initiating crystallization of supercooled SAT composites using these methods has not been reported previously in the literature.

2. Experimental method

2.1. Material cooling tests (freezer)

2.1.1. Materials

We used SAT of food grade (European standard 262i) from industrial supply [35]. The material was delivered in bags of 20 kg, 4 months after its production.

To stabilize the SAT, we used CMC and a prototype of a liquid polymer composition provided by Suzhou Hongde Co., Ltd. Jiangsu, China. Demineralised water was used to increase the water content of composites.

To study their influence on the minimum supercooling (nucleation) temperature, we also added graphite flakes and silicone oils to the samples. Table 1 presents the specifications of the materials used.

Table 1. Material properties.

	SAT	CMC	Graphite flakes	Liquid polymer	Silicone Oil
Type:	CAS 6131-90-4	Cekol 30000	- 10 mesh	HD 310	CAS 63148-62-9
Supplier:	IG Chemicals GmbH (GER)	CP Kelco (FIN)	Alfa Aesar	Sushou Hongde Co. Ltd. (CN)	Sigma Aldrich
Batch:	18.10.2015 (production date)	09.09.2013 (production date)	no. 43319	2016 (prototype)	-
Selected properties:	purity: >98.5%, water content: 36-42%wt., pH-value: 8-9.5	-	purity: 99% (metal basis)	Solid concentratoion: 40%, PH-value: 4.0, brookfield viscosity (at 20°C): 1.2	density: 963 kg/l at 25°C

After delivery, the exact water content of the SAT material was measured in dry-out tests. Four open glass cups were prepared with a thin layer (~2 mm) of crystalline SAT, resulting in four SAT samples with a weight of 20 g each. The SAT samples were put in a drying oven (Mettler UF55) with a temperature of 120 °C and air exchange to the ambient. Periodic weight measurements were conducted with a precision scale (linearity: +/- 30 mg) outside the drying unit. The dry-out tests were repeated with stored SAT 22 months after delivery.

Finely granulated SAT was melted in a closed glass jars at 90 °C for about 24 hours. To dissolve SA fully, about 1%wt. demineralized water was added to the granulated SAT. As explained in section 3.1, the SAT delivered showed differences from the molecular SA-water ratio (Fig. 8). The fully dissolved SA aqueous solution (40.5%wt. water) was referred to SAT.

Then the additives were added to the opened jars and the jars were weighed using a precision scale for individual material configurations. CMC was added in powder form during stirring, while extra water and liquid polymer were added using pipettes and the materials were thoroughly stirred afterwards. The mass of each additive was set to achieve a certain fraction of the gross-weight of the compositions (Eq.1):

$$\%wt. = \frac{m_{Additive}}{m_{SAT} + m_{Additive}} \quad (1)$$

The resulting compositions were re-heated in closed glass jars for about 12 hours. Then the samples were put into 50 ml glass jars with polymeric lids. Fig. 1 shows some of the SAT composite samples in solid state from the present work.

Fig. 1. SAT composite samples after crystallization (from left: SAT, SAT + water, SAT + CMC, SAT + HD 310).

The resulting sample weight depended on the fraction of additive in the mixed composition, based on 60 g of SAT. Graphite flakes and silicone oil were also added to similarly prepared samples. This approach was chosen to ensure an equal amount of SA in the samples for comparison.

To study the influence of rough metal surfaces, slices of corroded T-profiles of steel (welded connections) were cut with a metal saw and added to samples (Fig. 2b) in 2 batches for separate cooling tests.

An overview of all the samples prepared is presented in Table 2.

Table 2. The material samples investigated, distributed over five batches.

	Batch 1			Batch 2				Batch 3			Batch 4				Batch 5		
	SAT (g)	Add. 1 %wt.	Add. 2 %wt.	SAT (g)	Add. 1 %wt.	Add. 2 %wt.	Steel (g)	SAT (g)	Add. 1 %wt.	Add. 2 %wt.	SAT (g)	Add. 1 %wt.	Add. 2 %wt.	Steel (g)	SAT (g)	Add. 1 %wt.	Add. 2 %wt.
SAT	60	-	-	60	-	-	11	60	-	-	61	-	-	9	60	-	-
SAT + water	63	9	-	60	9	-	12	61	9	-	60	9	-	10	60	10	-
SAT + CMC	61	5	-	59	5	-	13	60	5	-	60	5	-	9	-	-	-
SAT + CMC + graphite flakes	60	5	5	60	5	5	13	60	5	5	60	5	5	10	60	5	5
SAT + HD 310	60	3	-	61	3	-	16	60	3	-	60	3	-	12	60	3	-
SAT + HD 310 + silicone oil	-	-	-	-	-	-	-	-	-	-	-	-	-	-	60	3	4

2.1.2. Test procedure

After sample preparation, TT-type thermocouples (copper/constantan) were attached to the outside of the glass jars. They were wrapped 3 times around the glass body and fixed with aluminium tape (Fig. 2a) to ensure good heat transfer. Next, the closed glass jars were evenly insulated with foam insulation (thickness: 15 mm) to avoid fast cooling in a cold environment and to achieve more uniform temperatures in the material samples.

The thermocouples positioned on the outer surfaces of the glass jars should have been able to measure the material temperatures because the thin glass wall (2 mm) had about 20 times greater thermal conductivity ($0.9\text{--}1.5 \text{ W m}^{-1}\text{K}^{-1}$) than the rather thick foam insulation ($0.04 \text{ W m}^{-1}\text{K}^{-1}$). Furthermore, the slow cooling rates ensured a uniform jar temperature, a precise measurement of sample temperatures, and a reduced risk of the glass jars breaking during the expansion and contraction of the samples. The bottom and the mantle insulation (as indicated in Fig. 2a) were glued together, while the cap insulation of the sample containers was fixed with tape.

The thermocouple wires of each batch (5 samples at a time) were connected to a portable 10-channel data logger (HIOKI LR8431-20) to ensure continuous logging of temperatures. The ambient air temperature was logged with an additional thermocouple (the 6th channel on the data logger). Measurements were logged at intervals of one second.

Fig. 2. (a) Sample containment for cooling tests; (b) Glass jar with SAT and sliced steel profile.

To ensure uniform material temperatures, samples were heated in the oven without insulation (placed side-by-side; Fig. 3a) for about 12 hours.

During a cooling down process, the samples were placed in foam insulation and rested, first at room temperature (about 20 °C). Then they were put into a freezer (temperature of about –30 °C) with 10 cm distance to the bottom and the walls, realised using a metal framework (Fig. 3b). The test conditions applied for each batch of samples are given in Table 3.

Fig. 3. Photos: (a) Sample placement during heating; (b) Sample placement during cooling.

The cooling process was stopped after 12 hours, when the sample temperatures were no more than 1 K above the air temperature in the freezer. Next, the samples were kept in their insulation for a period of 4 hours at room temperature before the insulation was finally removed.

Table 3. Test conditions.

	Batch 1	Batch 2	Batch 3	Batch 4	Batch 5
Charging temperature (°C)	90	90	90	90	85
Rest period	none	none	7 days	8 days	2 h
Temperature at freezer placement (°C)	>60	>60	~20	~20	<30
Steel profile?	no	yes	no	yes	no
No. of repetitions	1	1	1	1	4

With batches 1–4, we tested various SAT compositions with different rest periods, cooling temperatures, and with and without the presence of a rusty steel profile in the sample. We found that nearly all the glass jars containing a CMC composition broke due to thermal and mechanical stress. So we decided to produce five more samples, to reduce the heating temperature slightly, and to apply a minimum rest period of 2 hours at room temperature (until the samples cooled down to 30 °C) the samples were placed in the freezer. With these samples (Batch no. 5), it was possible to repeat the heating and cooling tests.

2.1.3. Analysis

Minimum supercooling temperatures, i.e. crystallization temperatures, were detected when sample temperatures started to rise due to the release of latent heat. We therefore measured the local temperature minima (T_{CRi}) reached in each sample during the cooling process. The first local minimum (T_{CR1}) was defined as the crystallization temperature. We also measured the increase in the temperature of each sample (ΔT_{CRi}) after each local minimum, so that we could compare the different samples.

The average logged temperature in the freezer during cooling of the supercooled samples was set as the reference (T_{ref}) for the cooling rate calculation. The start of the cooling periods (T_{C1}) was set to 45 K above the reference temperature, while the end of the cooling period (T_{C2}) was set to 20 K above the reference temperature. The resulting temperature difference (25 K) was divided by the duration of cooling in minutes (Eq. 2) to obtain the cooling rate:

$$cool-down\ rate\ (K\ min^{-1}) = \frac{(T_{C1} - T_{ref}) - (T_{C2} - T_{ref})}{t(T_{C1}) - t(T_{C2})} \quad (2)$$

For example, Fig. 4 shows the approach we chose applied to a cooling curve with SAT + 9%wt. water. The characteristic temperature levels are indicated.

Fig. 4. An example illustrating the evaluation method in the cooling tests (schematic).

2.2. Initialization of crystallization using CO_2 evaporation

Prototype heat storage units with SAT composites were heated up to 90 °C and actively or passively cooled down to ambient temperatures under controlled conditions, as described by Dannemand et al. [10]. The prototype heat storage units were 240 x 120 x 5 cm and contained 200–220 kg of SAT composite. In a series of six experiments (overview in Table 5 and section 3.6), the SAT composites in the units were stable in supercooled state at ambient room temperatures

(about 20 °C), before we initiated crystallization by evaporating CO₂ in a small chamber adjacent to the chamber with the supercooled PCM.

As shown in Fig. 5, a CO₂ cylinder with a dip tube was connected via a rubber tube to the 100 mL CO₂ chamber on one side of the heat storage unit. Another tube was connected to the outlet of the CO₂ chamber with a pressure-regulating valve including a manometer. The pressure-regulating valve was set to maintain a pressure of approximately 5.5 bars in the tubes and the CO₂ chamber. This was to allow the CO₂ to remain in liquid phase in the tubes and the CO₂ chamber.

The temperatures on the outside of the CO₂ chamber (T_{CO_2}), on the side of the PCM chamber ($T_{surface}$), and inside the PCM (T_{PCM}) were measured using thermocouples (TT-type). The temperature in the PCM was measured in the other side of the PCM chamber, approximately 240 cm away from the CO₂ chamber.

To start the crystallization of the supercooled SAT composite in the heat storage unit, the valve for the CO₂ cylinder was opened and the CO₂ was flushed through the CO₂ chamber until a temperature increase (indicating crystallization) was detected by the nearby temperature sensor ($T_{surface}$) on the side of the PCM chamber.

Fig. 5. Principle diagram of the nucleation method using cooling with CO₂.

2.3. Initialization of crystallization using Peltier elements

Based on previous work by Dröscher [36], we built a novel double-element Peltier device with cooling power of 8.8 W and 29.7 W (Fig. 6).

To ensure contact with the PCM, a steel extension pipe with a hole drilled in the bottom was mounted on the Peltier device. The extension pipe was filled with PCM. Rubber sealing ensured air-tightness in the construction. The lower Peltier element was in contact with the pipe via a steel plate. Thermal compound was used to improve the heat transfer to the steel plate. A thermocouple (type K) was placed below the lower Peltier element to measure the temperature of its cold side, which was attached to the steel plate ($T_{steel\ plate}$). Above the upper Peltier element, a heat sink was realised with an air cooler mounted, consisting of an electrically driven fan and aluminium fins. The temperature of the heat sink ($T_{heat\ sink}$) was measured with a thermocouple.

**Fig. 6. (a) Drawing of the Peltier device with extension pipe (dimensions in mm);
(b) Peltier triggering device with extension pipe (without the heat sink).**

The Peltier device was tested with a small-scale flat heat storage prototype, filled with 5.4 kg SAT + 6%wt. extra water. It was attached via a steel plate to the top-left flange of the heat storage unit (Fig. 7). An additional thermocouple was inserted into the opposite side of the heat storage unit to measure the temperature of the SA-water mixture (T_{PCM}).

The SA-water mixture was heated to 85 °C and then passively cooled down to room temperature. The power supplies of the two Peltier elements and the fan were turned on until an increase in the temperature T_{PCM} indicated the crystallization of the SA-water mixture.

Fig. 7. Photograph of the Peltier device mounted on the small-scale heat storage prototype.

3. Results and discussion

3.1. Water content of SAT from industrial supply

Fig. 8 shows the water loss over time during material dry-out. The various water-loss development characteristics of the samples were caused by differences in local temperature and humidity inside the drying oven:

Fig. 8. Drying of SAT samples.

In the case of samples 1–4, the drying tests were conducted after delivery (4 months after production), and they showed an average water loss of 39.5%wt. Based on the molar SA-water ratio, the water content of chemically pure SAT is calculated to be 39.72%wt. Taking the minimal purity of the material into account, the water loss was expected to be 39.12%wt. (lower benchmark). The influence of scale accuracy on the final weight measurement was calculated to be $\pm 0.248\%$ wt. The mass loss measured therefore indicates the complete dry-out of SAT containing impurities.

After a period of 22 months (26 months after production), samples 5–8 were taken from a SAT bag, which was stored in the laboratory. By this time, the material, which was originally finely granulated particles, had clumped together due to its hygroscopic properties. An average water loss of 40.80%wt. was measured. This indicates that the water content of SAT from industrial supply strongly depends on the storage conditions.

In this research, SAT was defined with a water content of 40.5%wt. This value was found to be necessary to fully dissolve SA in water during melting when SAT of food grade (purity $>98.5\%$) was used. The value also includes water vapour losses during the melting of the SA-water mixtures in closed glass jars.

3.2. Crystallization behaviour of SA-water mixtures

In the cooling tests of SAT and SAT with extra water, two stages of heat release were observed. Fig. 9 shows the temperature developments in the SA-water mixtures in four tests with similar conditions. After the initialization of crystallization was detected for the first time (T_{CR1}), the samples cooled down to a temperature below T_{CR1} , until a second temperature increase was measured at T_{CR2} .

Fig. 9. Crystallization behaviour (sample batch no. 5) in SAT (top) and SAT with extra water (bottom).

As listed in Table 4, with extra water the SA-water mixture cooled down to significantly lower temperatures than any of the other SAT composites (the mean value of T_{CR1} was about 2 K below the T_{CR1} in other samples). So, extra-water did lower the crystallization temperature. Moreover, the second release of latent heat happened at lower temperatures with this SAT composite.

When the second release of latent heat occurred at a temperature around -25°C , a rapid temperature increase was observed. In some cases, the second reaction was initialised at higher temperatures and ΔT_{CR2} was lower, followed by a longer cooling period. This behaviour indicates that the heat content released during the reaction was similar to when there was a rapid temperature increase.

3.3. Crystallization behaviour of SAT with stabilizers

The composites with CMC or liquid polymer did not have a second temperature peak during the cooling process. Fig. 10 shows, however, that samples containing stabilizers reached significantly higher temperatures after crystallization started (evaluated in Fig. 13). Afterwards, the SAT composites cooled down over a similar time period (about 8 hours) to the freezer temperature.

Fig. 10. Crystallization behaviour (sample batch no.5) in SAT +CMC + graphite flakes (top) and SAT +HD 310 (bottom).

The presence of silicone oil in samples containing liquid polymer did not influence temperature developments. As shown in Table 4, four similar heating and cooling cycles revealed no significant tendency for T_{CR1} and T_{CR2} to vary and neither did the type of stabilizer (CMC or liquid polymer) significantly influence T_{CR1} (basis: 12 tests).

3.4. Influence of steel profiles on the crystallization behaviour

In parallel tests with various samples containing steel profiles (Fig. 11 bottom), almost uniform crystallization temperatures were detected. As shown in Table 4, crystallization started at higher temperatures. In contrast, samples without steel profiles (Fig. 11 top) showed a big variation in crystallization temperatures, indicated by the time gaps between the crystallizations.

Fig. 11. Crystallization behaviour of sample batch no. 1 (without steel profiles; top) and sample batch no.2 (with steel profiles; bottom).

We found no significant difference in the crystallization behaviour of SAT composites cooled immediately (Fig. 11) and those first given one week of stable supercooling at 20 °C before cooling (Fig. 12). However, as shown in Table 4, lower crystallization temperatures were observed in samples containing rusty steel profiles that were given the rest period before cooling.

Fig. 12. Crystallization behaviour after one week rest sample batch no. 3 (without steel profiles; top) and sample batch no.4 (with steel profiles; bottom).

3.5. Cooling test analysis

Cooling rates varied only slightly in between 0.2 and 0.24 K min^{-1} . Samples containing extra water cooled down most slowly, while the SAT only samples showed the fastest cooling. Table 4 gives an overview of the local minimum temperatures in all the tests. Temperature values showing a moderate temperature increase are highlighted in grey.

The random measurement error of the thermocouples used was $\pm 0.5 \text{ K}$, assuming a uniform probability distribution. Because measurements were taken every second and the cooling rates were rather low, the temperature measurements are considered to be precise. The position of the sensors means the temperatures measured should be very close to the actual sample temperatures. Moreover, the slow cooling resulted in a graph with a very small slope, which enables us to say that crystallization usually started at about the temperature reported in the literature for pure SA aqueous solutions ($-15 \text{ }^{\circ}\text{C}$). However, significantly lower temperatures were measured with Batches 1 and 3.

During cooling of the samples without steel profiles present, we observed composite-specific ranges for T_{CR1} (and T_{CR2} where applicable). The crystallization of SAT and SAT + water samples (16 tests) started in the range of -11.5 to $-23.0 \text{ }^{\circ}\text{C}$, while in the samples with stabilizers (24 tests), T_{CR1} occurred in the range of -10.7 to $-17.2 \text{ }^{\circ}\text{C}$. The second release of heat for SAT and SAT + water (16 tests) ranged from -17.8 to $-30.4 \text{ }^{\circ}\text{C}$.

Table 4. Overview of crystallizations temperatures (T_{CR1}) and second local minimum temperatures (T_{CR2}).

	Batch 1	Batch 2	Batch 3	Batch 4	Batch 5			
	no rest	no rest steel	rest	rest steel	test 1	test 2	test 3	test 4
	(°C)	(°C)	(°C)	(°C)	(°C)	(°C)	(°C)	(°C)
T_{CR1}								
SAT	-21.8	-10	-23.8	-14.4	-14.2	-11.5	-15.8	-16.2
SAT + water	-22.9	-8.8	-12.6	-14.4	-16.9	-16.4	-18.5	-17.6
SAT + CMC	-15.3	-10.2	-13.7	-12.6	-	-	-	-
SAT + CMC + graphite flakes	-17.2	-10.9	-17.6	-14.5	-13.1	-15.9	-16.8	-16.9
SAT + HD310	-12.1	-10.3	-10.5	-11.7	-15	-15.8	-14.9	-16.6
SAT + HD 310 + silicone oil	n.a.	n.a.	n.a.	n.a.	-14.2	-16.6	-13.1	-10.7
T_{CR2}								
SAT	-28.9	-20.6	-30.4	-20.5	-27	-24.1	-21.1	-26.3
SAT + water	-23.0	-17.8	-26.7	-20.5	-27.9	-19.7	-25.9	-26.2

The samples with steel profiles (10 tests) showed a clear tendency to more uniform and higher T_{CR1} and T_{CR2} values. Crystallization temperatures between -8.8 and -10.9 °C were measured for samples cooled without a rest period (5 tests). The second local minimum resulted in a moderate temperature increase.

The samples with steel profiles that had a rest period before cooling (5 tests) crystallized at temperatures in a lower range (-11.7 to -14.5 °C). There were no significant differences in the local temperature minima in samples without steel profiles that had a rest period before cooling (5 tests) and those that did not (5 tests).

Several of the glass jars containing CMC broke. The tests were repeated with Batch 5, when the test procedure had been improved.

In Fig. 13, the blue bars show the average temperature increase in samples during the cooling tests. Here we have excluded the two tests with samples containing CMC and graphite flakes (Fig. 11 top and Fig. 12 top) that did not show a sharp temperature increase when crystallization started.

Because the material volumes of the composite samples (based on 60 g SAT) differed, ΔT_{CR1} can only be compared in a qualitative way. Nevertheless, a clear order of average temperature increase can be seen: SAT samples showed 36.6 K; SAT + water samples showed 40.3 K; SAT + CMC showed 46.3 K, and 46.5 K with added graphite; SAT + HD 310 showed 50.9 K, and 49 K with silicone oil added. This order is in good agreement with the heat content for SAT composites measured by Kong et al. [14] using a T-history method. This indicates that CMC and liquid polymer significantly improve the reactivity of SA with water during crystallization.

Fig. 13. Comparison of temperature increase during crystallization of material samples.

The second temperature increase (ΔT_{CR2} , red bar segments) was derived from samples that showed a sharp temperature increase (10 tests). In 6 out of 16 tests runs, the second temperature peak was reached slowly. The total temperature increase ($\Delta T_{CR1} + \Delta T_{CR2}$) was 41.6 K for SAT and 48.3 K for SAT + water samples. This indicates that, due to the very low temperature in these samples, further exothermic crystallization was enabled. This further potential of heat release was not observed by Kong et al. [14] when crystallization took place at 20 °C. We conclude that it can be only utilized when the SA-water mixtures (full volume) are exposed to low temperatures, as in the freezer (−30 °C).

The presence of steel profiles (10 tests) and periods of rest before cooling showed no systematic influence on the temperature increases.

3.6. Initialization of crystallization using CO_2 evaporation

Six experiments (Table 5) with the initiation of nucleation using cooling by evaporating CO_2 were carried out with the prototype units. A temperature range of −27 to −44 °C was measured on the outside on the CO_2 chamber. After 1.5–5.0 minutes of releasing CO_2 into the chamber, a temperature increase was detected at the nearby temperature sensor ($T_{surface}$) on the surface of the PCM chamber and the valve on CO_2 cylinder was closed. The crystallization probably started earlier, because it would take some time for the temperature to increase at the point next to the CO_2 chamber where the sensor was. Furbo et al. have reported that supercooled SAT nucleated within 30 seconds of applying the CO_2 evaporation method through a 2 mm metal plate. In their case, the nucleation was observed visually [37].

Table 5. Overview of our experiments with heat storage unit prototypes to initialize crystallization using CO₂ evaporation.

Experiment	1	2	3	4	5	6
PCM	SAT + 1%wt. CMC	SAT + 5%wt. water	SAT + 5%wt. water	SAT + 1%wt. CMC	SAT + 1 %wt. CMC	SAT +1%wt. CMC + 2%wt. graphite powder
Cooling method	active	active	active	passive	passive	active
Duration of stable supercooling	1 day	5 days	3 days	14 days	71 days	1 day
Minimum temperature CO ₂ chamber	-30 °C	-34 °C	-36 °C	-27 °C	-27 °C	-44 °C
Maximum PCM temperature during solidification	58 °C	53 °C	53 °C	58 °C	58 °C	58 °C

Fig. 14 shows the development of T_{PCM} , $T_{surface}$ and T_{CO_2} in the heat storage unit during the initialization of crystallization in the supercooled SAT composite in Experiment 1 (1%wt. CMC). As the PCM composite temperature was measured at the other end of the heat storage unit, approximately 240 cm from the CO₂ chamber, the crystallization and temperature increase occurred there approximately 12 minutes after the nucleation started next to the CO₂ chamber. The maximum temperature reached in the PCM after crystallization is also listed in Table 5.

For each nucleation, 700–900 g CO₂ was used. This could probably be significantly reduced with a faster response in relation to when nucleation actually started and an optimized design of the tube and CO₂ chamber size.

Fig. 14. Temperature development in the PCM, the CO₂ chamber and next to the CO₂ chamber.

3.7. Initialization of crystallization using Peltier elements

Two series of ten experiments each were carried out with the Peltier triggering device. After the SA-water mixture cooled to room temperature, the Peltier device was turned on. The temperature reached in the area of the lower Peltier element ($T_{steel\ plate}$) varied from -31 to -36 °C. Despite the limited cooling power of the Peltier elements, sufficiently low cold-side temperatures were possible when the SA-water mixture supercooled to temperatures below 30 °C.

In the first test series the number of necessary cooling attempts increased till the last experiment, when no crystallization was initiated. The time gap between turning on the Peltier device (first attempt) to measured temperature increase of the PCM varied from 5 minutes to more than six hours at the last experiment. The reason for unsuccessful attempts might be due to the SA-water mixture flowing out of the extension pipe so that no PCM was in contact with

the steel plate. So we reduced the diameter of the bore-hole at the front of the extension pipe from 1mm to 0.4 mm and the PCM filling volume of the extension pipe from 7.5 to 4.5 ml by adding inert material. The second test series showed 10 successful experiments, in each of which the SA-water mixture crystallized 2.5–7.5 minutes after the Peltier device was turned on.

Fig. 15 shows the temperature developments in Experiment 3 of the second test series. $T_{\text{steel plate}}$ decreased as $T_{\text{heat sink}}$ increased. When $T_{\text{steel plate}}$ reached $-33\text{ }^{\circ}\text{C}$, a temperature increase was observed (dotted line in Fig. 15) indicating that the SA-water mixture had started to solidify in the extension pipe. One minute later, the solidification was detected at the temperature sensor in the PCM container (T_{PCM}). During crystallization, the SA-water mixture reached a temperature of $56.5\text{ }^{\circ}\text{C}$ (T_{PCM}).

Fig. 15. Crystallization of SA-water mixture by local cooling using Peltier elements.

3.8. Economic aspects

For industrial use SAT is available in large quantities. Market prices depend on its purity and are typically below 0.5 Euros per kilogram. Therefore SAT is potentially an economically attractive heat storage material. For economic assessment of heat storage units additives, container, heat exchanger, hydraulics and devices for controlled activation of crystallization need to be taken into account. Rathgeber et al. [38] made a top-down evaluation of acceptable costs of energy storages. They found that acceptable storage capacity costs ranged from about 1 Euro per kWh for seasonal heat storage to 429 Euros per kWh for daily utilization in buildings. Englmair et al. [12] demonstrated that flat prototype heat storage units, utilizing stable supercooling of 200 kg SAT composites, enable combined short- and long-term heat storage in domestic solar heating systems. Their total heat storage capacity was 27.4 kWh. 12 storage cycles per year result by assuming monthly utilization. In this case acceptable costs for such heat storage units range from 200 to 460 Euros. Prototype heat storage units and activation devices used in the presented work need to be further developed to achieve this cost range. Cylindrical unit design [11] is expected to allow reduction of container and heat exchanger costs in comparison to flat units. Peltier elements and CO_2 evaporation can potentially initiate SAT composite solidification with inexpensive devices of reduced size.

4. Conclusions

All the SAT composites investigated supercooled to temperatures below $-8\text{ }^{\circ}\text{C}$, which means they can be considered useful for application as heat storage materials in households. The following temperatures were required for initialization of crystallization of SAT composites:

- $-24\text{ }^{\circ}\text{C}$ for SAT and SAT with 9%wt. water added
- $-18\text{ }^{\circ}\text{C}$ for SAT with 5% CMC and 3% of HD 310 added
- $-15\text{ }^{\circ}\text{C}$ for all SAT composites in closed steel containers

Extra water added to the SAT resulted in lower crystallization temperatures in comparison to SAT. Experiments on additional impurities in SAT compositions in the form of rusty steel profiles (which could resemble the metal surfaces of the PCM container) showed that crystallization started at higher and more uniform temperatures for all samples. Letting the samples rest at ambient temperature only affected the crystallization behaviour when steel profiles were present.

The cooling tests proved useful for investigating the supercooling stability of SAT composites:

- A second minor temperature peak occurred below $-20\text{ }^{\circ}\text{C}$ in the SAT composites with no stabilizers added, and the samples cooled further after the first crystallization. When it occurred below $-21.3\text{ }^{\circ}\text{C}$, a sharp temperature rise resulted. Otherwise, the temperature increase was slow.
- SAT with 9%wt. extra water showed higher temperature increases during crystallization than SAT on its own.
- SAT composites with stabilizers (carboxymethyl cellulose and liquid polymer HD 310) showed the highest temperature increase when crystallization was initiated, and there was no second temperature peak as a result of the release of heat.

When a large amount of loose crystalline SAT in paper bags is used to prepare composites for heat storage, its water content must be determined. During melting, 40.5%wt. water content was necessary to fully dissolve SAT of food grade.

To initiate crystallization of SAT composites in heat storage units in a controlled way, evaporating carbon dioxide in a chamber adjacent to a PCM chamber with supercooled SAT composites worked reliably. A device with Peltier elements mounted on a PCM chamber was also able to generate local cooling to sufficiently low temperatures to initiate crystallization for all the SAT composites investigated when the SA-water mixtures had supercooled to temperatures below $30\text{ }^{\circ}\text{C}$. With both methods, local cooling was realised rapidly, so that crystallization was initiated two to five minutes after the experiments were started.

Declaration of interest

None.

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