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## Freezing-induced topological transition of double-emulsion†

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Solidification of complex liquids is pertinent to numerous natural and industrial processes. Here, we examine the freezing of a W/O/W double-emulsion, *i.e.*, water-in-oil compound droplets dispersed in water. We show that the solidification of such hierarchical emulsions can trigger a topological transition; for example, in our case, we observe the transition from the stable W/O/W state to a (frozen) O/W single-emulsion configuration. Strikingly, this transition is characterised by sudden expulsion of the inner water drop from the encapsulating oil droplet. We propose that this topological transition is triggered by the freezing of the encapsulating oil droplet from the outside in, putting tension on the inner water drop thus, destabilizing the W/O/W configuration. Using high-speed imaging we characterize the destabilization process. Interestingly, we find that below a critical size of the inner drop,  $R_{in,crit} \approx 19 \mu\text{m}$ , the topological transition does not occur any more and the double-emulsion remains stable, in line with our interpretation.

## 1 Introduction

The characteristic physical feature common between the fabrication of functionalized ceramics,<sup>1–4</sup> cryo-biology,<sup>5–7</sup> frost-heaving in colder regions<sup>8–10</sup> and ice cream making is the interaction between insoluble dispersed particles in a melt and a moving solidification front. Due to its relevance and impact in both natural phenomena and technological applications, this interaction has been extensively studied both experimentally and theoretically.<sup>1,2,5–8,11–13</sup> The prominent focus of these investigations has been to determine the conditions under which dispersed particles are either engulfed or rejected by the advancing freezing front.<sup>11,12,14–16</sup> This aspect holds particular importance in material science, as the distribution of dispersed particles profoundly influences the resulting microstructure, thereby governing the functional properties of the solidified materials. Importantly, these collective efforts have revealed that the interaction between a particle and the moving solidification front is influenced by distinct physical processes operating across different length scales. This characteristic renders it a true multi-scale problem, with each decade of length scale presenting its own set of governing mechanisms and phenomena. For instance, at the particle level, the interaction is influenced by

factors such as thermal conductivity mismatches between the particle and the melt, as well as density changes during the liquid–solid phase transition. At smaller scales, van der Waals interactions and the flow of lubricating films in premelted layers dictate the local dynamics.<sup>17</sup> Furthermore, recent investigations involving “soft” particles, such as droplets or bubbles, have shown that the overall complexity is further amplified due to the mechanical deformation experienced by the particles during the interaction with the moving solidification front.<sup>18–20</sup>

The current understanding of this subject primarily stems from studying the interaction between uniform particles and a moving solidification interface. However, in modern technological applications, such as cryobiology and material science, complex and multi-component particles are frequently encountered. Inspired by these applications, in this Letter, we extend the envelope of the current state of the art on this subject to the interaction between the solidification front and multi-component particles. Using a dilute W/O/W double-emulsion as a model system, we report unexpected behaviors that emerge during its solidification. We show that dispersed compound droplets in a double-emulsion can undergo a topological change upon engulfment in the solidifying bulk. This topological transition is triggered by the partial solidification of the dispersed compound droplet. Our findings are relevant for exerting better control over cryopreservation procedures of bio-specimen as well as the fabrication of advanced materials.

## 2 Experimental procedure

The W/O/W double-emulsions used in our current experiments are prepared using a microfluidic setup similar to the one

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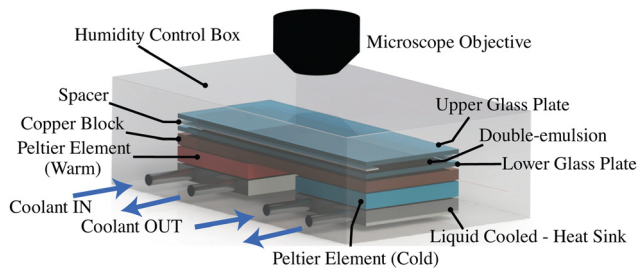


Fig. 1 Schematic of the experimental setup.

described in ref. 23. In our adopted setup, we exploit the process of spontaneous liquid–liquid phase separation of a ternary mixture (water, ethanol and oil) to achieve double-emulsions, as discussed in detail in ref. 24 and 25. Note that here we use diethyl-phthalate (DEP) as an oil phase and Pluronic-F127 as surfactant to stabilise the emulsion. The oil has a freezing point close to that of water, *i.e.*,  $T_{m,DEP} = -3$  °C. Since the process of emulsion preparation is extremely sensitive to the initial composition of the ternary mixture, we use a DEP/ethanol/water mixture (0.74/0.05/0.21 vol.) with which we achieve a stable double-emulsion. As a consequence of the adopted methodology of the double-emulsion preparation, both the inner water drop and the encapsulating oil droplet contain traces of ethanol.

The experimental assembly employed in current investigations is shown schematically in Fig. 1. In a typical experiment, the W/O/W double-emulsion is first injected into a horizontal Hele-Shaw cell of thickness 200  $\mu\text{m}$ , resting on a copper block. A thermal gradient  $G \sim 5 \text{ K mm}^{-1}$  is then applied to this copper block using Peltier elements (spaced 3 mm apart), such that the continuous phase of the double-emulsion (water) begins to solidify uniformly from the cold end. Solidification is initiated by placing a small piece of ice at the cold end of the cell when its temperature falls below the melting point of water. It is important to note that, due to the extremely slow freezing rates employed in our experiments, clear ice is formed with a refractive index (1.31) similar to that of water (1.33). The solidification rate  $V$  of the bulk medium is controlled by the applied thermal gradient  $G$ . The engulfment of the dispersed compound droplets into the solidified bulk is recorded in top-view using a camera (Nikon D850 or Photron NOVA S16) connected to a long working distance lens (Thorlabs, MVL12X12Z plus 2X lens attachment). The sample is illuminated with diffused cold-LED to avoid localised heating. For further experimental details and the protocol for the double-emulsion preparation, we refer to the ESI.†

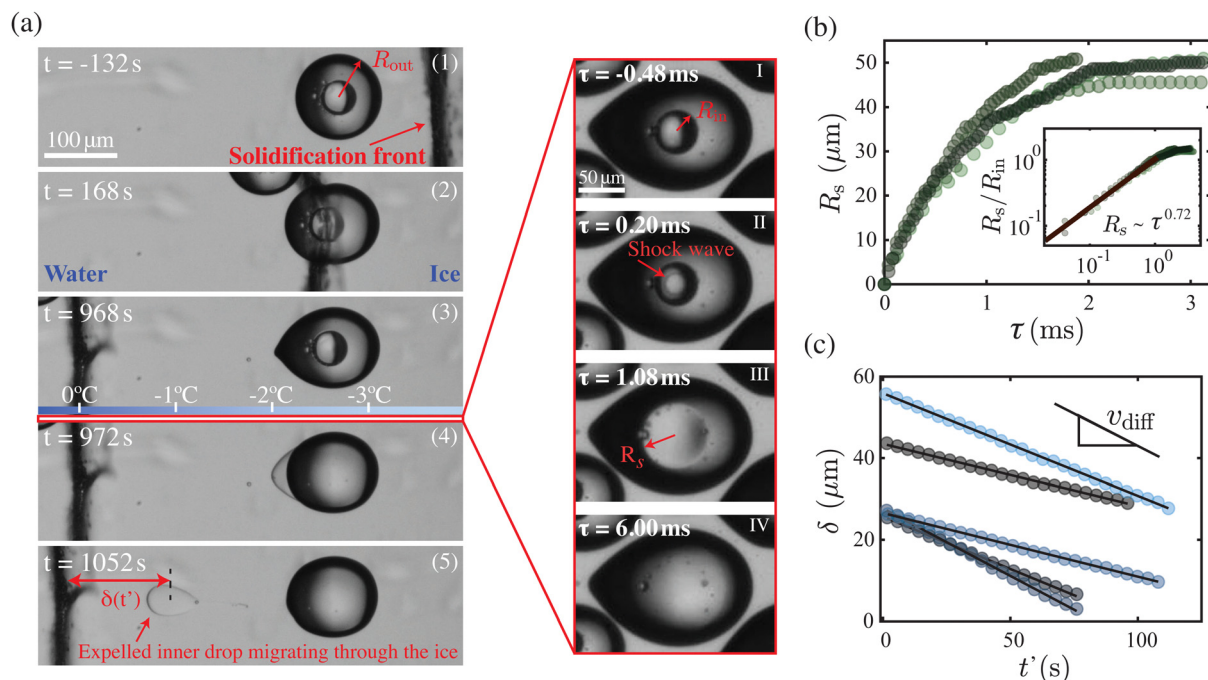
### 3 Results and discussion

The sequence of images in Fig. 2(a) highlights the events leading up to the sudden topological transition of a W/O/W double-emulsion when a dispersed water-in-oil compound droplet is engulfed into the solidifying bulk at a steady rate,  $V \approx 0.5 \mu\text{m s}^{-1}$ . The moment in time when the solidification front makes contact with the compound droplet is defined as  $t = 0$ . As shown in Fig. 2(a), while the solidification front passes over, the

encapsulating oil droplet severely deforms into a pointy, tear-like shape. Recent findings<sup>20</sup> have revealed that this deformation of the engulfed droplet is mediated by pressure variations in the nanometric thin liquid film, also referred to as premelted film,<sup>17</sup> sandwiched between the dispersed droplet and the solidifying bulk. In contrast, during this engulfment process, the inner water drop clearly remains unaffected. However, very unexpectedly, a few minutes ( $\sim 16$  min) after the dispersed compound droplet is completely engulfed in the solidified bulk (ice) and the solidification front is not in its vicinity, the inner water drop suddenly gets expelled outside the encapsulating oil droplet, altering the topology of the initially stable W/O/W double-emulsion. The expelled liquid gathers in the thin premelted film enveloping the oil droplet. Interestingly, the accumulation predominantly occurs at the warmer end, suggesting that the temperature gradient plays a role in this process. More importantly, as this topological transition occurs, the pointy, tear-like shape of the oil droplet relaxes to its original unperturbed (spherical) shape. Finally, we observe that the expelled liquid eventually begins to migrate towards the moving solidification front at a constant speed  $v_{\text{diff}} \sim 2 \mu\text{m s}^{-1}$ , see Fig. 2(c). This surprising migration of the liquid through the solidified bulk resembles a process akin to brine-pocket diffusion in sea ice that is driven by a delicate diffusive process of solute diffusion and its influence on the local phase equilibrium.<sup>26–28</sup> It is worth noting that, as a result of the chosen fabrication method for the double-emulsions used in this study, the dispersed compound droplets contain traces of ethanol, which eventually acts as a solute, thus enabling a diffusive process that leads to the migration of the liquid pocket. For a detailed discussion of the physical mechanisms leading to the migration of the liquid pocket, we refer to the ESI.† In the following, we focus on the main finding of this paper: the sudden topological transition of the double-emulsion.

Since the time scales associated with the topological change of engulfed compound droplets are much faster than the rate of solidification, we employ high-speed imaging to reveal the details of this apparently instantaneous event. As illustrated in the sequence of images in Fig. 2(a)(I)–(IV), the transition is triggered at the center of the inner drop, marked by the appearance of a radially expanding dark ring. Remarkably, this outward-moving dark ring is also accompanied by a localized increase in light intensity at its center, forming a discernible bright spot. Once the size of the bright spot exceeds the dimensions of the inner drop, it gradually fades (see ESI†), coinciding with the complete disappearance of the inner water drop. Following this event, the disappeared liquid collects outside the enveloping oil droplet within the premelted film in about 10–100 ms (not shown here). By tracking the temporal expansion of this dark ring,  $R_s(t)$ , we obtain a power-law-like behaviour  $R_s \sim \tau^{0.72}$  at the early stages of its growth when  $R_s/R_{\text{in}} < 1$ , see inset Fig. 2(b). Such scaling behavior evokes similarities with an adiabatically expanding spherical shock wave, which entails a discontinuity in density, pressure, and temperature.<sup>21,22</sup> Considering that the double-emulsions used in our current experiments contain traces of ethanol, we propose that all the above described features are triggered by





**Fig. 2** (a) (1)–(5) Sequence of images capturing the engulfment of a water-in-oil compound droplet by a solidification front, advancing at rate  $V = 0.5 \mu\text{m s}^{-1}$  (Movie 1, ESI†). The moment in time the front makes first contact with the droplet is set to  $t = 0$ . The third panel shows a sketch of the expected temperature distribution in the ice at a certain moment in time set by the strength of the applied thermal gradient  $G$ . (I)–(IV) High-speed images were acquired at 25 000 fps, showing the sudden topological transition (Movies 2 and 3, ESI†). At  $\tau = 0$  a shock wave appears at the center of the inner drop and an ethanol vapour cloud expands radially outward. (b) Radius  $R_s$  of the shock wave as a function of time for different transitions, indicated by different shades. The inset shows that at the early stages, when  $R_s/R_{\text{in}} < 1$ , it follows a power-law-like behaviour that is consistent with that of a diverging spherical shock wave.<sup>21,22</sup> (c) Distance  $\delta$  between the solidification front and the center of the expelled inner drop migrating through the ice. Note the  $\sim 5$  orders of magnitude difference in time scale in (c) as compared to (b). At  $t' = 0$  the expelled drop detaches from the oil. The migration velocity is determined by the velocity  $v_{\text{diff}}$  at which diffusion takes place within the expelled water–ethanol pocket.

its phase transition from liquid to vapor during the solidification of the surrounding oil droplet. Since ethanol is crucial for the stability of the double-emulsion, its sudden phase transition leads to the dissolution of the inner water drop, destabilizing the compound droplet. The process of dissolution, we argue, is optically screened-off by the vapour cloud (bright spot) itself and therefore not visible in Fig. 2(a)(I)–(IV). Note that the ethanol vapor allows greater transmission of light because of its reduced refractive index, which in turn rationalizes the appearance of a bright spot during the transition. Also, we will see later that the rate of dissolution indeed exceeds the rate at which the vapour cloud expands.

The hypothesized phase transition of ethanol during the solidification of the double-emulsion is rationalized as follows: following engulfment, as the solidification front steadily recedes away from the dispersed compound droplet, the local temperature in its vicinity progressively decreases in line with the applied thermal gradient (see sketch in Fig. 2(a)(3)). Upon reaching the freezing point of the oil ( $T_{\text{m,DEP}} = -3^\circ\text{C}$ ), the outer oil droplet begins to solidify, causing shrinkage in its volume, since  $\rho_{\text{oil-solid}} > \rho_{\text{oil-melt}}$ . Nevertheless, this volumetric shrinkage is prevented due to the confinement by the solidified bulk water. As a result the pressure inside the encapsulating oil droplet plummets, imposing tension on the inner water drop and causing it to stretch. An approximate estimate of the

pressure reduction experienced by the inner drop due to the solidification of the encapsulating oil droplet can be determined from the Gibbs–Duhem relation<sup>29</sup>

$$\rho_{\text{solid}} \mathcal{L} \frac{T_{\text{m}} - T}{T_{\text{m}}} = \Delta p \left( 1 - \frac{\rho_{\text{solid}}}{\rho_{\text{melt}}} \right). \quad (1)$$

Here,  $\mathcal{L}$  is the latent heat of solidification,  $T$  is the temperature and  $\Delta p$  the corresponding pressure change which we want to evaluate. Estimating  $\Delta p$  from eqn (1) with typical values,  $T_{\text{m}} - T = 0.1$  K and  $1 - \rho_{\text{solid}}/\rho_{\text{liquid}} = -0.1$ , yields  $\Delta p \sim -1$  MPa. Such a depression in pressure critically affects the local phase equilibrium of ethanol for which the saturated vapour pressure at  $T = -3^\circ\text{C}$  is  $p_{\text{v,eth}} \approx 1.2$  kPa. To verify our hypothesis, we performed additional experiments by replacing DEP with dibutyl-phthalate (DBP) as an oil phase, while keeping the procedure of the double-emulsion preparation the same (see ESI†). Since the freezing point of the DBP is much lower compared to DEP,  $T_{\text{m,DBP}} = -35^\circ\text{C}$ , which cannot be achieved in our experimental setup, no abrupt topological transitions are observed. For further details related to these experiments, we refer to the ESI†.

Since the dissolution process of the inner water drop is screened-off by the ethanol vapor cloud (bright spot) appearing at the center, it is extremely difficult to directly measure related





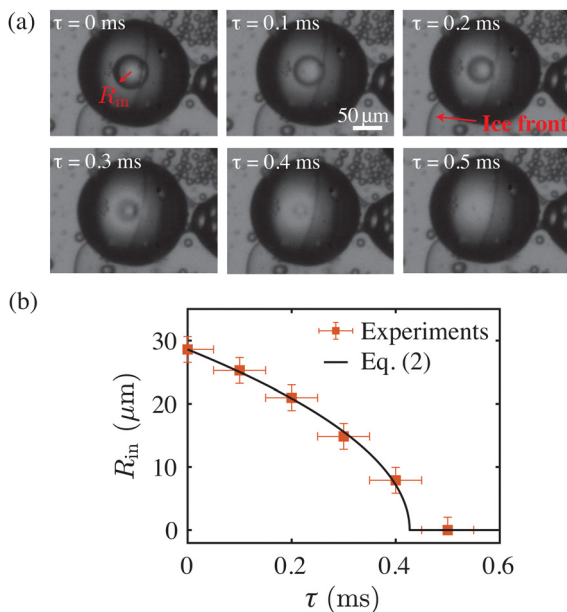


Fig. 3 (a) Experimental snapshots showing the sudden dissolution of the inner drop, here triggered by the ice front passing over the compound droplet (Movie 4, ESI†). (b) Radius  $R_{in}$  of the inner water drop as function of time (boxes). The rate of shrinking for several of such cases follows the steady Epstein–Plesset model of dissolution (solid line, eqn (2)).

characteristic features. For this, we rely on experiments performed with metastable W/O/W emulsions. In these instances, especially for relatively large compound droplets with an inner drop size greater than  $R_{in} \geq 30 \mu\text{m}$ , the dissolution process is directly measured without any hindrance from the ethanol vapour cloud, see Fig. 3(a). Note that the sudden dissolution of the inner drop in these barely stable compound droplets is triggered by disturbances in the ambient conditions. The sequence of images in Fig. 3(a) exemplifies one such case, where the inner drop dissolves while the compound droplet is getting engulfed in the solidifying bulk. The rate at which the inner drops dissolve, here directly measured, agrees well with the simplified Epstein–Plesset model,<sup>30,31</sup>

$$\frac{dR_{in}}{d\tau} = -\frac{\alpha}{\rho_w R_{in}(\tau)}, \quad \text{thus} \quad R_{in}^2 = R_{0,in}^2 - \frac{2\alpha}{\rho_w}\tau, \quad (2)$$

with  $\alpha = D\Delta c$ , where  $D$  and  $\Delta c$  are the unknown diffusion coefficient and concentration gradient, respectively and  $R_{0,in}$  is the initial drop size. eqn (2) is fitted through the experimental data (see Fig. 3 (b)) to obtain  $\alpha \approx 9.6 \times 10^{-4} \text{ kg m}^{-1} \text{ s}^{-1}$ . A typical dissolution time scale arises as  $\tau_d = \rho_w R_{in}^2 / 2\alpha \approx 0.3 \text{ ms}$ , which is significantly smaller compared to the time it takes for the ethanol vapour cloud to fully expand, see Fig. 2(b).

Beyond the freezing driven liquid-to-vapor phase transition of the constituent liquid, another surprising feature of our system is that the topological transition of the double-emulsions is strongly dependent on the size of the inner drop. We find that below a critical size,  $R_{in,crit} \approx 19 \mu\text{m}$ , the compound droplets remain stable, see Fig. 4. Note that due to the applied procedure of double-emulsion preparation  $R_{out}$  and  $R_{in}$  cannot be varied

independently and their ratio, here  $R_{out}/R_{in} \approx 3.3$  (see Fig. 4(a)), is governed by the ethanol mass transfer rate during preparation<sup>24,25</sup>. A second control parameter is the rate of freezing, characterised by the advancing velocity  $V$ . Due to experimental limitations we are not able to capture both the engulfment and sudden transition of each drop. To nevertheless still qualitatively address the rate at which freezing occurred, we couple the rate of engulfment  $V$  to the experienced drop deformation  $\Gamma$  once incorporated into the ice, as it has been established that both are inversely related.<sup>19,20</sup> Here, a large deformation (larger  $\Gamma$ ) is related to slower freezing (smaller  $V$ ) and *vice versa*, where for the range of advancing velocities is approximately  $0.2 \mu\text{m s}^{-1} \lesssim V \lesssim 2 \mu\text{m s}^{-1}$ . In Fig. 4(b) we thus show that the critical size dependence remains unaffected by the extend of the experienced deformation  $\Gamma$  and hence by the rate of engulfment  $V$ .

To examine this unexpected size dependence we consider that the pressure in the inner drop,  $p_{in}(t)$ , needs to fall below the vapour pressure of ethanol,  $p_{v,eth}$ , at  $T = -3 \text{ }^\circ\text{C}$ , to trigger the topological transition. Due to the tension exerted by the partially solidifying oil on the inner drop a reduction in pressure is expected (ideally) as  $\Delta p_{in}(t) = -K\Delta V_{in}(t)/V_{in}$ , where  $K$  is the elastic bulk modulus of water and  $\Delta V_{in}(t)/V_{in}$  the relative change in

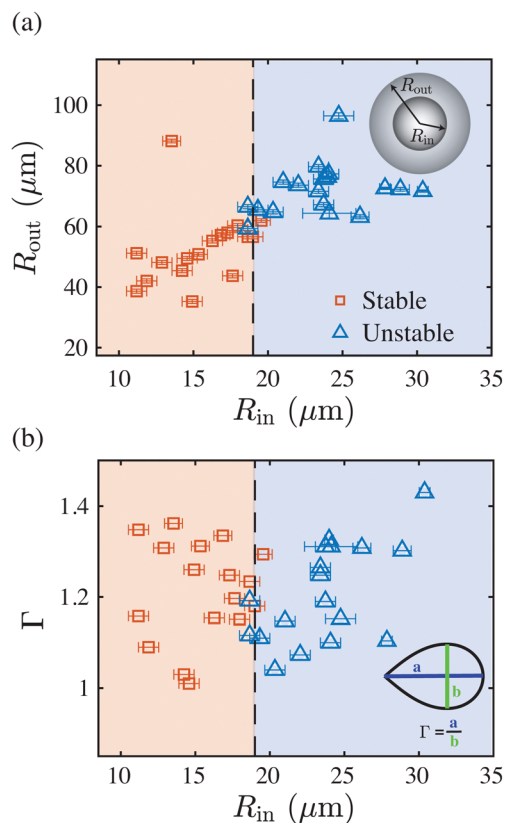


Fig. 4 Parameter space showing the dependence of the topological transition on (a) the size of the compound droplet and (b) its aspect ratio  $\Gamma$  after encapsulation by the ice. The latter is directly related to the (adjustable) advancing velocity  $V$  of the solidification front, *i.e.*,  $\Gamma \sim V^{-1}$  for pure oil drops in water.<sup>19,20</sup> Below a critical inner drop size,  $R_{in,crit} \approx 19 \mu\text{m}$ , the compound droplets remain stable. The rate of freezing does not affect this critical size.



volume over time. The initial pressure in the drop is given by the sum of ambient and Laplace pressure, resulting in the condition that the compound droplet remains stable if  $p_{\text{in,ref}} + 2\sigma/R_{\text{in}} - K\Delta V_{\text{in}}(t)/V_{\text{in}} > p_{\text{v,eth}}$ , or, in terms of the inner drop size, if  $R_{\text{in}} < 2\sigma/(p_{\text{v,eth}} + K\Delta V_{\text{in}}(t)/V_{\text{in}} - p_{\text{in,ref}})$ , where  $\sigma$  is the interfacial surface tension. Given the critical drop size,  $R_{\text{in,crit}}$ , observed experimentally (see Fig. 4) we conclude that the experienced change in volume over time,  $\Delta V_{\text{in}}(t)/V_{\text{in}}$ , stagnates and that therefore the amount of tension applied to the inner drop is limited. We suspect that, given the limitations of the experimental set-up and the bounded applied thermal gradient, sufficiently low temperatures could not be achieved and a thermal equilibrium is reached, causing only a small portion of the encapsulating oil to solidify.

## 4 Conclusion

In summary, in this Letter, we discussed that a dilute double-emulsion, *i.e.*, water-in-oil compound droplets in water, can undergo the topological transition from its W/O/W state to an O/W state after being frozen. The sudden topological transition, we argue, is triggered by the partial solidification of the encapsulating oil, putting tension on the inner drop, causing the trace amounts of liquid ethanol inside to undergo a liquid-to-vapour phase transition. As the vapour cloud expands, ethanol is depleted, making the compound drop unstable, and the inner drop dissolves. The then expelled interior accumulates at the warmer end and migrates though the ice in the direction of motion of the advancing solidification front. The topological transition critically depends on the size of the inner drop and does not depend on the engulfment velocity  $V$ . We found that below  $R_{\text{in,crit}} \approx 19 \mu\text{m}$  the topological transition does not occur and the double-emulsion remains stable. We rationalised that this size dependence originates from the limited amount of tension applied to the inner drop. Our findings on the individual compound droplets can be extrapolated to those in a more densely packed double-emulsion, where the same phenomena is observed (see ESI†).

The surprising topological transition we found is relevant when freezing multi-component systems and in the context of the cryopreservation procedures of food emulsions and bio-specimen. In addition, our findings highlight the effect of phase changing liquid inclusions in a solidifying material and how this might alter their topology in unexpected ways.

## Conflicts of interest

There are no conflicts to declare.

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