Theory of Specific Heat in Glass Forming Systems

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Experimental measurements of the specific heat in glass-forming systems are obtained from the linear response to either slow cooling (or heating) or to oscillatory perturbations with a given frequency about a constant temperature. The latter method gives rise to a complex specific heat with the constraint that the zero frequency limit of the real part should be identified with thermodynamic measurements. Such measurements reveal anomalies in the temperature dependence of the specific heat, including the so called “specific heat peak” in the vicinity of the glass transition. The aim of this paper is to provide theoretical explanations of these anomalies in general and a quantitative theory in the case of a simple model of glass-formation. We first present new simulation results for the specific heat in a classical model of a binary mixture glass-former. We show that in addition to the formerly observed specific heat peak there is a second peak at lower temperatures which was not observable in earlier simulations. Second, we present a general relation between the specific heat, a caloric quantity, and the bulk modulus of the material, a mechanical quantity, and thus offer a smooth connection between the liquid and amorphous solid states. The central result of this paper is a connection between the micro-melting of clusters in the system and the appearance of specific heat peaks; we explain the appearance of two peaks by the micro-melting of two types of clusters. We relate the two peaks to changes in the bulk and shear moduli. We propose that the phenomenon of glass-formation is accompanied by a fast change in the bulk and shear moduli, but these fast changes occur in different ranges of the temperature. Lastly, we demonstrate how to construct a theory of the frequency dependent complex specific heat, expected from heterogeneous clustering in the liquid state of glass formers. A specific example is provided in the context of our model for the dynamics of glycerol. We show that the frequency dependence is determined by the same α-relaxation mechanism that operates when measuring the viscosity or the dielectric relaxation spectrum. The theoretical frequency dependent specific heat agrees well with experimental measurements on glycerol. We conclude the paper by stating that there is nothing universal about the temperature dependence of the specific heat in glass formers - unfortunately one needs to understand each case by itself.

I. INTRODUCTION

The traditional measurements of the specific heat $C_v$ at constant volume or $C_P$ at constant pressure involve cooling (or heating) the sample at a constant rate [1, 2]. When applied to glass-forming systems, this approach has an inherent difficulty. Since glass-forming system tend to relax to equilibrium slower and slower as the temperature is lowered, at some point the ‘constant rate’ of cooling becomes too high for the system to respond to, and then the system does not reach equilibrium. Typically the specific heat then drops abruptly, giving rise to the “specific heat peak” at some temperature which is sometimes identified as the glass transition temperature $T_g$. Needless to say, such a definition of transition temperature is less than compelling, since it clearly depends on the rate of cooling and is not inherent to the system properties.

In an attempt to overcome this difficulty Birge and Nagel [3] introduced ”specific heat spectroscopy”. In this technique one keeps the sample close to a temperature $T$ at all times, but perturbs it periodically with a small-amplitude oscillation of frequency $\omega$. Linear response theory then relates the amount of heat exchanged at that frequency, $\delta Q(\omega)$ to the oscillatory temperature perturbation $\delta T(\omega)$ via the relation

$$\delta Q(\omega) = C(\omega)\delta T(\omega) \quad (1)$$

where $C(\omega)$ is the frequency dependent specific heat that can be measured at either constant volume or constant pressure. In order to find the thermodynamic specific heat one needs to extrapolate data to the $\omega \to 0$ limit. Whether or not this extrapolation overcomes the above-mentioned worry of sufficient relaxation time is an issue that has not been fully clarified in the literature.

In this paper we concern ourselves with the theoretical calculation of the specific heat in glass-forming systems and in the relation of the specific heat to other material properties. To this aim we focus on one simulational example (a binary mixture of point particles interacting via an $r^{-12}$ repulsive potential) and one experimental example (glycerol). In the context of the first example we present results of new simulations that exhibit two distinct peaks in the curve of the specific heat vs. the temperature. We present for this example various theoretical results, culminating with a new scenario to explain the specific heat peaks, i.e. the micro-melting of clusters. We believe that this is the central point of the present paper. To understand the nature of the specific heat anomalies one must understand the physics that is behind the glassy behavior of this model in general and
the existence of the two specific heat peaks in particular. When the temperature is lowered at a fixed pressure this system \([4]\) (as well as many other glass-formers \([5–10]\)) tends to form micro-clusters of local order. In the present case large particles form long-lived patches of hexagonal ordering first (starting at about \(T = 0.5\), and at lower temperatures (around \(T = 0.1\)) also the small particles form long-lived hexagonal clusters. The clusters are not that huge, with at most \(O(100)\) particles, (cf. Fig. 1), depending on the temperature and the aging time. But we have shown that the long time properties of correlation functions are entirely carried by the micro-clusters [4]. Below we will refer to the micro-clusters as curds and the liquid phase as whey. We will argue that the specific heat responds to the micro-melting of the clusters - those of small particles at the lowest temperatures and those of the larger particles at higher temperatures. The large increase in the number of degrees of freedom when a particle leaves a crystalline cluster and joins the liquid background is the basic reason for the increase in entropy that is seen as a specific heat peak.

In the context of the second example we show that the calculation of the frequency dependent specific heat is easy when we have a reasonable model of the glassy relaxation. Having such a model for glycerol [11], we demonstrate in section IV that the information gained from the frequency dependent specific heat is very similar to that learned from other linear response functions like broad-band dielectric spectroscopy. We will be able therefore to present spectra of the frequency dependent specific heat in close correspondence with experiments.

The specific heat has interesting relations with the mechanical moduli of the material, and we present relations (which pertain to any system with an \(r^{-n}\) potential) to the bulk and shear moduli. As a result of our thinking we conclude that the bulk and shear moduli change rapidly in the temperature range of the two distinct specific heat peaks mentioned above. The relation to the bulk modulus is explicit, and is shown rigorously in Subsect. II C. The relation to the shear modulus is less explicit, simply because one does not have an equation of state with strain (a quantity that is ill defined in the context of glasses). Nevertheless we present a conjecture that the bulk and shear moduli in generic glasses may change rapidly at two different temperature ranges.

The structure of the paper is as follows: In Sect. II we present the model glass consisting of a binary mixture of point particles interacting via soft potentials, and discuss its thermodynamic properties. We derive exact equations for its specific heat at constant volume, which are correct at all temperatures through the glass transition. We compare these results to molecular dynamics simulations in which special care had been taken to equilibrate the system, summarizing a computational effort of about two years. The main conclusion of this section is that the details of the interaction potential are crucial in determining what the specific heat does in the vicinity of the glass transition, and there is nothing universal about it. For a simple enough potential we can derive a theory that is in excellent agreement with simulations up to the first specific heat peak. To explain both peaks we must present a theory that takes into account explicitly the tendency of the system to form micro-clusters [4]. The state of the system then becomes like curds of local crystalline order embedded in a whey of disordered fluid. It is the freezing or melting of these curds that account quantitatively for the specific heat peaks, as is shown in Subsect. III. Below we use interchangeably the words ‘clusters’ and ‘curds’. In Sect. IV we turn to discussing the frequency-dependent complex specific heat. To construct a theory of this object one needs a model of the dynamics of the system under study, be it glycerol or any other material. We demonstrate, using our dynamical model of glycerol [11], how this measurement is equivalent in terms of its dynamical contents to any other linear response to an oscillatory perturbation. We present theoretical spectra and show satisfactory agreement with the experiments. The paper ends in Sect. V where we draw conclusions and summarize the results and the implications of our calculations.

II. THE BINARY MODEL AND ITS SPECIFIC HEAT

The model discussed here is the classical example \([12, 14]\) of a glass-forming binary mixture of \(N\) particles in a 2-dimensional domain of area \(V\), interacting via a soft \(1/r^{12}\) repulsion with a ‘diameter’ ratio of 1.4. We refer the reader to the extensive work done on this system.
[12, 14–17]. The sum-up of this work is that the model is a *bona fide* glass-forming liquid meeting all the criteria of a glass transition.

In short, the system consists of an equimolar mixture of two types of particles, “large” with ‘diameter’ $\sigma_2 = 1.4$ and “small” with ‘diameter’ $\sigma_1 = 1$, respectively, but with the same mass $m$. In general, the three pairwise additive interactions are given by the purely repulsive soft-core potentials

$$\phi_{ab}(r) = \epsilon \left( \frac{\sigma_{ab}}{r} \right)^n, \quad a, b = 1, 2, \quad (2)$$

where $\sigma_{aa} = \sigma_a$ and $\sigma_{ab} = (\sigma_a + \sigma_b)/2$. The cutoff radii of the interaction are set at $4.5\sigma_{ab}$. The units of mass, length, time and temperature are $m, \sigma_1, \tau = \sigma_1 \sqrt{m/\epsilon}$ and $\epsilon/k_B$, respectively, with $k_B$ being Boltzmann’s constant. In numerical calculations the stiffness parameter of the potential (2) was chosen to be $n = 12$.

We turn now to the analysis of the specific heat of this model as a function of the temperature.

### A. Specific heat (simulations)

The specific heat capacity (specific heat) at constant volume is defined by:

$$\frac{C_V}{N} = \frac{d}{2} + \frac{\partial}{\partial T} \frac{\langle U \rangle}{N} \bigg|_V, \quad (3)$$

where $d$ is the space-dimension and the potential energy of a binary mixture is given by:

$$U = \frac{1}{2} \sum_{i \neq j} \phi_{ab}(r_{ij}). \quad (4)$$

Here $r_{ij}$ is the distance between particles $i$ and $j$. The average value of the potential energy is defined by averaging over configurational space $\Gamma$:

$$\langle U \rangle = \int U \exp\left(-\frac{U}{T}\right) d\Gamma \int \exp\left(-\frac{U}{T}\right) d\Gamma. \quad (5)$$

Substitution of (5) into (3) yields the following expression for the specific heat:

$$\frac{C_V}{N} = \frac{d}{2} + \frac{\langle U^2 \rangle - \langle U \rangle^2}{NT^2}. \quad (6)$$

The specific heat of our binary mixture model was measured at constant volume in [13, 14, 18] and by us. In simulations one can measure the specific heat directly from its definition (3) or (6). We have used the last equation which allows one to estimate the specific heat in a single run of the canonical ensemble Monte Carlo simulations. At each temperature the density was chosen in accordance with the simulation results in an NPT ensemble as described in [14] with the pressure value fixed at

$P = 13.5$. As the initial configuration in the Monte Carlo process the last configuration of the molecular dynamics run for this model at given temperature after $1.3 \times 10^8$ time steps was used. After short equilibration the potential energy distribution functions were measured during $2 \times 10^6$ Monte Carlo sweeps. The acceptance rate was chosen to be 30%. Simulations were performed with $N = 1024$ particles in a square cell with periodic boundary conditions.

Examples of the spline interpolation of the potential energy distribution for a few temperatures are shown in Fig.2. The first and second moments of these distributions define the average value of the potential energy (Fig.3) and the specific heat (Fig.4). We stress that these results were computed at constant volume, such that the volume corresponds to simulations in NPT ensemble with the pressure $P = 13.5$ [14] at each temperature.

One can see from these figures that the behavior of both quantities, the first and second moments of the distribution, change abruptly in the vicinity of $T \sim 0.5$. The specific heat displays a maximum in the temperature dependence. Our simulations appear to provide trustable values of $C_V$ down to lowest temperatures where the value of the specific heat coincides with that of two-dimensional solid, i.e. $C_V = 2$. What could not be seen in earlier simulations is that there is a much smaller second peak of the specific heat at lower temperatures. To resolve it to the naked eye we present in Fig. 4 a blow-up of the region of lowest temperatures where the second peak is more obvious. To understand the nature of the specific heat anomalies we turn now to a theoretical analysis of the caloric equation of state in order to study the specific heat using the definition (3). The physical origin of the two peaks will be explained in Subsect. III. The
FIG. 3: Color online: in dots: the temperature dependence of the average potential energy per in the binary mixture model, computed at constant volume, such that the volume agrees with the pressure $P=13.5$ [14] at each temperature. The continuous line represents the approximation furnished by the virial expansion, which obviously fails for $T < 0.5$.

FIG. 4: Color online: in dots: the temperature dependence of the specific heat in the binary mixture model, computed at constant volume, such that the volume agrees with the pressure $P=13.5$ [14] at each temperature. The data indicate the existence of two specific heat peaks, one prominent at about $T = 0.5$ and a smaller one at about $T = 0.1$, and see the inset for finer detail. The continuous line represents the approximation furnished by the virial expansion, which obviously fails for $T < 0.5$.

B. Specific heat (series expansions)

The general expression for the pressure (thermal equation of state) obtained from the virial theorem is given by:

$$P = \rho T - \frac{\rho}{2d} \cdot \frac{1}{N} \sum_{i \neq j} r_{ij} \frac{\partial \phi_{ab}(r_{ij})}{\partial r_{ij}} ,$$

(7)

where $\rho$ is the particle number density. The potential is a homogenous function of degree $-n$ (Eq. (2)), therefore:

$$r \frac{\partial \phi_{ab}(r)}{\partial r} = -n \phi_{ab}(r) .$$

(8)

Due to this property of the interaction potential we find a connection between the pressure and the temperature and mean energy:

$$P = \rho T + \frac{n}{d} \rho \langle U \rangle / N .$$

(9)

This equation is exact for one component and multicomponent systems and is valid at all temperatures, from liquid to solid.

The next simplification for systems with an inverse power inter-particle interaction consists in the dependence of all excess thermodynamic properties relative to the ideal gas on a single density-temperature variable [19, 20]. To see why, recall that for a one component system the canonical partition function $Z_N$ is defined by:

$$Z_N = \frac{1}{N! \Lambda dN} \int \exp \left( - \sigma n \frac{\epsilon}{k_B T} \sum_i \frac{1}{r_{ij}} \right) d^d r .$$

(10)

Here $\Lambda = \hbar/(2\pi mk_BT)^{1/2}$ is the thermal de Broglie wavelength. The typical distance between particles is given by

$$l = \left( \frac{V}{N} \right)^{1/d} .$$

(11)

Thus one can use new variables in the integral (10), $\vec{s} = \vec{r}/l$, and the canonical partition function can be rewritten as:

$$Z_N = \frac{V^N}{N! \Lambda^{dN}} \frac{1}{N^N} \int \exp \left( - \rho n/d \frac{\epsilon}{k_B T} \sum_i \frac{1}{s_{ij}} \right) d^d s .$$

(12)

where the dimensionless particle number density is defined by $\rho = \frac{N}{V} \sigma^d$. This way of writing the partition function underlines the existence of the ideal gas contribution before the configurational integral, and the dependence of the configurational integral, in the case of one component, on a natural parameter, $\Gamma = \rho (k_B T)^{d/n}$.

In the case of a multicomponent system the properties of the mixture can be approximated by those of a one
component reference fluid [21] with an effective diameter defined by:

$$\sigma_e^d = \sum_{a,b} x_a x_b \sigma_{ab}^d,$$

where $x_a = N_a/N$ is the particle number concentration. Therefore, the properties of a mixture are defined by the effective parameter:

$$\Gamma_e = \Gamma \left( \frac{\sigma_e}{\sigma_1} \right)^d.$$

Substitution of the equation (16) to (9) yields the caloric equation of state, the specific heat is calculated after that by (3). The density corresponding to a given temperature is defined as the solution of the equation (16) at $P = 13.5$.

Results of the calculations in the frame of the virial approach are compared with the simulation results in Figs. 3 and 4. Clearly, the virial expansion (16) cannot be trusted for temperatures lower than $T = 0.5$ since it was computed from simulations that did not go below that temperature. This temperature is precisely the point at which small micro-clusters become significant, and a pure liquid homogenous state description of the glassy phase breaks down. Indeed, down to that temperature the prediction of equation (3) together with the virial expansion fits the data excellently well. To understand what happens at lower temperatures we must await Sect. III where the existence of micro-clusters is taken explicitly into account. Some comments about the limit of temperature going to zero and the relation to the Madelung constant can be found in Appendix A.

### C. Specific heat (mechanical approach)

In this subsection we connect the specific heat to the bulk modulus of the system. To this aim we begin with the microscopic definition of the stress-tensor in an NVT ensemble (see, e.g. [22]):

$$\sigma_{\alpha\beta} = \frac{1}{V} \left( \sum_i p_i^\alpha p_i^\beta - \frac{1}{2} \sum_{j \neq i} \frac{\partial\phi_{ab}(r_{ij})}{\partial r_{ij}} r_{ij}^\alpha r_{ij}^\beta \right),$$

where $p_i^\alpha$ is the $\alpha$ component of the dimensionless momentum of particle $i$ and $r_{ij}^\alpha$ is the $\alpha$ component of the vector joining particles $i$ and $j$. The first invariant of the stress tensor is its trace:

$$V (\sigma_{xx} + \sigma_{yy}) = \left( \sum_i (p_i^x)^2 + \sum_i (p_i^y)^2 \right) + n U.$$

In order to average (18) one has to take into account that:

$$\left( \sum_i (p_i^x)^2 \right) = N \langle (p_i^x)^2 \rangle = N \cdot T.$$

Thus the average of the first invariant (18) is:

$$\langle (\sigma_{xx} + \sigma_{yy}) \rangle = 2 \rho T + n \rho \frac{(U)}{N}.$$

The pressure is defined as $P = \langle (\sigma_{xx} + \sigma_{yy}) \rangle/2$ and (20) yields (9).

The square of (18) is:

$$V^2 (\sigma_{xx}\sigma_{xx} + \sigma_{yy}\sigma_{yy} + 2\sigma_{xx}\sigma_{yy}) = \left( \sum_i (p_i^x)^2 + \sum_i (p_i^y)^2 \right)^2 + 2n \left( \sum_i (p_i^x)^2 + \sum_i (p_i^y)^2 \right) U + n^2 U^2.$$

To compute the average of this equation we need to use the fact that $\langle (p_i^x)^2 \rangle = T$ and $\langle (p_i^y)^4 \rangle = 3 T^2$. After averaging (21) is written as:

$$V^2 \left( \langle \sigma_{xx} \sigma_{xx} \rangle + \langle \sigma_{yy} \sigma_{yy} \rangle + 2 \langle \sigma_{xx} \sigma_{yy} \rangle \right) = 4 N T^2 + 4 N^2 T^2 + 4 N T \langle U \rangle + n^2 \langle U \rangle^2.$$

The square of (20) is given by:

$$V^2 \left( \langle \sigma_{xx} \rangle^2 + \langle \sigma_{yy} \rangle^2 + 2 \langle \sigma_{xx} \rangle \langle \sigma_{yy} \rangle \right) = 4 N^2 T^2 + 4 N T \langle U \rangle + n^2 \langle U \rangle^2.$$
After subtraction of (23) from (22) we have:
\[
V^2 \left[ (\langle \sigma_{zx} \sigma_{zx} \rangle - \langle \sigma_{zx} \rangle^2) + (\langle \sigma_{yy} \sigma_{yy} \rangle - \langle \sigma_{yy} \rangle^2) + 2(\langle \sigma_{zx} \sigma_{yy} \rangle - \langle \sigma_{zx} \rangle \langle \sigma_{yy} \rangle) \right] = 4NT^2 + n^2(\langle U^2 \rangle - \langle U \rangle^2).
\] (24)

This equation has a well-defined thermodynamic limit since the quantity in square brackets scales like \(V^{-1}\). This is seen explicitly in Eq. (25).

These results allow us now to find exact relationships between the specific heat, a caloric quantity, and the elastic constants which are mechanical quantities. To do so we recall the definitions of the elastic constants through the stress fluctuations (see, e.g., [23]):
\[
\frac{V}{T} (\langle \sigma_{a}^\alpha \sigma_{a}^\beta \rangle - \langle \sigma_{a} \rangle \langle \sigma_{a} \rangle) = 2\rho T (\delta_{a\alpha} \delta_{a\beta} + \delta_{a\beta} \delta_{a\alpha}) + (C_{a\alpha\beta\delta} - C_{a\beta\alpha\delta}),
\]
(25)

where \(C_{a\alpha\beta\delta}\) are the elastic constants and \(C_{a\alpha\beta\gamma\delta}\) are the so-called Born terms which determine the instantaneous elastic constants for any given configuration.

Substitution of (25) to (24) yields:
\[
n^2(\langle U^2 \rangle - \langle U \rangle^2) = 4NT^2 + V \left[ (C_{x\alpha\alpha}^B + C_{y\beta\beta}^B + 2C_{x\alpha\alpha}^B) - (C_{x\alpha\alpha} + C_{y\beta\beta} + 2C_{x\alpha\alpha}) \right]
\] (26)

The compression (bulk) modulus in two-dimensional systems is:
\[
K = \frac{1}{4} (C_{x\alpha\alpha} + C_{y\beta\beta} + 2C_{x\alpha\alpha}^B).
\] (27)

Recalling Eq. (6) and substituting it and Eq. (27) into (26) yields:
\[
\frac{C_V}{N} = 1 + 4K^\infty - K
\] (31)

where the bulk modulus in the infinite frequency limit \(K^\infty = \rho T + K^B\) and the Born term is defined by the interparticle interactions [24]:
\[
K^B = \frac{1}{4V} \sum_{i \neq j} r_{ij} \left( \frac{\partial^2 \phi_{a b}(r_{ij})}{\partial r_{ij}^2} - \frac{\partial \phi_{a b}(r_{ij})}{\partial r_{ij}} \right).
\] (29)

For the present model the “infinite frequency” term (cf. [24]) is given by:
\[
K^\infty = \rho T + \frac{n(n + 2)}{4} \rho \langle U \rangle / N.
\] (30)

This is as much as one can go using exact identities. We reiterate that Eq. (28) is very interesting, allowing us to connect the bulk modulus to the specific heat. In fact, this connection implies that the specific heat measures the difference between the bulk modulus and its infinite frequency limit. At low temperatures this difference in the harmonic approximation as given by:
\[
K^\infty - K = \frac{n^2}{4} \rho T,
\] (31)

independent of the solid structure in contrast to the shear modulus (cf. [17]).

The bulk modulus \(K\) cannot be computed exactly using identities, and we need further information to evaluate it. Fortunately we can estimate the bulk modulus from the virial expansion (7) at \(T > 0.5\), since:
\[
K = \frac{\partial P}{\partial \rho}.
\] (32)

Having done so we can compare the measurements to what we expect theoretically. The specific heat as predicted from the virial expansion is shown in Fig. 4 as the blue (continuous) line. We should stress that computing \(C_V\) from either Eq. (3) or Eq. (28) (using the virial expansion (16), yield essentially identical results that cannot be distinguished in the blue line in Fig. 4 for \(T > 0.5\).

III. SPECIFIC HEAT - THE PHYSICAL EXPLANATION

In this section we propose the physical picture behind the existence of the specific heat peaks. We argue that the specific heat responds to the micro-melting of the clusters - those of small particles at the lowest temperatures and those of the larger particles at higher temperatures. The large increase in the number of degrees of freedom when a particle leaves a crystalline cluster and joins the liquid background is the basic reason for the increase in entropy that is seen as a specific heat peak. We argue that the specific heat peaks should be computed from either Eq. (3) or Eq. (28) (using the virial expansion (16), yield essentially identical results that cannot be distinguished in the blue line in Fig. 4 for \(T > 0.5\).

A. Mechanical Equation of State

In this subsection we employ the mechanical equation of state derived above from which the specific heat will be computed. To start we define \(\epsilon_{c_w}^L\), \(\epsilon_{c_w}^S\), \(\epsilon_{c_l}^L\) and \(\epsilon_{c_l}^S\) respectively as the volume of large particle in the whey, small particle in the whey, large particle in the solid and small particle in the solid. Similarly we denote by \(\epsilon_{c_w}^L\), \(\epsilon_{c_w}^S\), \(\epsilon_{c_l}^L\) and \(\epsilon_{c_l}^S\) the energy of a large and small particle in the in the whey and in the crystalline phase respectively.
Needless to say, all these quantities are temperature and pressure dependent; we will therefore explicitly use our low temperature knowledge concerning $v_c^g$ and $v_w^g$ in the crystalline phase, but treat the difference $v_c^g - v_w^g$ and $v_s^g - v_c^g$ as constants that we estimate below from our simulation knowledge. Similarly we estimate $c_c^g$ and $c_c^w$ from our knowledge of the hexagonal lattices at $T = 0$. We assume that $c_w^g \approx c_c^g$ and similarly $c_w^s \approx c_c^s$ since our simulations indicate a very small change in these parameters, see Table I. It should be stressed that the enthalpy change at these pressures are almost all due to the $PV$ term. This will result in a semi-quantitative theory as presented in [16] to get more accurate estimates

$$p_c^g(P, T) \approx \frac{1}{1 + g_w^g e[(c_c^g - c_c^w) + P(c_w^g - c_w^g)]/T} \quad (34)$$

$$p_c^s(P, T) \approx \frac{1}{1 + g_w^s e[(c_c^g - c_c^w) + P(c_w^g - c_w^g)]/T} \quad (35)$$

It is important to note that the combination of Eq. (33) together with Eqs. (34) and (35) provides a mechanical equation of state that is alternative to the virial expansion presented above. Whereas the latter is best at temperature higher than $T \approx 0.5$ we expect the present one to be best at low temperatures because only the present approach takes into account the formation of clusters explicitly. The virial expansion by construction is a liquid theory. We will now compute $C_v$ directly from Eq. (28). The peaks in the specific heat are determined by the temperature dependence of $p_c^g(P, T)$ and $p_c^s(P, T)$ each which has a temperature and pressure derivatives that peaks at a different temperature, denoted as $T^g(P)$ and $T^s(P)$, and see below for details. As said above we take $\Delta v^g \equiv v_c^g - v_w^g$ and $\Delta v^s \equiv v_s^g - v_c^g$ as approximately constants (as a function of temperature and pressure). The constants are estimated from the condition that the second temperature derivative of $p_c^g(P, T)$ and $p_c^s(P, T)$ should vanish. Explicitly:

$$\Delta v^g \approx T^g(P^*) \ln g_w^g/P^* \quad, \quad \Delta v^s \approx T^s(P^*) \ln g_w^s/P^* \quad (36)$$

where $P^*$ is the pressure for which the peaks in the derivatives are observed (13.5 in our simulations). This is equivalent to a linear dependence of the specific heat peaks as a function of pressure, $T^g(P)/T^g(P^*) = P/P^*$ and similarly for the small particles.

In terms of these objects we can rewrite

$$v = v_c(P, T) + \Delta v^g(1 - p_c^g) + \Delta v^s(1 - p_c^s) \quad, \quad \left(\frac{\partial v}{\partial P}\right)_T = \left(\frac{\partial v_c}{\partial P}\right)_T - \Delta v^g\left(\frac{\partial p_c^g}{\partial P}\right)_T - \Delta v^s\left(\frac{\partial p_c^s}{\partial P}\right)_T \quad (37)$$

To compute the temperature dependence of $\left(\frac{\partial v}{\partial P}\right)_T$ we need first to determine its $T \to 0$ limit, which is determined by the first term on the RHS of Eq. (37) as the other terms on the RHS decay exponentially fast when $T \to 0$. Since we have already exact results for the bulk modulus for the present model, we return to Eqs. (28) and (30). We know on the one hand that $\lim_{T\to0}C_v = 2$ and that $(U)/N \approx 2.94$ over the whole interesting temperature range, cf. Fig. 3. The compressibility $\kappa$ is related to the bulk modulus via $\kappa = -\left(\frac{\partial v}{\partial P}\right)_T/v = 1/K$ and therefore easily estimated as $T \to 0$ since there $\left(\frac{\partial v}{\partial P}\right)_T \approx -1/(123.5 - 35T)$. We use this approximation up to $T \approx 0.5$.

Having all the ingredients we can compute $C_v/N$. The parameters used were estimated from the numerical simulation and are summarized in Table I. Since the aim of this subsection is only semi-quantitative, we do not make any attempt of parameter fitting, and show the result of the calculation in Fig. 5.

| $v_c^g$ | $v_c^w$ | $v_w^g$ | $v_w^s$ | $v_s^g$ | $v_s^w$ |
|--------|--------|--------|--------|--------|--------|
| 3.69   | 2.07   | 3.76   | 2.16   | 1.43   | 0.92   | 1.58   | 0.94   |

Table I: Parameters used in the calculation of the specific heat.

As the condensed phase consists of clusters of large and small particles, we use the notation $N_c^p$ for the number of clusters of $n$ large particles and $N_w^p$ for the clusters of $m$ small particles. In the next subsection we write the energy of our system explicitly in terms of these cluster numbers. Here however we only need the intensive variables $p_c^g = 2 \sum_n N_c^p/N, p_w^g = 2 \sum_m N_w^p/N, p_c^s = 2 N_c^s/N$ and $p_w^s = 2 N_w^s/N$ which stand for the fraction of large particles and small particles in the curds, and large particles and small particles in the whey, such that $p_c^g + p_w^g = 1$ and $p_c^s + p_w^s = 1$. Using these variables we can write an expression for the volume per particle $v \equiv V/N$:

$$v = \frac{v_w^g + v_c^g - v_c^w}{2} + \frac{v_c^g - v_w^g}{2} p_c^g + \frac{v_s^g - v_c^s}{2} p_c^s$$

(33)

At this point we need to derive expressions for $p_c^g$ and $p_c^s$. To do so we need to remember that in the relevant range of temperatures the large particles in the whey can occupy either hexagonal or heptagonal Voronoi cells, whereas small particles can occupy only pentagonal or hexagonal cells [4, 15, 16]. Accordingly there are $g_w^g \approx (2^6 - 1)/6 + 2^7/7$ ways to organize the neighbours of a large particle in the whey (neglecting the rare large particle in heptagonal neighbourhood), but only one way in the cluster. Similarly, there are $g_w^s \approx (2^6 - 1)/6 + 2^5/5$ ways to organize a small particle in the whey. We note that this estimate assumes that the relative occurrence of the different Voronoi cells is temperature independent. While reasonable at higher temperatures [4], at lower temperature one should use the full statistical mechanics as presented in [16] to get more accurate estimates of $g_w^g$ and $g_w^s$. This is not our aim here; we aim at a physical understanding of the specific heat peaks rather than an accurate theory. We thus end up with the simple estimates
Indeed, the theoretical calculation exhibits the existence of two, rather than one, specific heat peaks. We can now explain the origin of the peaks as resulting from the derivatives \( \frac{\partial E}{\partial T} \), and \( \frac{\partial E}{\partial P} \). These derivatives change most abruptly when the micro-clusters form (or dissolve), each at a specific temperature determined by \( (h^w_w - h^s_w) / \ln g^w_w \) and \( (h^c_w - h^s_c) / \ln g^c_w \). Note that there can be pressures (both upper and lower boundaries) where the sign of \( (h^w_w - h^s_w) \) or \( (h^c_w - h^s_c) \) change sign and the peak can be lost.

**B. Caloric Equation of State**

In this subsection we present a more general approach which does not take direct input from results derived for the inverse power potential. Thus although we use below some parameters read from the simulation, the derivation is very general any pertains to any distribution of clusters. To this aim we derive a second equation of state, a caloric one. It is quite standard to have two equations of state, only for ideal gas and inverse potentials the two equations of state degenerate into one.

Denote the total energy of the system as a sum of \( E_c \), the energy of the clusters (or curds), and \( E_w \), the energy of the liquid background (or whey), i.e:

\[
E = E_c + E_w .
\]

These energies are sums over the degrees of freedom - translational, rotational, vibrational and configurational:

\[
E_c = E_{tr,c} + E_{rot,c} + E_{vib,c} + E_{conf,c} ,
\]

\[
E_w = E_{tr,w} + E_{vib,w} + E_{conf,w} .
\]

To estimate \( E_{tr,c} \) we consider the number \( N^c_n \) of clusters of \( n \) large particles and \( N^s_m \) of clusters of \( m \) small particles and write

\[
E_{tr,c} = T\left[\sum_n N^c_n + \sum_m N^s_m\right] .
\]

On the other hand in the whey we follow Eyring [25] and Granato [26] and write

\[
E_{tr,w} = T[N^c_w + N^s_w]f .
\]

where \( f = 1 - V^{(s)}_w / V_w \) is defined as the fraction of free volume in the liquid phase compared to the equivalent solid crystalline phase. In other words,

\[
V_w = N^c_w V^c_w + N^s_w V^s_w .
\]

Similarly, we write

\[
E_{rot,c} = \frac{1}{2} T\left[\sum_n N^c_n + \sum_m N^s_m\right] ,
\]

\[
E_{vib,c} = 2T\left[\sum_n N^c_n + \sum_m N^s_m m\right] ,
\]

\[
E_{vib,w} = 2T[N^c_w + N^s_w](1 - f) ,
\]

\[
E_{conf,c} = \epsilon^c_c \sum_n N^c_n + \epsilon^s_c \sum_m N^s_m ,
\]

\[
E_{conf,w} = \epsilon^c_w N^c_w + \epsilon^s_w N^s_w .
\]

In terms of these variables we can rewrite

\[
E_{tr} = \frac{N}{2} \sum_n \frac{p^c_n}{\langle n \rangle} + \frac{p^s_n}{\langle m \rangle} + \langle p'_w + p''_w \rangle f ,
\]

\[
E_{rot,c} = \frac{NT}{4} \left( \frac{p^c}{\langle n \rangle} + \frac{p^s}{\langle m \rangle} \right) ,
\]

\[
E_{vib} = NT\left[2(1 - f) + \langle p'_c + p''_c \rangle \right] ,
\]

\[
E_{conf} = \frac{N}{2} \left[ \epsilon^c_c + \epsilon^s_c + (\epsilon^s_c - \epsilon^c_w)p'_c + (\epsilon^s_c - \epsilon^c_w)p''_c \right] .
\]

Summing up all the contributions we need to pay attention to the order of magnitude of the various terms. Since we expect the average size of clusters, at the temperatures of interest, to be of the order of 30 or so, we can neglect safely all the terms that have average size clusters in the denominator. With this in mind the expression for the energy of the system takes the form

\[
\frac{E}{N} = T[2 - \frac{2 - p'_c - p''_c}{2} f] + \frac{\epsilon^c_c + \epsilon^s_c}{2} + \frac{\epsilon^c_w - \epsilon^s_w}{2} p'_c + \frac{\epsilon^c_w - \epsilon^s_w}{2} p''_c .
\]
This is the caloric equation of state that we were after. Using it, we can compute \( C_v \) directly from the thermodynamic identity

\[
\frac{C_v}{N} = \frac{C_p}{N} - T \frac{\nu \alpha^2}{\kappa}, \tag{56}
\]

where the thermal expansion coefficient is

\[
\alpha = \left( \frac{1}{v} \frac{\partial v}{\partial T} \right)_p = \frac{1}{v} \left[ \frac{\nu (v_c^f - v_w^f)}{2} \left( \frac{\partial p_c^f}{\partial T} \right)_p + \frac{\nu (v_c^s - v_w^s)}{2} \left( \frac{\partial p_c^s}{\partial T} \right)_p \right],
\]

and the compressibility is

\[
\kappa = -\left( \frac{1}{v} \frac{\partial v}{\partial P} \right)_T = -\frac{1}{v} \left[ \frac{\nu (v_c^f - v_w^f)}{2} \left( \frac{\partial p_c^f}{\partial P} \right)_T + \frac{\nu (v_c^s - v_w^s)}{2} \left( \frac{\partial p_c^s}{\partial P} \right)_T \right].
\]

Having all the ingredients we can sum up the terms in Eq. (56). The parameters used were estimated from the numerical simulation and are summarized in Table 1. As before, we did not make any attempt for parameter fitting, and show the result of the calculation in Fig. 6.

\[
\frac{C_P}{N} = 2 - \frac{2 - p_c^f - p_c^s}{2}f + T \left( \frac{\partial p_c^f}{\partial T} \right)_p \left[ \frac{f}{2} + \frac{h_c^f - h_c^s}{2T} \right] + T \left( \frac{\partial p_c^s}{\partial T} \right)_p \left[ \frac{f}{2} + \frac{h_c^s - h_w^s}{2T} \right] - \frac{2 - p_c^f - p_c^s}{2} T \left( \frac{\partial f}{\partial T} \right)_p \tag{59}
\]

C. Discussion

The bottom line of the simple theory described in the previous subsection is that there are two important ranges of temperature, first around \( T \approx 0.5 \) where clusters of large particles begin to form, and a second around \( T \approx 0.1 \) where clusters of small particles begin to appear. The first important change is also seen in the bulk modulus; this is not surprising, since the crystalline clusters have a bulk modulus very different from the fluid. Nevertheless between the clusters we still have appreciable fluid regions which act as lubricants for the response to shear. We thus expect the shear modulus to change appreciably only when the small particles begin to cluster, in the vicinity of the smaller specific heat peak. We therefore conjecture that the two specific heat peaks are also associated with changes in the bulk and shear modulus respectively. We expect that any measurements of the glass properties connected with bulk and shear moduli will show different transition temperatures if these quantities do not reach simultaneously their \( K^\infty \) counterparts.

We cannot at this point assess how general is this split between bulk and shear moduli, and whether it will be seen in generic glasses. We thus leave this point for further research, stressing that we expect this phenomenon to appear whenever there exist micro-clusters of preferred ordering in the scenario of glass-forming.

Having an effective equation of state, albeit approximate, we can easily compute any thermodynamic derivative of interest. We wrote above explicit expressions for the compressibility and the thermal expansion coefficient. Others are as easily calculated. The point to stress however is how non-universal the thermodynamics is. In this model we have two specific heat peaks, in others we might have one or several. We would also expect a strong pressure dependence for these peaks.

IV. FREQUENCY-DEPENDENT SPECIFIC HEAT

By applying time dependent heat fluxes \( \delta Q(t) \) to the liquid and measuring the resulting temperature fluctuations \( \delta T(t) \), the specific heat can be measured \( \delta Q(t) \rightarrow C S \delta T(t) \). As mentioned in the introduction, measurements of the specific heat of glassy fluids at low temperatures can in principle be made under conditions of either constant volume (isochoric conditions) or constant pressure (isobaric conditions), but experimentally isobaric conditions are the norm.

The first and best known measurement of the fre-
imhomogeneities will also contribute to anomalies in the
caloric equation of state supplied by Eqs. (56)-(59). In
agreement with the simulations and the theory based on the
mechanical equation of state (Cf. Fig. 5) the theory predicts
the two peaks to both \( C_s \) and \( C_v \) which are associated with
the micro-melting or micro-freezing of the clusters of large
and small particles respectively. Note that \( C_p \geq C_v \) as
is expected from thermodynamics.

The reader is referred to [11] for an introduction to the
theory on an isobaric ensemble where the appropriate
Boltzmann distribution is \( \exp(-\beta H)/Z \), with the en-
thalpy given by \( H = E + PV \). Nielsen and Dyre [27]
find that the frequency dependent specific heat is given
by the form

\[
C_p(T, \omega) = \frac{\langle \Delta H^2 \rangle}{k_B T^2} + \frac{i \omega}{k_B T^2} \int_0^\infty \langle \Delta H(t) \Delta H(0) \rangle e^{+i\omega t} dt.
\]

(61)

In Eq. (61) \( \Delta H(t) = H(t) - \langle H \rangle \) is an enthalpy fluctua-
tion away from equilibrium. Therefore we write

\[
H(t) = \sum_s N_s(t) h_s + M(t) h_l
\]

(62)

where \( N_s(t) \) is the number of clusters consisting of \( s \)
molecules in the glassy phase and \( M(t) \) are the remain-
ing molecules in the mobile liquid phase. \( h_s \) is the en-
thalpy per molecule in the mobile phase.

Now in equilibrium we can write

\[
\langle H \rangle = \sum_s \langle N_s \rangle h_s + \langle M \rangle h_l
\]

(64)

and we also have the sum rule

\[
\sum_s s N_s(t) + M(t) = M
\]

(65)

where \( M \) is the total number of molecule in the system.

We can write the enthalpy fluctuations away from equi-
librium at time \( t \) as

\[
\Delta H(t) = H(t) - \langle H \rangle = \sum_s (N_s(t) - \langle N_s \rangle) h_s = \sum_s \Delta N_s h_s
\]

(66)

where

\[
h_s = H_s - \langle h_s \rangle = (\epsilon_c - \epsilon_l) s + P(v_c - v_l) s + \sigma s^{2/3}.
\]

(67)

Let us first calculate the equilibrium fluctuations \( \langle (\Delta H)^2 \rangle \). To this end we assume that there are no cor-
relations between the dynamics of clusters of different
sizes, implying \( \langle \Delta N_s \Delta N'_s \rangle = 0 \). Then, using the ex-
pression Eq. (66) for the enthalpy fluctuations, we can
immediately write that

\[
\langle (\Delta H)^2 \rangle = \sum_s \langle (\Delta N_s)^2 \rangle h_s^2.
\]

(68)

Similarly for the time dependent enthalpy fluctuations

\[
\langle \Delta H(t) \Delta H(0) \rangle = \sum_s \langle \Delta N_s(t) \Delta N_s(0) \rangle h_s^2.
\]

(69)
We have thus reduced the correlation functions for the enthalpy fluctuations into expressions involving the fluctuations in cluster number for clusters of different sizes $s$.

To estimate these correlation functions we proceed as follows. First we note that the number of molecules $N_c(t)$ in the clusters can be written as a sum over the clusters as $N_c(t) = \sum_s N_s(t)s$, and consequently the fluctuations in the total number of particles away from equilibrium are $\Delta N_c(t) = \sum_s \Delta N_s(t)s$. Then, assuming Gaussian fluctuations we estimate the mean square fluctuations of the number of particles within some small volume

$$\langle (\Delta N_c)^2 \rangle \approx \langle N_c \rangle$$

or rewriting

$$\sum_s \langle (\Delta N_s)^2 \rangle s^2 = \sum_s \langle N_s \rangle s.$$  

(71)

From this equation we therefore see that

$$\langle (\Delta N_s)^2 \rangle = \langle N_s \rangle / s.$$  

(72)

For the time dependent fluctuations therefore, assuming an independent Debye relaxation for each cluster,

$$\langle \Delta N_s(t) \Delta N_s(0) \rangle = \langle N_s \rangle e^{-t/\tau_s} / s$$

(73)

where $\tau_s$ is the lifetime of a cluster of size $s$. Thus we get our final expression for the enthalpy fluctuations in equilibrium in terms of the cluster size distribution as

$$\langle (\Delta H)^2 \rangle = \sum_s \langle N_s \rangle (h_s^2 / s).$$

(74)

and for their time dependent correlations

$$\langle \Delta H(t) \Delta H(0) \rangle = \sum_s \langle N_s \rangle e^{-t/\tau_s} (h_s^2 / s).$$

(75)

We now substitute these expressions into Eq. 61 with the result

$$C_p(T, \omega) = \frac{1}{k_B T^2} \sum_s \langle N_s \rangle (h_s^2 / s) \frac{1}{1 - i\omega \tau_s}$$

(76)

or splitting the specific heat into its real and imaginary parts

$$\Re C_p(T, \omega) = \frac{1}{k_B T^2} \sum_s \langle N_s \rangle (h_s^2 / s) \frac{1}{1 + (\omega \tau_s)^2}$$

$$\Im C_p(T, \omega) = \frac{1}{k_B T^2} \sum_s \langle N_s \rangle (h_s^2 / s) \omega \tau_s \frac{1}{1 + (\omega \tau_s)^2}$$

(77)

We can now use our previous results for the cluster distributions in the case of glycerol to find the real and imaginary specific heat anomalies in the case of glycerol. We do not re-fit any of the parameters used in the calculation of the BDS spectra [7], but we simply use the previous knowledge at the temperatures indicated, and plot the results, fitting only the heat conductivity of glycerol. We approximate $h_s^2 / s \approx (h_c - h_l)^2 / s$, such that Eq (76) is rewritten as

$$C_p(T, \omega) \approx \frac{(h_c - h_l)^2}{k_B T^2} \sum_s \langle N_s \rangle s \frac{1}{1 - i\omega \tau_s}.$$  

(78)

Splitting the specific heat into its real and imaginary parts

$$\Re C_p(T, \omega) \approx \frac{(h_c - h_l)^2}{k_B T^2} \sum_s \langle N_s \rangle s \frac{1}{1 + (\omega \tau_s)^2},$$

$$\Im C_p(T, \omega) \approx \frac{(h_c - h_l)^2}{k_B T^2} \sum_s \langle N_s \rangle s (\omega \tau_s) \frac{1}{1 + (\omega \tau_s)^2}.$$  

(79)

(80)

The resulting curves multiplied by the thermal conductivity are shown in Figs. 7 and 8. These should be compared to Fig.2 of [3]. The reader can convince himself that the theory captures the experimental results quantitatively.

We would like to stress at this point that the results obtained here are equivalent in dynamical contents to the computation of the dielectric spectra in [11]. In that calculation one focused on the dielectric response $\epsilon(\omega)$ and here decomposed it into its real and imaginary parts $\epsilon(\omega) = \Re \epsilon(\omega) + i \Im \epsilon(\omega)$. It was found in [11] that without the dc contribution (which is absent in the case of specific heat) we could write

$$\Re \epsilon(\omega) - \epsilon_{\infty} = \frac{1}{(\epsilon_0 - \epsilon_{\infty})} \sum_s \langle N_s \rangle s / [1 + (\omega \tau_s)^2],$$

$$\Im \epsilon(\omega) = \frac{1}{(\epsilon_0 - \epsilon_{\infty})} \sum_s \langle N_s \rangle s (\omega \tau_s) / [1 + (\omega \tau_s)^2].$$

(81)

Once normalized the specific heat spectra are identical these spectra. The reason for the identity is in the assumptions that $\langle m_m \cdot m_m \rangle \sim s$ and $\langle h_p^2 \rangle \sim s^2$, cf. Eq. (67). On the other hand the role of the relaxation times $\tau_s$ and the distribution of cluster sizes are exactly the same in the two expressions.
other systems, or may be an even richer scenario can
two specific heat peaks discussed above may be seen in
would be very useful for the near future.
exactly the same temperature, but such an experiment
the spectra measured from specific heat can be divided
sequence of our model for glycerol is the prediction that
methods of linear response like Broad Dielectric Spec-
revealed by the complex specific heat compared to other
the context of the example of glycerol, that the physics
the complex specific heat we have shown, in
higher temperatures when the first type of clusters form,
ranges for the increase in moduli, the bulk modulus at
in the case of the binary mixture indicates two different
for the increase in moduli, the bulk modulus at
shear modulus when the other type of clusters
Higher temperatures when the first type of clusters form,
pronounced that the appearance of two specific heat peaks
the potential energy (4) of the binary mixture model (2)
is given by:
Due to the scaling properties of the inverse power poten-
tial it is possible to normalize the interparticle distances
by the typical distance (11) \( r_{ij} \rightarrow s_{ij} = r_{ij}/l \) [29] ;
where the constant \( c_M = (1/2N) \sum_{ij} 1/s_{ij}^n \) is independent of the density. Note that this constant is known as
the Madelung constant in solid-state physics. Taking into
account (A4) the average potential energy of a harmonic
solid (A2) can be rewritten as:

The value of the constant \( c_M \) can be calculated simu-
lationally using Eq. (A5); the results are shown in Fig.9.

V. SUMMARY AND DISCUSSION

Probably the most glaring consequence of the calculations presented in this paper is that the specific heat is a valuable indicator of the interesting physics that occurs during the glass transition, but this transition is in no way universal. The temperature dependence of the specific heat is determined by details like inter-particle potentials and micro-melting or micro-formation of clusters. In this sense any hope for universality is untenable. Nevertheless we have shown that the specific heat peaks herald interesting new physics, leading to fast changes in the mechanical moduli which are also associated with fast changes in the inhomogeneities that are crucial for the glassy behavior, i.e. the formation of micro-clusters. We propose that the appearance of two specific heat peaks in the case of the binary mixture indicates two different ranges for the increase in moduli, the bulk modulus at

It is interesting to see in future research whether the two specific heat peaks discussed above may be seen in other systems, or may be an even richer scenario can appear, with more peaks, when more types of clusters intervene in the process of glass-formation.

APPENDIX A: THE ZERO TEMPERATURE LIMIT

It follows from the simulation results that at lower temperatures the specific heat is at least close to the value of the solid. Therefore, we can write the energy of the system in the harmonic approximation:

\[
U = U_0 + \sum_{i,j} a_{ij} q_i q_j,
\]

where \( U_0 \) is the potential energy of the system in the reference state, \( a_{ij} \) are expansion coefficients and \( q_i \) is a Cartesian coordinate of the deviation of the current position of a particle from equilibrium. There are \( dN \) degrees of freedom (neglecting translations and rotations of the system) in (A1), therefore it follows from the equipartition theorem that the average potential energy of the system in the solid state is [28]:

\[
\langle U \rangle / N = U_0 / N + dT / 2.
\]

Substituting (A2) in (3) immediately yields \( C_V = 2 \) for two dimensional solids. At the equilibrium configuration the potential energy (4) of the binary mixture model (2) is given by:

\[
U_0 = \frac{1}{2} \sum_{i,j,l} (\sigma_{ab} / r_{ij})^n
\]

Due to the scaling properties of the inverse power potential it is possible to normalize the interparticle distances by the typical distance (11) \( r_{ij} \rightarrow s_{ij} = r_{ij}/l \) [29];

\[
\langle U \rangle / N = c_M \rho \phi + \frac{d}{2} T.
\]

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One can see that below $T = 0.5$ the change of this constant is small. Nevertheless, this change reflects the fact that our calculations at the smallest values of the temperature are not fully relaxed to equilibrium even though we took extreme care. Typically at the lowest temperatures the system can be trapped for incredibly long times in a local minimum of the energy surface, where each local minimum having slightly different equations of state [16]. While we expect the Madelung constant to be unique for a given crystal, our system here contains clusters of preferred structures with random orientations [30], and therefore the analog of the Madelung constant is not strictly defined. It may very well depend on the protocol of cooling. The present best estimate of the value of this parameter at the lowest temperatures is $c_M = 14.649$.

The caloric equation of state (A5) substituded to the virial equation (7) gives the following thermal equation:

$$
\frac{P}{\rho T} = (1 + \frac{n}{2}) + \frac{n}{d}(c_M/\sigma_c)\Gamma e^{n/d}. 
$$

The value of the renormalized constant $c_M/\sigma_c = 1.394$ can be compared with the result for the two dimensional one-component system with hexagonal crystal, which is 1.268 [31]. The fact that this constant is expected to increase in an amorphous solid was anticipated in [29].

Finally, we note from (A6) that in contrast to a crystalline solid the thermal (caloric) equation of state here remains ambiguous because the value of $c_M$ depends on the preparation protocol. With this in mind it becomes fruitless to seek the anharmonic corrections to equation (A6) as in the case of a one component system with a well-defined reference state at low temperatures. Nevertheless we stress that the specific heat at constant volume does not suffer from any ambiguity and therefore can be taken good as a good indicator of the solidification.

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