Analysis of a Lennard-Jones fcc structure melting to the corresponding frozen liquid: differences between the bulk and the surface

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Abstract

We computed a Lennard Jones frozen liquid with a free surface using classical molecular dynamics. The structure factor curves on the free surface of this sample were calculated for different depths knowing that we have periodic boundary conditions on the other parts of the sample. The resulting structure factor curves show an horizontal shift of their first peak depending on how deep in the sample the curves are computed. We analyze our resulting curves in the light of spatial correlation functions during melting. The conclusion is that the differences between bulk and surface are quite small during melting and that at the end of melting, only the very surface happens to be less dense than the bulk. This result is intrinsic to the shape of the Lennard Jones potential and does not depend on any other parameter.

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I. INTRODUCTION

The aim of this study is to analyze the structure of the surface of a Lennard Jones frozen liquid (within a depth of a few tens of Lennard Jones units). Our Lennard Jones liquid may be compared to a model of glass for which the glass transition would have been obtained instantaneously from the liquid.

Different studies deal with the analysis of glass surface. They mainly concern the chemical nature of the surface. In the case of silicate and silica glasses, there are different numerical models, see for that the following references [1, 2, 3, 4, 5]; these previous references studied the structure, attack of water and the presence of free oxygen atoms on the surface of silicate glasses (with dangling bonds, as there are lacking hydrogens) as well as the variation of composition at the surface.

But very few numerical study of the surface of simple models of liquids or glasses have been made up to now [6]. Stallons and Iglesia [6] studied the free surface of a simple model of silica glass using the Stillinger Weber potential and found that the surface of their glasses was less dense than the bulk.

We deal here with the surface of a Lennard Jones frozen liquid. Here the action of the atmosphere and the presence or not of dangling bonds have not been studied. We are interested in the change in the structure of the free surface of a simple model of frozen liquid (Lennard Jones liquid) without any external action, only the fact that there are no periodic boundary conditions in one direction: the free surface. After obtaining the Lennard Jones structure with a free surface by classical molecular dynamics computation, we analyzed the free surface structure with the calculations of structure factors corresponding to different depths within the sample. This analysis is completed by the computation of the density as a function of depth within the sample and by computation during melting, of the spatial correlation function in the bulk and at the surface.

The computational method used for the sample preparation as for the calculation of the structure factor curves are presented in section II. Section III shows the results which are discussed. And finally, section IV is the conclusion.
II. NUMERICAL PROCEDURE

A. Preparation of the numerical sample

We used the classical molecular dynamics to compute a Lennard Jones liquid. The potential was a Lennard Jones 6:12 potential cut at a distance 2.5 Lennard Jones units. We began the simulation with a fcc structure which was melt in \(2.5 \times 10^{13}\) s (250 Molecular Dynamics-MD- steps). This is the time needed to obtain total melting (i.e. no further melting). The computation was done out of equilibrium: i.e. due to the free surface we could not impose a NVT or NPT equilibrium during melting. We chose an instantaneous cooling (in 1 MD step) in order to prevent the sample from shearing phenomena which would rearrange the structure even at its surface. Thus our sample is just a frozen Lennard Jones liquid. The sample has periodic boundary conditions in all directions except on two free surfaces on the bottom and on the top of it. As the aim of this study is the analysis of the free surface of Lennard Jones frozen liquids, we had to add these two free surfaces surrounded by vacuum. The initial box (not taking into account the boundary conditions) has a dimension of 50X50X50 Lennard Jones (LJ) units corresponding to 320000 LJ particles.

In order to analyze only the action of the free surface on the corresponding structure factors, we had to cut the initial box containing the LJ frozen liquid into two, according to its depth. Unless, there would be action of the two free surfaces on the bottom and on the top of the simulated LJ frozen liquid, and the resulting structure factor curves would be to difficult to analyze. Furthermore it would not be a physical model of the surface of frozen liquids. The fact that we have a free surface before having melt and frozen the sample allows us to obtain a free relaxed surface on the top of the bulk of the Lennard Jones frozen liquid.

B. Calculation of the structure factor

We used a similar calculation as in reference [7] to obtain the structure factor. The difference is that the domain of computation is limited by the upper surface surrounded by vacuum and a given depth within the numerical sample. Indeed, the wave vector may be written:

\[
q = k_f - k_i \tag{1}
\]
The calculation of the structure factor curves is the following:

\[ S(q) = \frac{\epsilon^6}{V} \sum_r |\delta(r) - c| \exp{i \mathbf{q} \cdot \mathbf{r}}^2 \]  

(2)

where \( \epsilon^6 \) comes from the replacement of the integral by discrete sum over the LJ particles, \( V \) is the scattering volume (in order to normalize all different results depending on the analyzed depth). \( c \) is the concentration of the sample which we have calculated as function of the depth of the sample. This last feature ensures that \( S(q) \) tends to zero when \( q \) tends to zero.

Then we introduce a double sum to expand the square, and, assuming isotropy in the plane (but not in the depth of the sample), we average over all directions of \( q \) to get:

\[ S(q) = \frac{\epsilon^3}{N_c} \sum_{r_1} \sum_{r_2} (\delta(r_1)\delta(r_2) - c[\delta(r_1) + \delta(r_2)]) + c^2)N \sin \frac{qr}{qr} \]  

(3)

where

\[ r = |r_1 - r_2| \]  

(4)

Then, transforming the double sum into a sum over all possible values for the distance \( r \) and separating the \( r = 0 \) contribution from the others, we obtain:

\[ S(q) = \epsilon^3(1 - c + \sum_{r \neq 0} [F_a(r) - cF_b(r)])\sin \frac{qr}{qr} \]  

(5)

where \( F_a(r) \) is the mean number of LJ particles at a distance \( r \) of a given LJ particle (pair correlation function) and where \( F_b(r) \) is the same quantity for a density equal to 1.

III. RESULTS

Figure 1 exhibits the typical structure factor for different depths in LJ units. At this scale it is not possible to make any difference between the curves. Anyway, let us remark that the curves are typical of a glassy material with several peaks. The first peak located at approximately at \( q = 0.34 \) i.e. at a distance equal approximately to 3 LJ units. This result shows that there is, in our Lennard Jones frozen liquid, a short range order up to 3 LJ units. However, the intensities of the structure factors have been normalized by the analyzed volume (see section II.B for details concerning this normalization). This last feature explains that we have no vertical shift for different structure factors.
Figure 2 presents a zoom of the previous curves (figure 1) for $0.3 < q < 0.39$ LJ units$^{-1}$. One may see that there is a difference for the curve corresponding to a depth of 5 LJ units compared to the others.

Figure 3 is also a zoom of the curves presented in figure 1 but for a different location than figure 2, here $0.24 < q < 0.32$ LJ units$^{-1}$. One may see again that the curve corresponding to the depth equal to 5 LJ units is different from the others.

Finally, and in order to be able to understand our computed structure factor curves we plotted the number of LJ particles as a function of the depth from the free surface of the sample (see figure 4). The numerical simulation box begins at height $z = 1$ LJ units but as explained in section II, we cut the sample into two, so the bottom of the sample corresponds to $z = 25$ LJ units and the free surface of the sample to $z = 50.1$ LJ units.

Figure 5 exhibits the distribution of the LJ particles whose center of mass is contained in the slice for $49.5 < z < 50.1$. This may be compared to figure 6 which is the distribution of the centers of mass of the LJ particles for $20.5 < z < 21.1$, i.e. for the same width following $z$. The fact that there are zones without particles in figure 6 is explained by the width of the analyzed slice: it is less than 1 LJ unit. In figure 5, the void zones are much larger, as expected from the variation of the number of particles presented in figure 4. What can be observed is that distribution is rather homogeneous. In order to go further in this analysis, we computed the two points correlation function:

$$g(r) = \frac{N(r)}{n^2}$$

where $n$ is the number of particles in the slice and $N(r)$ is the number of particles at distance $r$ from another particle. We computed the evolution of this two points correlation function as a function of computation times of the sample structure, during melting. We chose a depth of this slice equal to 0.3 because it corresponds to the depth where the surface has a lower density. Although the depth of the slice is only 0.3 LJ particles, we chose the slices at the surface and in the bulk so that we get half of a fcc elementary cell in the slice at the beginning of computation. Figures 7-a (left and right) are the correlation functions at the beginning of computation (for the fcc structure) at the very surface and in the bulk. One may see a peaked distribution function. This is due to the crystalline structure of the fcc initial distribution. Fig.7-b,7-c and 7d correspond respectively to 100 MD time steps, 200 MD time steps and 250 MD time steps (corresponding to the frozen liquid). We can observe
that the right and left figures 7 are very similar and we will discuss the small differences between bulk and surface later in the discussion. The correlation function (for times t=100 MD and 200 MD time steps) becomes similar to a random packing except that there are one peak at a distance equal to 1 LJ units. This means that there is short range correlation during melting and that aggregates most likely remain during melting. This peak almost disappears when the liquid is frozen for 250 MD time steps. Fig.7d shows that there is no more a short range correlation and that the LJ particles are more equally distributed over the slice.

And finally, figure 8 is the plot of the number of particles per LJ unit volume for a slice of depth 0.3 LJ units at the very surface and in the bulk.

IV. DISCUSSION

We observed that the density of our LJ frozen liquid near the surface is lower only at the very surface for a depth included between $49.8 < z < 50.1$ LJ units. There is an increasing density for $z < 49.5$ LJ units and this is much less than the correlation length of about 3 LJ units (see above). Therefore the mean density for a depth of 5 LJ units is lower than for larger depths for which the low density of the free surface is negligible by averaging the density over the volume. This explains that we have no differences between the structure factors for lower depths, i.e. $z = 10, 15, 20$ and 25 LJ units and very small differences for $z = 5$ LJ units. We cut the sample in the $x$ direction and checked that the density did not vary at $x = 0$ and at $x = 50$ compared to the bulk. To better understand the origin of these small differences it is interesting to compare the evolution of the density at the very surface and in the bulk during melting.

The evolution of the density within the bulk and at the very surface, represented in figure 8, shows that there is not a large difference between the two curves. This means that the LJ particles belonging to the surface have sufficient connections with other particles in the bulk and within the surface to prevent them to lower the density at the very surface and to prevent them from evaporation. One can observe that in the bulk, the density decreases during the first MD computation steps and remains constant for all others computation times. On the other hand, there is another lowering of density of the upper surface during the last freezing step.
Considering now the two points correlation function, and comparing the left curves of figure 7 (at the very surface) and right curves (in the bulk), we can observe a few differences. First in figure 7b, there are regular oscillations for 100MD time steps in the figure corresponding to the bulk, indicating that correlations between particles in the bulk remains on a larger time scale than at the very surface; these last disappear for 200 MD time. In figure 7d, one can observe that at the end of melting, there are more particles at shorter distances in the bulk than at the very surface. This is well evidenced in figures 5 and 6.

Using data of figures 5 and 6, we computed the mean distance between the particles eliminating the void zones (i.e., distances larger than 1.5). We obtained almost the same mean distance for the bulk and for the surface of about 1.1 LJ unit. The particles stop to move during melting when they reach a mean distance between particle of 1.1 LJ unit which is the maximum of the attractive part of the LJ potential (if we eliminate the void zones). This means that the arrangement of particles are almost the same than in the bulk, but that there is a kind of “roughness” at the very surface due to large void zones.

We can say that the free surface of a LJ frozen liquid has also surface defects (i.e., the surface is less dense as shown in figure 4 and as deduced from figures 2 and 3). There are some larger “voids” at the very surface and a few particles rising above the mean surface. But these effects concern only the region very next to the surface (for a depth lower than 1 LJ unit). The effect of the free surface induce changes in the structure only for the first LJ particles layer. Moreover, the bulk conserves aggregates during melting while the surface loses this order more rapidly (in figure 7, the peak at abscissa 1 is smaller for the surface for 100MD and 200MD than for the bulk).

As a conclusion, the shape of the LJ potential leads to a surface where the density is of course lower because of contact with vacuum. The surface has a structure of homogeneously distributed particles which interaction is the largest attractive possible with the LJ potential surrounded by void zones: the surface is rough.

V. CONCLUSION

We computed the structure factors as a function of the depth near the surface of a Lennard Jones frozen liquid. The results are that very close to the surface the density of the numerical frozen liquid is lower. This is compared with the analysis of the density as a
function of depth into the numerical frozen liquid and explains the shift toward lower $q$ of the structure factor curves for lower depths. Our Lennard Jones frozen liquid is less dense near its surface which means that the set up of the sample itself, i.e. with a free surface leads to a rearrangement of the LJ particles without any external action (like the attack of water on real glasses). The interesting conclusion of our work is that even in a Lennard Jones frozen liquid (i.e. without any action of water: presence of atmosphere or without any presence of dangling bonds) the density and structure modification induced by the free surface only concern a very small depth of the sample (smaller than 1 LJ unit). During melting in the LJ fcc then LJ liquid, the LJ particle rearrange to become very rapidly a quasi two dimensional amorphous material on the surface.
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FIG. 1: The structure factor curves for different depths: at this scale it is not possible to make a difference between the different curves.
FIG. 2: Zoom of the different structure factor curves for different depths and for $0.3 < q < 0.39$ LJ unit$^{-1}$. 
FIG. 3: Zoom of the different structure factor curves for different depths and for $0.24 < q < 0.32$ LJ units$^{-1}$. 
FIG. 4: Number of Lennard Jones particles as a function of the height of the sample. Left figure: vertical coordinates from 25 to 50.1 LJ units which is the surface of the sample, right figure: zoom on the number of Lennard Jones particles number close to the surface for a height equal to 50.1 LJ units.
FIG. 5: Top view of the centers of mass for the free surface for $49.5 < z < 50.1$
FIG. 6: Top view of the centers of mass of the LJ particles for the bulk for $20.5 < z < 21.1$
FIG. 7: Two points correlation function for a slice at the free surface (left) and in the bulk (right) within a depth of 0.3 LJ units. (a): for the fcc structure at the beginning of computation (0 MD time step); (b): for 100 MD time steps; (c): for 200 MD time steps; (c) for 250 MD time steps when the liquid is frozen
FIG. 8: Number of LJ particles per LJ unit volume in the bulk and at the very surface both for a depth of 0.3 LJ units