Exposing nanobubble-like objects to a degassed environment

Robin P. Berkelaar, Erik Dietrich, Gerard A. M. Kip, E. Stefan Kooij, Harold J. W. Zandvliet and Detlef Lohse

The primary attribute of interest of surface nanobubbles is their unusual stability and a number of theories trying to explain this have been put forward. Interestingly, the dissolution of nanobubbles is a topic that did not receive a lot of attention yet. In this work we applied two different experimental procedures which should cause gaseous nanobubbles to completely dissolve. In our experiments we nucleated nanobubble-like objects by putting a drop of water on HOPG using a plastic syringe and a disposable needle. In method A, the nanobubble-like objects were exposed to a flow of degassed water (1.17 mg l$^{-1}$) for 96 hours. In method B, the ambient pressure was lowered in order to degas the liquid and the nanobubble-like objects. Interestingly, the nanobubble-like objects remained stable after exposure to both methods. After thorough investigation of the procedures and materials used during our experiments, we found that the nanobubble-like objects were induced by the use of disposable needles in which PDMS contaminated the water. It is very important for the nanobubble community to be aware of the fact that, although features look and behave like nanobubbles, in some cases they might in fact be induced by contamination. The presence of contamination could also resolve some inconsistencies found in the nanobubble literature.

1 Introduction

Sub-micron bubbles on hydrophobic interfaces were thought to be the cause of the long-range hydrophobic interaction and this resulted in the emergence of a completely new field, that of surface nanobubbles. The seminal work was performed by Parker and Attard in 1994, who observed long-range attractive forces using a surface force apparatus and attributed this to the existence of nano-scale gas bubbles at the interface. Actual images of nanobubbles had to wait until advancements in atomic force microscopy (AFM) immersed in liquids resulted in the observation of soft spherical cap shaped features by Lou et al. and Ishida et al. in 2000. Unfortunately, the AFM tip disturbs these soft features and properly imaging nanobubbles is not a trivial task. Nanobubbles have been observed on a wide variety of surfaces and found to be stable under a broad range of conditions like elevated temperature, low pH and salt solutions. The first convincing proof for the gaseous nature of these features came from Zhang et al., showing gas-enrichment near the interface using ATR-IR measurements in 2007. During this period, the gaseous nature of these bubbles was also indirectly inferred by degassing the liquids used to nucleate nanobubbles and by degassing the nanobubble covered substrate. Also the growth of nanobubbles by rectified diffusion using an acoustic field suggests their gaseous nature. Although an abundance of experiments have been performed on these nanobubbles since 1994, mixed results on a number of topics were found. For example, the contact angle depends in some studies on the radius of curvature whereas in other experiments the contact angle is found to be constant. The presence of a gas layer at the solid–liquid interface is observed in several experimental studies where in other studies such a gaseous phase is not found. Also, nanobubbles are sometimes found in ethanol while others observe pristine surfaces when immersed in ethanol. And finally, nanobubbles are in some cases imaged on HOPG just by immersing the substrate in water while others need to perform an ethanol–water exchange to induce nanobubble nucleation.

Despite the inconsistencies, what the nanobubbles all have in common is their long term stability. The fact that these bubbles can be observed is quite a remarkable feat on its own, they have been measured to be stable for as long as several days. For small bubbles the Laplace pressure dominates, and this drives the dissolution of gas from the bubble into the environment.
liquid. Bubbles with a radius of curvature $R_c$ less than 1 μm should thus dissolve on a timescale of $\tau \sim R_c^2/D$, where $D = 1 \times 10^{-9}$ m$^2$s$^{-1}$, i.e. in microseconds.\textsuperscript{38,39} The existence of stable bubbles with radii of a few hundred nanometers and heights in the order of ten nanometers, hence the name nanobubbles, sparked the interest to what the mechanism behind this remarkable stability could be.

Since the discovery of nanobubbles a number of theories explaining this surprising behavior have been proposed. Just after the discovery of nanobubbles it was argued that they might not be bubbles, but contamination, e.g. resulting from polymers used to hydrophobize the surface (theory 1).\textsuperscript{40} However, this was soon to be discarded by the assertion that the bubble contained gas. A new theory followed, in which the presence of contamination at the bubble gas–liquid interface lowered the surface tension, and thus lowered the Laplace pressure, which in effect reduced the dissolution of the bubble (theory 2).\textsuperscript{41} In addition, calculations of the contamination concentration needed for a sufficiently low surface tension to match the measured contact angle for nanobubbles resulted in a layer thickness which greatly hinders the gas out-flux.\textsuperscript{42} Also the calculations from Das et al.\textsuperscript{41} suggest that a possible contamination lowers the surface tension and the gas-flux through the interface, but this was insufficient to stabilize the bubble. Experiments using a surfactant to remove a hypothetical contamination layer by Zhang et al.\textsuperscript{19} showed that nanobubbles remain stable and do not dissolve when exposed to the SDS surfactant (which should wash away contaminations), a result confirmed by Peng et al.\textsuperscript{43} As these authors showed, the detergents do help to mechanically remove surface nanobubbles with the AFM tip. Both used a surfactant concentration below the critical micelle concentration (CMC). In other studies a concentration above the CMC was used; Ducker\textsuperscript{44} showed the dissolution of nanobubbles in this particular case. However, in a more recent study Zhang et al.\textsuperscript{44} observed again stable nanobubbles even for surfactant concentrations above the CMC.

As the stability could not be explained by contamination, there was a need for a new and completely different approach, which resulted in the dynamic equilibrium theory by Brenner and Lohse\textsuperscript{18} (theory 3). The main idea of this theory is that the gas out-flux of the bubble is compensated by a gas in-flux at the three-phase contact line. This theory was later extended and specified by Seddon et al.,\textsuperscript{46} the gas inside a nanobubble fulfills the requirements for a Knudsen gas, meaning that the mean free path of the gas molecules is larger than the distance to the interface of the bubble. Therefore, gas-molecules desorbing from the gas–solid interface will hit the gas–liquid interface and transfer momentum along a preferred direction perpendicular to the solid–liquid interface. This then drives a circulatory flow around the nanobubble transporting a stream of gas rich water to the three-phase line of the nanobubble, where the gas absorbs onto the surface and diffuses back into the nanobubble. Using alternate formulations of this theory it made possible to predict the temperature and gas saturation dependency of nanobubbles.\textsuperscript{47} What, however, remains unclear in this theory is what energetically drives the flow and therefore a non-equilibrium situation has to be assumed.

Recently another theory was proposed by Weijts and Lohse\textsuperscript{48} (theory 4), which does not suffer from the difficulty that the dynamic equilibrium theory has. The theory combines the assumption, which is in some cases observed experimentally\textsuperscript{47} that the contact line of a nanobubble is pinned together with the retardation of gas diffusion in a liquid compared to air. The moment a small amount of gas leaves the nanobubble, the contact angle will have to decrease in order to accommodate the reduction in volume, which in return lowers the Laplace pressure and hence slows down the dissolution of the nanobubble. The gas molecules, which just dissolved from the bubble into the liquid, increase the gas saturation around the nanobubble and will take a significant time to diffuse towards the interface of the water film and leave the system. The increased gas saturation around the nanobubble, resulting from these gas molecules and those from neighboring nanobubbles, lowers the out-flux of new gas molecules from the nanobubble and thus enhances the stabilization. The combined effect of contact-line pinning and diffusion retardation in liquids results in considerably longer lifetimes, dependent on the liquid-film thickness $\tau \sim \ell^2/D$, where $\ell$ is the liquid film thickness and $D$ the diffusion coefficient of gas in liquid.

The number of experimental studies that focuses on verifying or disproving one or more of the above theories is rather limited and the results from these studies are often inconsistent. A recent experiment showed that nanobubbles were stable in degassed water which was refreshed every 20 minutes, and the authors concluded that this was most likely due to contamination.\textsuperscript{31} However, Zhang et al.\textsuperscript{25} have shown the localized disappearance of nanobubbles after degassing, though some regions remain covered with nanobubbles after degassing.

The scope of the present study is to try to contribute to a clarification of the puzzling situation. In the way to produce nanobubbles or nanobubble-like objects, we will restrict us to the case of liquid deposition on hydrophobic flat surfaces. We will not address the most popular method for nanobubble formation, namely ethanol–water exchange, or more generally, solvent-exchange. We investigate whether the nanobubble-like objects “communicate” with the surrounding liquid by the diffusion of gas molecules using two different methods. In method A, the nanobubble-like objects are exposed to a continuous flow of degassed water and in method B the ambient pressure is reduced. This should unambiguously result in a significant reduction of the lifetime, if nanobubbles are stabilized by theory 3, theory 4 or any other mechanism where the gas can diffuse through the gas–liquid interface of the bubble. In both cases the nanobubble-like objects, which develop at deposition, remain stable after prolonged exposure to degassed water. We conclude that therefore they are either actually not gaseous or have a gas impermeable shell, which could be in accordance with theory 1 or 2, i.e. due to contamination, or the result of an, so far, unknown physical mechanism.

2 Experimental details

Nucleation of nanobubble-like objects was acquired by immersing an HOPG (ZYA grade, MikroMasch) substrate in
water. The substrate was freshly cleaved prior to each experiment and subsequently clamped between two Teflon rods in an all Teflon liquid-cell. The liquid-cell was cleaned in a Piranha solution (a 3:1 \( \text{H}_2\text{SO}_4 \) to 30% \( \text{H}_2\text{O}_2 \) mixture) and rinsed with copious amounts of water. Purification of the water was performed using a Simplicity 185 system (Millipore) up to a resistivity of 18.2 MΩ cm. The liquid-cell was filled with 3–4 ml water using a new 5 ml sterile plastic disposable syringe (Discardit, BD) and a disposable needle (Microlance, BD). The Teflon liquid cell was then placed within an Agilent 5100 atomic force microscope. The AFM nose-cone was rinsed thoroughly with ethanol (EmSure ≥99.9% purity, Merck) and dried in a \( \text{N}_2 \) gas flow before imaging. The immersed HOPG surface covered with nanobubble-like objects was imaged by the AFM operated in intermittent contact mode. The liquid-cell was subsequently removed from the AFM and sealed with a SiO\(_2\) wafer, which intermittently adjusted close to 100% in order to limit the deformation of the objects by the tip. The liquid-cell was thereafter removed from the AFM and closed using a SiO\(_2\) substrate. Degassed water was then injected into the liquid cell with a continuous flow of 1.5 ml min\(^{-1}\). As a result, the measured \( \text{O}_2 \) saturation inside the liquid cell during degassed water flow was <13% (1.17 mg l\(^{-1}\)). The diffusion coefficients at 20 °C in water for the other major constituents of air (nitrogen \( \text{D} = 2.6 \times 10^{-9} \text{ m}^2 \text{ s}^{-1} \) and argon \( \text{D} = 2.3 \times 10^{-9} \text{ m}^2 \text{ s}^{-1} \)) are comparable to that of oxygen (\( \text{D} = 2.3 \times 10^{-9} \text{ m}^2 \text{ s}^{-1} \)). The measured \( \text{O}_2 \) saturation can thus be regarded as the absolute gas saturation of the water. Flowing degassed water has a number of advantages compared to other degassing techniques. Firstly, a continuously low gas-saturation can be guaranteed, even when the liquid-cell is not sealed properly. Secondly, the flow will cause convection and thus better mixing compared to statically filling the liquid cell with degassed water. Thirdly, there are no detrimental effects from macroscopic bubbles expanding and sweeping clean the area of interest, as can be the case for degassing by reducing the ambient pressure. After exposing the nanobubble-like objects to the degassed water flow for 96 hours the liquid-cell was placed back in the AFM, and the same area was imaged once more.

Surprisingly, the nanobubble-like features had not vanished, quite the opposite, they appear even larger, Fig. 1B. All effort was taken to exclude deformation of the objects by the tip and having similar scanning parameters for all images, such as amplitude (nm) and set-point. Still we have the impression that

### 3 Results & discussion

The stability of our nanobubble-like objects was first challenged using method A: degassed water was flowed over the objects for a prolonged time. A freshly cleaved HOPG surface was clamped into an all Teflon liquid cell and immersed in water. The liquid-cell was then mounted into the AFM, where the surface was scanned in intermittent-contact mode. This procedure resulted in a substantial coverage of surface nanobubble-like objects, as can be observed in Fig. 1A. The larger objects have an asymmetrical appearance, generally referred to as parachuting, due to the set-point being close to 100%. The set-point was intentionally close to 100% in order to limit the deformation of the objects by the tip. The liquid-cell was thereafter removed from the AFM and closed using a SiO\(_2\) substrate. Degassed water was then injected into the liquid cell with a continuous flow of 1.5 ml min\(^{-1}\). As a result, the measured \( \text{O}_2 \) saturation inside the liquid cell during degassed water flow was <13% (1.17 mg l\(^{-1}\)).

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a parameter has changed, such as liquid height or the difference in effective spring-constant between the cantilevers used in the two images (though freshly taken from the same box), and this increase does not represent an actual change in the objects’ height. In any case, we are not so much concerned about the actual geometry of the nanobubble-like objects, rather the fact that they are still present after prolonged exposure to a degassed environment is of interest. Comparing the images before and after degassed water flow results in virtually no change regarding the number and position of the nanobubble-like objects. The only viable explanation is the absence of mass transfer through the interface between the nanobubble-like objects and the water.

We also investigated the gas out-flux using the above mentioned method B, in which the nanobubble-like objects are exposed to a reduced pressure for a prolonged period of time. The liquid cell with the immersed HOPG sample was removed from the AFM and inserted into a glass pressure vessel. A few centimeters high water layer was present in the pressure vessel to prevent complete evaporation of the water in the liquid cell. The pressure was gradually dropped from atmospheric pressure to $\approx 20$ mbar in the course of 24 h. It is essential that the pressure drop is slow, in order to restrain the formation of macroscopic bubbles on the HOPG interface, since the moving contact line of a growing macroscopic bubble will efface the nanobubble-like objects from the surface. The pressure remained at a low pressure of $\approx 20$ mbar for 30 min before increasing it back to atmospheric conditions within 5 min. The liquid-cell was subsequently returned to the AFM for imaging. The results are similar to that of the degassed water flow experiment. Comparing the same area before and after degassing reveals that again the number and size of the nanobubble-like objects remain virtually unchanged, see Fig. 2. Some of the smallest objects do not appear on the image after degassing (an example is pointed out using arrows), Fig. 2B, which is presumably due to reduced resolution. This is in agreement with the results from the degassed water flow; there does not seem to be any mass-transfer between the alleged nanobubble-like objects and the liquid.

A bubble with a gas-impermeable shell should still expand in volume, which could result in changes in the nanobubble coverage as discussed in Appendix A. However, there we show that the changes in radius or contact angle are too small to result in any lasting modifications by coalescence in nanobubble coverage after the pressure is reduced and subsequently increased back to atmospheric conditions.

The results from the depressurizing experiment are in complete agreement with the degassed water flow experiment: in both cases the nanobubble-like objects do not dissolve. This again implies that there is no mass transfer between the nanobubble-like object and the liquid. Two stability theories (3 & 4), the dynamic equilibrium theory and limited diffusion theory, both depend on a mechanism that involves gas in- and out-flux. Therefore these two theories are in contradiction with the present results for the analyzed features and, for these nanobubble-like objects created by deposition, we have to turn our attention to the two remaining theories. Either these nanobubble-like objects have a gas-impermeable shell or these objects are simply not bubbles, but droplets of contamination. Both theories depend on a certain concentration of contaminants present in the system. Investigating the literature revealed that a variety of contamination sources could possibly play a role. These sources include, but are not limited to: glue from the adhesive tape used for cleaving HOPG, poor quality solvents, plastic syringes, flexible tubes, and air quality. Also, when employing the ethanol-exchange to nucleate nanobubbles a lot of care has to be taken as ethanol is especially susceptible to distribute any organic contaminants present in the nucleation procedure.

The procedures and materials used in our deposition experiment were scrutinized for any possible contaminant sources. Nonetheless, in our case it turned out that the use of sterile disposable plastic syringes and/or disposable needles was a crucial step for the nucleation of the nanobubble-like objects. We checked this finding by depositing a drop of water on freshly cleaved HOPG using either a glass syringe and a full-metal needle or a plastic syringe and a disposable needle. Fig. 3 shows six experiments, labeled and performed in the order I–VI, in which a glass (in experiments I, III, and V) or a disposable plastic syringe and a disposable needle (in experiments II, IV, and VI) were used to deposit the water on the substrate. It clearly shows that no nanobubble-like objects were observed if a glass syringe was used, however, in the case of a plastic syringe with a disposable needle, objects looking like nanobubbles are found. In both cases different positions on the HOPG sample were imaged with similar results. The water was kept in plastic syringes for durations ranging from a few minutes up to a day, which resulted in no significant changes in the nanobubble-like coverage. However, refilling the plastic syringe with water
several times does result in a reduced surface coverage with nanobubble-like objects. These degassing results are different from what was observed by Zhang et al. after degassing, where they show regions on the HOPG substrate where nanobubbles have disappeared. This can be explained by having used a procedure that does not introduce contamination and produces gaseous nanobubbles. However, this does not explain why nanobubbles remained stable in other regions.

Clearly, in our experiments a contaminant is present in the plastic syringes and/or disposable needles that results in the formation of these features on the surface. The question that remains is: what is the chemical nature of the contamination? To answer this question we performed X-ray photoelectron spectroscopy (XPS) on an HOPG sample on which a droplet of water, deposited using a plastic syringe combined with a disposable needle, was dried. In the resulting XPS spectrum more peaks show up than the normal carbon peak as would be the case for a clean HOPG surface, see Fig. 4A, so there clearly is some contamination on the surface. Table 1 shows a comparison of the atomic percentage, binding energies and O/Si peak ratio of our measurements with XPS data on PDMS from the literature. Comparing the peak positions, relative intensity and especially the valence electron spectrum with the literature it is possible to chemically characterize the contamination layer as polydimethylsiloxane (PDMS). Since no nanobubble-like objects are observed when using glass syringes with full-metal needles, the contamination has to be induced by the plastic syringe and/or the disposable needle. To confirm this, the metal cannula of the disposable needle was measured using XPS, see Fig. 4B. Also in this case a spectrum very similar to that of the dried HOPG was measured, which can be attributed to a 5 nm thick layer of PDMS on the cannula since no metal is visible in the spectrum. Interestingly, XPS measurements on the inside of the plastic syringe do not show any silicon peaks and therefore the syringe itself seems PDMS free, see Fig. 4C. Drying a drop of water, deposited using a plastic syringe and without a disposable needle, on HOPG resulted in a clean XPS spectrum without PDMS contamination.

The formation of nanobubble-like PDMS droplets would be quite consistent with the observation of Evans et al. In order to confirm whether PDMS contamination is responsible for the nanobubble-like objects we observe, we deliberately added PDMS to our system to confirm the formation of nanobubble-like objects by this polymer. For this we mixed 0.1 ml of PDMS (Sylgard 184, Dow Corning) with 0.4 l water by stirring vigorously. A droplet of the PDMS water mixture is then applied to the HOPG substrate using a glass syringe and imaged using the AFM. The resulting AFM images are strikingly similar to that of the nanobubble-like objects produced using plastic syringes with clean Millipore water, see Fig. 5A, or from any other...
nanobubble study for that matter. One of the characteristics of nanobubbles is that it can be moved and coalesced using the AFM tip by increasing the tip–sample interaction. We applied the same technique on the PDMS droplets by scanning a 2μm² area with increased force (highlighted using a dashed square) and imaging consecutively the same area using normal scanning conditions. Besides the appearance of these PDMS droplets their behavior during increased loads is also strikingly similar to that of nanobubbles reported in the literature.7,9–11,56,57 The PDMS droplets were moved by the AFM tip and some of the bubbles coalesced, see Fig. 5B. The resolution of the AFM image after the scan with an increased tip–sample interaction was reduced due to changes of the AFM tip.

The literature on nanobubbles states that their apparent shape is very much dependent on the set-point used for scanning the surface in the AFM.7–10,56 The set-point dependence of PDMS droplets was therefore compared to that of nanobubble-like objects created using plastic syringes in combination with disposable needles, see Fig. 6. The radius of curvature, \( R_c \), and the contact angle, \( \theta \), are \( R_c = 90 \text{ nm} \) and \( \theta = 50° \) for the object in Fig. 6A and \( R_c = 360 \text{ nm} \) and \( \theta = 22° \) for the object in Fig. 6B, both acquired from the 94% set-point measurement. In both cases the apparent height of the features is highly dependent on the AFM set-point. A decreasing set-point results in increased tip–sample interactions and soft features like bubbles and droplets are therefore easily deformed by the AFM tip as is the case for the features in this study.

The height, radius and contact angle of these features are similar to what is stated in the literature on nanobubbles. In addition, these features are soft, could be swept away using an AFM tip and disappeared after drying the surface.35 This finding, in retrospect, also may or may not affect some of our and others’ previous work in which plastic syringes in combination with disposable needles were used.7,10,18,34,35,56,58 Whether the features observed in these experiments are actual nanobubbles with a gas-impermeable shell induced by the PDMS or are in fact PDMS droplets is something to be investigated and lies outside the scope of this work. This purpose of the present study is to attain awareness in the nanobubble community, for possible sources of contamination that might in some cases distort experimental results. This could also resolve the mixed results found in the literature on a number of subjects related to nanobubbles.

Table 1: Atomic percentage, binding energy and O/Si peak ratio taken from the XPS measurements on freshly cleaved HOPG, HOPG on which a droplet of water had dried, deposited using a plastic syringe and a disposable needle, and data of an unused disposable needle. Literature data of XPS on PDMS are shown as a comparison and show a clear similarity to our data on the metal needle and contaminated HOPG.

|                         | Cl1s | O1s | Si2p |
|-------------------------|------|-----|------|
|                         | At.% | E_b [eV] | At.% | E_b [eV] | At.% | E_b [eV] | O/Si |
| Our data                |      |       |      |         |      |         |      |
| Clean HOPG             | 100  | 284.8 | 26.5 | 532.0   | 23.5 | 102.0   | 1.13 |
| Drying stain on HOPG   | 50.0 | 284.4 | 26.6 | 532.0   | 23.2 | 102.0   | 1.15 |
| Disposable needle       | 50.2 | 284.4 | 26.6 | 532.0   | 23.2 | 102.0   | 1.15 |
| Literature data PDMS   |      |       |      |         |      |         |      |
| Beamson and Briggs54   | 50.3 | 285   | 27.1 | 532.00  | 22.6 | 101.5   | 1.20 |
| Owen and Smith55       |      |       |      |         |      |         |      |
|                        | 50.3 | 284.4 | 26.6 | 532.00  | 22.6 | 101.5   | 1.20 |

Fig. 5: AFM images of an HOPG surface under a droplet of water mixed with PDMS and deposited using a glass syringe and a full-metal needle. Small PDMS droplets are settled on the substrate and have a similar appearance to nanobubbles (A). After scanning a 2 × 2 μm² area with increased force (highlighted using the dashed square), objects inside this area have moved to another position or coalesced (B). The z-range is 21 nm.

Fig. 6: The dependence of the geometry on four different AFM set-points for a free amplitude of 19 nm is shown for a nanobubble-like object in water on HOPG produced using a plastic syringe and a disposable needle (A). The geometry dependence on four different AFM set-points for a deliberately added PDMS droplet (using a glass syringe and a full-metal needle) on HOPG in water using a free amplitude of 19 nm shows similar results (B). In both cases the apparent height of the objects is heavily dependent on the set-point.
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We think that it is of utmost importance for the nanobubble from disposable needles, in the presence of contamination in some studies, originating on a variety of subjects. This variance could be resolved by the literature. The literature on nanobubbles is not in agreement.

The radius of a nanobubble with a radius of 1000 nm and less. In the case of a pinned curvature (fixed radius at $r = 250$ nm, i.e. fixed footprint area) the contact angle increases only by a fraction of its original value (B). The contact angle never exceeds 90°.

4 Conclusion

We have studied the resistance of nanobubble-like objects obtained by droplet deposition on HOPG against a gas-depleted environment using two different experimental techniques. First, the nanobubble-like objects were exposed to a degassed water flow and secondly the ambient pressure was decreased to 20 mbar. In both cases the coverage of the nanobubble-like objects remained virtually unchanged. An in-depth study of possible contamination sources in the procedures and materials used during the experiment showed that in our case the sterile disposable needles were the source of contamination.

The chemical nature of the contamination was concluded to be PDMS. Both the nanobubble-like objects and the deliberately formed PDMS droplets can be moved and coalesced using the AFM tip and their apparent shapes highly depend on the used set-point. The nanobubble-like objects that we nucleated in this way behave not differently from the nanobubbles discussed in the literature. The literature on nanobubbles is not in agreement on a variety of subjects. This variance could be resolved by the presence of contamination in some studies, originating from disposable needles, influencing the experimental results.

We think that it is of utmost importance for the nanobubble community to be aware of the subtlety of contamination sources.

5 Appendix A

Assuming fully gaseous nanobubbles, a reduction of the ambient pressure should lead to an expansion of the bubbles' volume. A sufficient increase in volume could result in contact and coalescence of nanobubbles. The question is: can we expect coalescence for these sizes at such low pressures? To answer this question we calculate the effect of a reduced pressure on nanobubbles, using the assumptions that: (i) there is no mass-transfer between the bubble and liquid, (ii) the temperature is constant, and (iii) that either the radius or the contact angle is fixed. Assumption (i) is, in this specific case, justified by the results obtained from the degassed water flow experiment. Assumption (ii) is not completely true since the evaporation of the liquid lowers the temperature, however, this results in only a small change in absolute temperature. Using the aforementioned assumptions reduces the ideal gas-law to the equation:

$$P_1V_1 = P_2V_2. \quad (1)$$

$P$ is the pressure inside the bubble and $V$ the volume of the bubble, where subscripts 1 and 2 denote the atmospheric and reduced pressure conditions respectively. The pressure in the bubble is the result of the combination of ambient pressure and Laplace pressure, $P_{\text{lap}} = 2\gamma/R_c$ (the hydrostatic pressure is negligible, for the 5 nm water column it is only $P_{\text{hyd}} = \rho g h = 0.5$ mbar), i.e. eqn (1) can be rewritten as:

$$\left(\frac{P_{e,1} + 2\gamma}{R_{c,1}}\right)V_1 = \left(\frac{P_{e,2} + 2\gamma}{R_{c,2}}\right)V_2. \quad (2)$$

$P_{e,1}$ and $P_{e,2}$ are the ambient pressures for atmospheric and reduced pressure conditions respectively. $R_{c,1}$ and $R_{c,2}$ are the radii of curvature of the nanobubble under atmospheric and reduced pressure conditions, and $\gamma$ is the surface tension of water, $\gamma = 72$ mN m$^{-1}$. Using the spherical cap geometry of a nanobubble to acquire a relationship for the nanobubble volume results in the following equations:

$$V(R_c, h) = \frac{1}{3} \pi h^2 \left(3 R_c - h\right), \quad (3)$$

$$R_c(r, \theta) = \frac{r}{\sin \theta}, \quad (4)$$

$$h(r, \theta) = \frac{r}{\sin \theta} - r \cot \theta, \quad (5)$$

where $h$ is the height of the nanobubble, $r$ is the radius of the contact line, and $\theta$ is the contact angle (measured in the gas-phase). A combination of eqn (3)–(5) results in the following expression for the nanobubble volume:

$$V(r, \theta) = X(\theta)r^3, \quad (6)$$

$$X(\theta) = \frac{1}{3} \pi \left(\frac{2}{\sin \theta} + \cot \theta\right) \left(\frac{1}{\sin \theta} - \cot \theta\right)^2. \quad (7)$$

So with relationships eqn (4), (6) and (7), eqn (2) is transformed into a function of $r$ and $\theta$:

$$\left(\frac{P_{e,1} + 2\gamma \sin \theta_1}{r_1}\right)X(\theta_1)r_1^3 = \left(\frac{P_{e,2} + 2\gamma \sin \theta_2}{r_2}\right)X(\theta_2)r_2^3. \quad (8)$$

Assuming a fixed contact angle of 20° (160° in the liquid phase), which is a typical value for nanobubbles, we numerically calculate the change in radius as a function of pressure from 1 atm to 1 mbar (100 Pa) for different initial radii, see Fig. 7A. Remarkably, for typical nanobubbles, which have a radius between 100 and 1000 nm, the increase in radius is rather limited. For the second case of a fixed radius of the
nanobubble (fixed at 250 nm), the expansion in volume is achieved by an increase in contact angle, see Fig. 7B. Also in this case the contact angle increases only with a fraction of the original value, always remaining smaller than 90°, so the lateral size remains constant. The limited increase in contact angle or radius can be explained by the enormous Laplace pressure for small bubbles; a reduction in ambient pressure leads to only a small reduction in the bubbles’ internal pressure. Therefore, it is not surprising that no coalescence of surface nanobubbles is observed at a reduced pressure of 20 mbar.

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References

1 M. A. Hampton and A. V. Nguyen, Adv. Colloid Interface Sci., 2010, 154, 30–55.
2 V. S. J. Craig, Soft Matter, 2011, 7, 40–48.
3 J. R. T. Seddon and D. Lohse, J. Phys.: Condens. Matter, 2011, 23, 133001.
4 J. L. Parker, P. M. Claesson and P. Attard, J. Phys. Chem., 1994, 98, 8468–8480.
5 S.-T. Lou, Z.-Q. Ouyang, Y. Zhang, X.-J. Li, J. Hu, M.-Q. Li and F.-J. Yang, J. Vac. Sci. Technol., B: Microelectron. Nanometer Struct.–Process., Meas., Physom., 2000, 18, 2573–2575.
6 N. Ishida, T. Inoue, M. Miyahara and K. Higashitani, Langmuir, 2000, 16, 6377–6380.
7 W. Walczyk, P. M. Schön and H. Schönherr, J. Phys.: Condens. Matter, 2013, 25, 184005.
8 C. W. Yang, Y. H. Lu and I. S. Hwang, J. Phys.: Condens. Matter, 2013, 25, 184010.
9 B. Zhao, Y. Song, S. Wang, B. Dai, L. Zhang, Y. Dong, J. Lü and J. Hu, Soft Matter, 2013, 9, 8837–8843.
10 B. M. Borkent, S. de Beer, F. Mugele and D. Lohse, Langmuir, 2010, 26, 260–268.
11 S. Wang, M. Liu and Y. Dong, J. Phys.: Condens. Matter, 2013, 25, 184007.
12 M. Holmberg, A. Kühle, J. Garnaes, K. A. Morch and A. Boisen, Langmuir, 2003, 19, 10510–10513.
13 A. Agrawal, J. Park, D. Y. Ryu, P. T. Hammond, T. P. Russell and G. H. McKinley, Nano Lett., 2005, 5, 1751–1756.
14 S. Yang, S. M. Dammer, N. Bremond, H. J. W. Zandvliet, E. S. Kooij and D. Lohse, Langmuir, 2007, 23, 7072–7077.
15 Y. Wang, B. Bhushan and X. Zhao, Nanotechnology, 2009, 20, 045301.
16 X. Zhang and N. Maeda, J. Phys. Chem. C, 2011, 115, 736–743.
17 M. Guan, W. Guo, L. Gao, Y. Tang, J. Hu and Y. Dong, ChemPhysChem, 2012, 13, 2115–2118.
18 R. P. Berkelaar, J. R. T. Seddon, H. J. W. Zandvliet and D. Lohse, ChemPhysChem, 2012, 13, 2213–2217.
19 X. H. Zhang, N. Maeda and V. S. J. Craig, Langmuir, 2006, 22, 5025–5035.
20 M. Mazumder and B. Bhushan, Soft Matter, 2011, 7, 9184–9196.
21 X. H. Zhang, A. Khan and W. A. Ducker, Phys. Rev. Lett., 2007, 98, 136101.
22 X. H. Zhang, A. Quinn and W. A. Ducker, Langmuir, 2008, 24, 4756–4764.
23 X. H. Zhang, X. D. Zhang, S. T. Lou, Z. X. Zhang, J. L. Sun and J. Hu, Langmuir, 2004, 20, 3813–3815.
24 X. H. Zhang, X. Zhang, J. Sun, Z. Zhang, G. Li, H. Fang, X. Xiao, X. Zeng and J. Hu, Langmuir, 2007, 23, 1778–1783.
25 X. H. Zhang, G. Li, N. Maeda and J. Hu, Langmuir, 2006, 22, 9238–9243.
26 A. Brotchie and X. H. Zhang, Soft Matter, 2011, 7, 265–269.
27 J. Yang, J. Duan, D. Fornasiero and J. Ralston, J. Phys. Chem. B, 2003, 107, 6139–6147.
28 M. A. J. van Limbeek and J. R. T. Seddon, Langmuir, 2011, 27, 8694–8699.
29 J. D. Miller, Y. Hu, S. Veeramasuneni and Y. Lu, Colloids Surf., A, 1999, 154, 137–147.
30 R. Steitz, T. Gutberlet, T. Hauss, B. Klösgen, R. Krastev, S. Schemmel, A. C. Simonsen and G. H. Findenegg, Langmuir, 2003, 19, 2409–2418.
31 A. Poynor, L. Hong, I. K. Robinson, S. Granick, Z. Zhang and P. A. Fenter, Phys. Rev. Lett., 2006, 97, 266101.
32 M. Mezger, H. Reichert, S. Schöder, J. Okasinski, H. Schröder, H. Dosch, D. Palms, J. Ralston and V. Honkimäki, Proc. Natl. Acad. Sci. U. S. A., 2006, 103, 18401–18404.
33 A. C. Simonsen, P. L. Hansen and B. Klösgen, J. Colloid Interface Sci., 2004, 273, 291–299.
34 B. M. Borkent, S. Dammer, H. Schönherr, G. J. Vancso and D. Lohse, Phys. Rev. Lett., 2007, 98, 204502.
35 R. P. Berkelaar, H. J. W. Zandvliet and D. Lohse, Langmuir, 2013, 29, 11337–11343.
36 X. Zhang, A. Kumar and P. J. Scales, Langmuir, 2011, 27, 2484–2491.
37 X. Zhang, D. Y. C. Chan, D. Wang and N. Maeda, Langmuir, 2013, 29, 1017–1023.
38 P. S. Epstein and M. S. Plesset, J. Chem. Phys., 1950, 18, 1505–1509.
39 S. Ljunggren and J. C. Eriksson, Colloids Surf., A, 1997, 130, 151–155.
40 D. R. Evans, V. S. J. Craig and T. J. Senden, Phys. A, 2004, 339, 101–105.
41 W. A. Ducker, Langmuir, 2009, 25, 8907–8910.
42 S. Das, J. H. Snoeijer and D. Lohse, Phys. Rev. E: Stat., Nonlinear, Soft Matter Phys., 2010, 82, 056310.
43 H. Peng, M. A. Hampton and A. V. Nguyen, Langmuir, 2013, 29, 6123–6130.
44 X. Zhang, M. H. Uddin, H. Yang, G. Toikka, W. Ducker and N. Maeda, Langmuir, 2012, 28, 10471–10477.
45 M. P. Brenner and D. Lohse, Phys. Rev. Lett., 2008, 101, 214505.
46 J. R. T. Seddon, H. J. W. Zandvliet and D. Lohse, Phys. Rev. Lett., 2011, 107, 116101.
47 N. D. Petsev, M. S. Shell and L. G. Leal, *Phys. Rev. E: Stat., Nonlinear, Soft Matter Phys.*, 2013, **88**, 010402.
48 J. H. Weijs and D. Lohse, *Phys. Rev. Lett.*, 2013, **110**, 054501.
49 D. Wise and G. Houghton, *Chem. Eng. Sci.*, 1966, **21**, 999–1010.
50 B. Rezania, M. Dorn, N. Severin and J. P. Rabe, *J. Colloid Interface Sci.*, 2013, **407**, 500–504.
51 A. Häbich, W. Ducker, D. E. Dunstan and X. H. Zhang, *J. Phys. Chem. B*, 2010, **114**, 6962–6967.
52 J. A. Carr, K. S. Nalwa, R. Mahadevapuram, Y. Chen, J. Anderegg and S. Chaudhary, *ACS Appl. Mater. Interfaces*, 2012, **4**, 2831–2835.
53 G. R. Buettner, B. D. Scott, R. E. Kerber and A. Mügge, *Free Radical Biol. Med.*, 1991, **11**, 69–70.
54 G. Beamson and D. Briggs, *High Resolution XPS of Organic Polymers: The Scienta ESCA300 Database*, John Wiley & Sons, 1992.
55 M. J. Owen and P. J. Smith, *J. Adhes. Sci. Technol.*, 1994, **8**, 1063–1075.
56 W. Walczyk and H. Schönherr, *Langmuir*, 2013, **29**, 620–632.
57 S. Yang, E. S. Kooij, B. Poelsema, D. Lohse and H. J. W. Zandvliet, *EPL*, 2008, **81**, 64006.
58 E. Dietrich, H. J. W. Zandvliet, D. Lohse and J. R. T. Seddon, *J. Phys.: Condens. Matter*, 2013, **25**, 184009.
59 B. M. Borkent, H. Schönherr, G. L. Caër, B. Dollet and D. Lohse, *Phys. Rev. E: Stat., Nonlinear, Soft Matter Phys.*, 2009, **80**, 036315.