

# 3D Sub-Nanometer Analysis of Glucose in an Aqueous Solution by Cryo-Atom Probe Tomography

**Tim Schwarz**

University of Stuttgart <https://orcid.org/0000-0001-9348-4160>

**Carolin Dietrich**

University of Stuttgart

**Jonas Ott**

University of Stuttgart

**Eric Weikum**

University of Stuttgart

**Robert Lawitzki**

University of Stuttgart

**Helena Solodenko**

University of Stuttgart

**Efi Hadjixenophontos**

University of Stuttgart

**Baptiste Gault**

Max Planck Institute for Iron Research <https://orcid.org/0000-0002-4934-0458>

**Johannes Kästner**

University of Stuttgart, Institute for Theoretical Chemistry <https://orcid.org/0000-0001-6178-7669>

**Guido Schmitz**

University of Stuttgart

**Patrick Stender** (✉ [patrick.stender@mp.imw.uni-stuttgart.de](mailto:patrick.stender@mp.imw.uni-stuttgart.de))

University of Stuttgart <https://orcid.org/0000-0002-2324-5479>

---

## Article

**Keywords:** Atom Probe Tomography (APT), cryogenic workflows, cryogenically-frozen solutions, biological materials

**Posted Date:** February 19th, 2021

**DOI:** <https://doi.org/10.21203/rs.3.rs-245717/v1>

**License:** © ⓘ This work is licensed under a Creative Commons Attribution 4.0 International License.

[Read Full License](#)



# Abstract

Atom Probe Tomography (APT) is currently a well-established technique to analyse the composition of solid materials including metals, semiconductors and ceramics with up to near-atomic resolution. Using an aqueous glucose solution, we now extended the technique to frozen solutions. While the mass signals of the common glucose fragments  $C_xH_y$  and  $C_xO_yH_z$  overlap with  $(H_2O)_nH$  from water, we achieved stoichiometrically correct values via signal deconvolution. Density functional theory (DFT) calculations were performed to investigate the stability of the detected pyranose fragments. This paper demonstrates APT's capabilities to achieve sub-nanometre resolution in tracing whole glucose molecules in a frozen solution by using cryogenic workflows. We use a solution of defined concentration to investigate the chemical and spatial resolution capabilities as a step toward the measurement of biological molecules in solution in 3D with sub-nanometre resolution by using cryo-APT. Our analyses take analytical techniques to a new level, since chemical characterization methods for cryogenically-frozen solutions or biological materials are limited.

## Introduction

Atom probe tomography (APT) <sup>1-5</sup> has become a key technique to investigate chemical structures at the near-atomic scale across a broad variety of materials systems <sup>6</sup>. Depending on the controlled removal of individual atoms from a sharp needle-like emitter in the form of ions, this destructive method transforms the studied sample volume into a 3D computer model with individual ions characterized chemically. One of the current trends in APT is to gain a deeper understanding of soft matter in general and liquids in particular <sup>7-16</sup>. While thin layers of ice, created by condensed vapour on metallic tips, have been studied previously <sup>17-21</sup>, the newly available cryo-transfer systems <sup>22,23</sup> facilitate the preparation and transport of APT specimens from frozen liquids with large sampled volumes. Such a step-change in capability may enable fundamental atomic-scale studies of solutions, bio(macro)molecules in their natural environment, and potentially catalytic reactions and other liquid-solid interfaces, e.g., wet corrosion.

Challenges arising from analysing soft matter by APT are not only the complex preparation routes, but also the rather complex nature of the field evaporation process. While single atoms typically evaporate from metallic samples, molecular ions of varying size <sup>24-26</sup> manifesting in a large variety are typically observed for non-metallic samples, with just a small fraction of atomic ions sometimes arising from the dissociation of metastable molecular ions during the flight. Earlier measurements of thin water layers <sup>17,27,28</sup> and recent observations from thick ice layers <sup>15,16</sup> show a strong tendency in the creation of protonated water clusters of varying size. Atomic hydrogen and oxygen are rarely detected and mostly fragments from the dissociation of molecular ions <sup>29-31</sup>. This does not only limit the spatial resolution but also gives a higher probability of peak from different species overlapping, hindering the detection of signals stemming from matter incorporated or dispersed in the water matrix.

In principle APT is a calibration-free technique to determine the chemical composition by simply counting each atom, but deviations are reported for carbides<sup>32–34</sup>, boron<sup>35</sup>, nitrides (AlN,GaN)<sup>29,31,36,37</sup>, oxides<sup>29,36,38</sup>, alkali halides and other semiconducting materials<sup>39–41</sup>. In the case of the latter, the composition measured by APT is not as accurate, and the specific loss of oxygen and nitrogen is commonly attributed to neutral species formed by molecular dissociations<sup>29–31</sup>, thereby impeding their detection. These additional uncertainties should be accounted for in discussions otherwise the derived conclusions can be erroneous. If such effects cannot be avoided, understanding the loss factors and, in the simplest cases, defining and using correction factors becomes of paramount importance. For measurements of frozen liquids or solutions, or objects dissolved in a liquid environment, standards and possible corrections are not yet available.

The importance of water in biological processes is undisputable. Most biochemical processes are enabled by the unique properties of water as a polar medium. Central examples of biologically relevant molecules are carbohydrates, glucose being one of the simplest ones. They are crucial for the energy balance and integral to most important biological information storage, RNA and DNA<sup>42</sup>. Glucose is a monosaccharide consisting of a pyranose ring, which contains five carbon atoms and one oxygen atom, with a hydroxymethyl group at C5. The molecular formula for glucose is C<sub>6</sub>O<sub>6</sub>H<sub>12</sub>. In its solid form, the ring is usually closed, while in solution a small fraction of the molecules transforms into an open chain. The most dominant representations of this type are α- and β-pyranose<sup>43</sup>. Its molar mass amounts to 180.16 g mol<sup>-1</sup> and the maximum solubility in water is 470 g l<sup>-1</sup> at room temperature. This equates to a solubility of 2.608 mol l<sup>-1</sup> or, on an atomic scale, 2 glucose molecules per 42 water molecules.

Equipped with the background knowledge from the prior analysis of pure Mili-Q water<sup>15</sup>, we focus in this study on the differences in the field evaporation behaviour between a saturated glucose solution, bulk glucose and pure water to broaden the discussion basis and advance the understanding and map the possibilities and limitations inherent to the application of state-of-the-art cryo-APT to the analysis of solutions, molecules and biological materials.

## Results And Discussion

We acquired through laser-assisted APT data from 7 specimens yielding at least 30 million atoms each. Stable measurement conditions, with a homogeneous detector pattern and a steady smoothly increasing voltage curve over time, were obtained with a detection rate of 3 ions per 1000 laser pulses, at a repetition rate of 100 kHz and a pulse energy between 40 and 80 nJ. Chemical information is derived from the measured time of flight (ToF). The ToF measurement is continuously restarted by the emission of a new laser pulse, defining the maximum opening time of the ToF measurement window. While the maximum detectable mass for a flight length of 120 mm at 6 kV and a repetition rate of 200 kHz accounts to 200 u e<sup>-1</sup>, the range can be extended at 100 kHz to about 800 u e<sup>-1</sup>. Such large masses are normally of limited interest for APT, but at 100 kHz we could detect significant signals stemming from large molecules up to a mass of 350 u e<sup>-1</sup>. For high laser repetition rates these large fragments will have not enough time to

reach the detector before a new laser pulse will be initiated. Besides, the decreased repetition rate enables samples with low thermal conductivity to cool down between laser pulses manifesting in an improved signal to noise ratio <sup>16</sup>.

The mass spectrum of the saturated glucose solution (Fig. 1) reveals more than 140 mass peaks, a clear increase compared to the pure water spectra <sup>15</sup>. Peak identification is ambiguous due to the large number of possible combinations of charge states and their elemental combinations. We generally favoured the simplest explanation for a specific mass signal, using the lowest possible charge state option to assign a molecular identity to each signal, since the effective evaporation field for water is low <sup>15,16</sup> and thus lower charge states are more likely to be observed. Different approaches to assign individual peaks will be discussed below and compared to the expected ratio of water to glucose molecules.

### **Mass spectra comparison of glucose solution to pure water**

Since water is the majority compound in the saturated glucose solution (470 g L<sup>-1</sup> at 20°C), it is reasonable to expect a mass spectrum with significant similarity to pure water <sup>15</sup>. We observe the same protonated water-containing ions (H<sub>2</sub>O)<sub>n</sub>H<sup>+</sup> with  $n = 1-5$ , with peaks at  $m/q = 19, 37, 55, 73$  and  $91 \text{ u e}^{-1}$ , as for pure water. Comparing the peak shapes of the signals match to the previous measurement of water with regard to their peak width, relative intensity and the pronounced tailing <sup>15</sup>. Furthermore, higher charged signals corresponding to the composition (H<sub>2</sub>O)<sub>n</sub>(OH)<sub>m</sub><sup>2+</sup> and their satellite peaks ( $\Delta m = +1$ ) match as well. Distinct differences become visible for larger atomic masses. Periodic signals at masses larger than  $m/q = 100 \text{ u e}^{-1}$  are readily visible in Fig. 2. These high-mass signals have a distinct mass difference of  $18 \text{ u e}^{-1}$ . All signals stemming from  $m/q$  ratios above  $91 \text{ u e}^{-1}$  could in principle be explained by large protonated water clusters (H<sub>2</sub>O)<sub>n</sub>H<sup>+</sup> with  $n > 5$ , which, however, are not observed in this quantity in earlier measurements of pure water (Fig 2).

The glucose molecule has a mass of  $180 \text{ u e}^{-1}$ . A reasonably strong signal appears at  $m/q = 181 \text{ u e}^{-1}$  and can be also interpreted as a protonated glucose molecule. Glucose contains seven hydrogen atoms and five OH-groups attached to different carbon atoms. The periodic peaks with lower masses than  $181 \text{ u e}^{-1}$  can be explained by the simultaneous removal of an OH group and one H-atom from the carbon atoms forming the pyranose ring. The removal of (HO+H)<sub>n</sub> with  $n = 0 - 5$  would result in the distinct spacing of mass peaks less than  $181 \text{ u e}^{-1}$  with a mass difference of  $\Delta m/q = 18 \text{ u e}^{-1}$  (H<sub>2</sub>O<sup>+</sup>). This periodicity stops once all OH groups are removed (Fig. 2b), at  $m/q = 91 \text{ u e}^{-1}$ . To complete the chain of evidence, relative peak intensities are determined for protonated molecules (Fig. 3 a). The relative intensities of the water cluster signal at  $m/q = 37, 55, 73,$  and  $91 \text{ u e}^{-1}$  agree with the intensity distribution determined for pure water, implying a similar effective evaporation field strength for both liquids. Beyond  $m/q = 91 \text{ u e}^{-1}$  the relative intensities increase again in the case of glucose solution ( $m/q = 91, 109, 127, 145, 163, 181, \text{ u e}^{-1}$ ). Since measurement conditions are comparable and such large clusters of water

molecules were absent in the case of pure water, all signals with high masses above  $91 \text{ u e}^{-1}$  are therefore attributed to glucose.

### Molecular ions consideration

DFT calculations have been performed to calculate the structure and the stability of the formed glucose fragments. The molecular ions smaller than the glucose molecule are depicted in Figure 4a. In each case, the equivalent of  $n$  water molecules  $n = 1 - 5$  has been removed compared to the structure at  $m/q = 181 \text{ u}$ . At the mass-to-charge state ratios of  $m/q = 127 \text{ u e}^{-1}$  and  $m/q = 109 \text{ u e}^{-1}$  isomers with planar 6-membered rings were found. These 6-membered pyrylium rings are aromatic, which renders these isomers particularly stable. The last molecule at  $m/q = 91 \text{ u e}^{-1}$  has a linear structure.

After removal of OH+H the remainder of the glucose molecule undergoes a molecular rearrangement. Due to their inherent mobility, the remaining hydrogen atoms relocate and bind to a different carbon atom. In the case of  $\text{C}_6\text{O}_6\text{H}_{13}^+$  ( $m/q = 181 \text{ u e}^{-1}$ ), it is possible to remove an  $\text{H}_2\text{O}$  from one carbon atom.  $\text{H}_2\text{O}$  is removed from C2 and subsequently one hydrogen relocates from C1 to C2 resulting in the structure shown in Figure 4a. It is unknown which OH molecule and which hydrogen atom are removed in the smaller fragments.

Each of the structures has been found to be stable against the abstraction of the equivalence of one  $\text{H}_2\text{O}$  from the ring structure except for the clusters at  $m/q = 181 \text{ u e}^{-1}$  and  $m/q = 163 \text{ u e}^{-1}$ , which are therefore metastable. However, in those cases the calculated dissociation energy is rather small,  $1.5 \text{ kJ mol}^{-1}$  for  $m/q = 181 \text{ u e}^{-1}$  and  $8 \text{ kJ mol}^{-1}$  for  $m/q = 163 \text{ u e}^{-1}$  (table 1).

Molecular structures with alternative compositions were found for  $m/q = 145, 127$  and  $109 \text{ u e}^{-1}$  (see Figure 4a marked with \*). In contrast to the structures shown in Figure 4a and 4b they contain only 5 instead of 6 carbon atoms. The formation of these structures is, however, considered unlikely as it would require a fragmentation of the carbon skeleton of the molecule, followed by the simultaneous removal of one carbon, one oxygen and several hydrogen atoms. A direct comparison of the energy to those structures shown in Figure 4b is impossible due to the different composition.

The reaction energies listed in table 1 are calculated using

$\Delta E = E(x) - E(x - 18) - E(n)$	(1)
--------------------------------------	-----

Where  $E(x)$  denotes the electronic energy + Zero-point correction from the DFT calculations and  $x$  is equal to the respective mass to charge state ratio  $m/q$ .  $E(n) = E(x=18)$  being either  $E(\text{H}_2\text{O})$  or  $E(\text{H+OH})$ .

Table 1: Calculated energy values by DFT for selected glucose molecules and the respective reaction energies after removal of H<sub>2</sub>O or HO+H.

m/q=x	electronic energy + ZPE in Hartree	reaction energy in kJ/mol: E(n) = E(H <sub>2</sub> O)	reaction energy in kJ/mol: E(n) = E(HO + H)
H <sub>2</sub> O	-76,342912		
H+OH	-76,159834		
91	-304,999354		
109	-381,442318	-262.7	-743.7
127	-457,868141	-217.7	-698.4
145	-534,211090	-0.1	-480.8
163	-610,553428	1.5	-479.2
181	-686,893155	8.4	-472.3
199	-763,270441	-90.3	-570.9
217	-839,654259	-107.4	-588.1

Molecules with a mass-to-charge state ratio higher than  $m/q = 181 \text{ u e}^{-1}$  following with a  $\Delta m/q = 18 \text{ u e}^{-1}$  are detected in considerable quantity and most likely correspond to a glucose molecule with one or more additional cations ( $\text{H}^+$ ,  $\text{H}_3\text{O}^+$  and  $\text{H}_5\text{O}_2^+$ , depicted in Figure 4b) bound via hydrogen bonds to the glucose ring. An analysis of the partial charges shows how the charges are distributed within the molecule. Oxygen is more electronegative than carbon and accumulates negative charge. The most weakly-charged oxygen atom, with  $-0.6 \text{ e}$ , is located within the closed ring structure, while the oxygen in the hydroxyl groups is more strongly charged with  $-0.7 \text{ e}$  and the oxygen atoms in  $\text{H}_3\text{O}^+$  or  $\text{H}_5\text{O}_2^+$  carry between  $-0.9$  and  $-0.8 \text{ e}$ . All carbon atoms on the other hand only carry small charges ( $-0.09$  to  $+0.4 \text{ e}$ ). For this reason, the hydrogen atoms have different amounts of positive charge depending on their binding partner: around  $0.5 \text{ e}$  for hydrogen attached to oxygen and  $0.2 - 0.3 \text{ e}$  for hydrogen attached to carbon. In the case of the molecular cluster at the mass-to-charge ratio of  $m/q = 217 \text{ u e}^{-1}$ ,  $\text{H}_5\text{O}_2^+$ , also known as Zundel cation, is bound to the cluster via hydrogen bonds. 79% of the overall charge of the cluster is located in the  $\text{H}_5\text{O}_2$  group. Most of the molecule's positive charge is therefore located in the water component of the cluster and this portion increases with the size of the molecule. For the biggest cluster depicted in Figure 4b, the percentage increases to 97 % in the water component.

In a saturated glucose solution, there are 21 water molecules available to solvate each glucose molecule. Further investigations of molecules at higher mass-to-charge ratio with more than two additional water molecules with the formula  $C_6O_6H_{12}+(H_2O)_nH^+$  with  $n = 2 - 9$  could also be observed (see supplementary material Fig. A 5). The respective stabilities and the arrangement of larger water cluster around the glucose molecule were also computed. The DFT calculations show that it is energetically favourable to build compact water cluster bound via hydrogen bonds to the glucose molecule instead of arranging water molecules around the glucose molecule as a hydration shell (Fig. 4 b).

The apparent distribution of relative intensities above  $m/q = 91 \text{ u e}^{-1}$ , assumed to stem from glucose molecules show a distribution with the maximum centred at  $m/q = 127 \text{ u e}^{-1}$  (Fig. 3b). From the analysis of multi-hit events<sup>36</sup> on the detector no preferential co-evaporation of HO, H<sub>2</sub>O, H<sub>3</sub>O with a large glucose cluster could be observed. Even in the spectra for the single, double, triple etc. events, no significant difference in the detected species is observed (see supplementary part Fig. A 1-4). The loss of HO+H does not occur simultaneously. A way to explain the observed distribution is to assume a step-by-step evaporation process with certain evaporation probabilities influenced by the remaining molecular fragment. Mass 181 is assumed to be generated in the first evaporation attempt  $k = 1$ .

To match the observed frequency distribution with this simple concept, the probability for evaporation has to increase continuously after each removal of an HO+H fragment ( $k > 1$ ). Or in other word, the further removal of another HO+H event becomes harder (see Table 1). Once all OH groups have been released, the molecule opens into a linear structure (Fig. 4a) as this structure is energetically more stable. Fragments smaller than C<sub>6</sub> can be explained by the splitting of the C<sub>6</sub>H<sub>3</sub>O<sup>+</sup> chain. To derive these probabilities, the number of occurring complete molecules in the range between  $91 - 181 \text{ u e}^{-1}$  are normalized. The relative number of occurring complete molecules at  $m/q = 181 \text{ u e}^{-1}$  is equal to the probability of evaporating the respective molecule. The detected relative distribution is used to match the respective probabilities in each step according to:

$P_{Total}(1) = p_{evap}(1) = 0.12$	3
-------------------------------------	---

The probability to split off an HO+H fragment is even:

$p_{Split}(k) = 1 - p_{evap}(k)$ with $k = \text{evaporation step } 1 - 5$	4
--	---

In general, the probability to evaporate the glucose fragment in step  $k$  is described by

$p_{Total}(k) = p_{evap}(k) \cdot \prod_{n=1}^{k-1} p_{Split}(n)$ with $k \in \{2,3,4,5\}$	5
--	---

Table 2: Calculated evaporation and split-off probability of the glucose molecules assuming a simple multi-step random process.

Mass-to-Charge state ratio ( $u e^{-1}$ )	181	163	145	127	109	91
Normalized relative occurrence	0.12	0.13	0.21	0.23	0.175	0.148
Evaporation attempt $k$	1	2	3	4	5	6
Evaporation probability $p_{evap}(k)$	0.12	0.153	0.28	0.42	0.55	0.9999
H+OH Split-off probability $p_{split}(k)$	0.88	0.85	0.72	0.579	0.448	0.0001

Being a model, this simplified approach has limitations, but the decrease in split-off-probability is supported by DFT calculations and correlates with experiments (see Fig. 3c), since the remaining fragment become more stable against further dissociation with every water or HO + H molecule removed (see Table 2).

Between the main peaks at  $m/q = 109$  and  $217 u e^{-1}$ , small peaks appear that are separated by a distance of  $1 u e^{-1}$ , corresponding to a single additional hydrogen atom. There are always three very prominent peaks between the two main peaks, with a distance of  $\Delta m/q = 5, 7, 12 u e^{-1}$  from the previous peak. The peaks between the mass-to-charge state ratio of  $m/q = 109-181 u e^{-1}$  can be associated to  $C_{4-6}O_xH_y$ .

However, the intermediate peaks from  $m/q = 181-217 \text{ u e}^{-1}$  can only correspond to the molecular ions  $\text{C}_5\text{O}_5\text{H}_{11-14} + (\text{H}_2\text{O})_x\text{H}_y^+$  with  $x = 2 - 3$ .

The identification of an ion only by its mass, of course, does not tell anything about the internal structure of the detected species. In this specific case, the structure and the identity of the injected molecules are a priori known. But it would not be possible to differentiate common monosaccharides like galactose and fructose based on the molecular mass. Since current detectors used for APT have an efficiency around 80%, it is still doubtful, whether current reconstruction algorithms could assign the original molecule even when it evaporated atomically. This would require a more sophisticated analysis of the data, and likely statistical comparison with a database of fragments, as is done in proteomics<sup>44</sup>. Whether the observed frequency distribution of fragments as described in table 2 is unique for glucose and allows a clear distinction from other monosaccharides is unknown and must be evaluated in the future. Typically, the respective evaporation field scales with the sublimation energy, which is increasing with the melting temperature of the substance<sup>2,45</sup>. The monosaccharides at least differ in their melting temperatures (glucose 146°C, fructose 106°C, galactose 164°C). In a first view, one may speculate that the melting point mostly affects the probability to evaporate the complete molecule, while the probability to split OH or H in both cases covalently bond might be less influenced. Thus, the relative fractions of the high-mass peaks would change, which may result in a distinguishable pattern.

The exact identification of fragments for lower masses is even more complicated. For all lower mass peaks combinations of different  $\text{C}_x\text{H}_y / \text{C}_x\text{O}_y\text{H}_z$  permutations in various charge state modifications can be found. A full interpretation list can be found in Table A 1 of the supplementary part.

### **Comparison of a saturated glucose solution to bulk glucose**

In the measurements presented before, water is the majority component. Thus the situation becomes less transparent, as signals stemming from water claim the same positions within the spectrum than the sugar molecules. To avoid such superposition, we also investigated pure bulk glucose sample to identify peak positions of probably highest overlap. Still, the ionization field strength of frozen water is rather low (2 – 6 V/nm)<sup>17,28,46</sup> while we can expect for solid sugar field strengths of some tens of V/nm which may hinder a direct comparison.

Samples were prepared from pure solid bulk glucose by FIB lift-out (see Materials and Methods section)<sup>47</sup>. Comparing the mass spectrum of bulk glucose plotted in Fig. 5 with that of glucose in aqueous solution (Fig. 1), significant differences are readily visible. First, for bulk glucose no signals above  $m/q = 80 \text{ u e}^{-1}$  are detected, indicating that the average field condition is sufficient to split the pyranose rings from the glucose molecule. Focusing on lower mass signals, most peaks pertain to molecular ions, although small but significant C (12 u e<sup>-1</sup>) and H (1 and 2 u e<sup>-1</sup>) atomic signals are present as reported for other carbon-based system with varying finger prints<sup>48-51</sup>

Nevertheless, series of peaks appear that are spaced by  $1 \text{ u e}^{-1}$  up to a mass-to-charge state ratio of  $m/q = 80 \text{ u e}^{-1}$ . A full interpretation list can be found in Table A2 of the supplementary part. The most intense peaks are at  $m/q = 15 \text{ u e}^{-1}$ , corresponding to  $\text{CH}_3^+$ ,  $m/q = 29 \text{ u e}^{-1}$  attributed to  $\text{C}_2\text{H}_5^+$  or  $\text{COH}^+$ , and at  $m/q = 44 \text{ u e}^{-1}$  giving multiple identification options ( $\text{C}_3\text{H}_8^+$ ,  $\text{C}_2\text{OH}_4^+$  or  $\text{COO}^+$ ).

Various peaks can be explained by combinations of carbon, oxygen and hydrogen ( $\text{C}_x\text{O}_y\text{H}_z$ ). For example, in the range of  $m/q = 28 - 31 \text{ u e}^{-1}$   $\text{COH}_z$  combinations can fit. The interpretation is not univocal, many possibilities exist for each peak (see table A 2 supplementary part), and because of the electric field conditions, only singly charged molecules are considered. However, a comparison of all shown measurements reveals a very different fragmentation behaviour of the glucose depending on whether it is dissolved in water or evaporated as bulk material. Bulk glucose shows a high fragmentation rate, whereas glucose dissolved in water has a lower fragmentation rate and frequently evaporates as a complete molecule. This observation must be traced back to the different bonding conditions in the solid and dissolved state and the respective electrical field strength necessary to cause field evaporation.

A further difference can be observed regarding the intensities of the detected molecules  $\text{H}_2\text{O}^+$  and  $\text{H}_3\text{O}^+$ . In the aqueous medium the molecule  $\text{H}_3\text{O}^+$  is dominant, but in the solid state the molecule  $\text{H}_2\text{O}^+$  is much more present, which could be due to a cleaving effect of the HO and H parts of the pyranose ring.

This significant difference in the ratio of  $\text{H}_2\text{O}^+$  to  $\text{H}_3\text{O}^+$  seems to be a reasonable marker in between the discussed measurements. Generally, the mass signals stemming from the solid samples display sharper peaks and less tailing compared to the aqueous solution samples. A simple explanation by a difference in heat conductivity  $\lambda$  is not feasible, since the reported values for a saturated aqueous glucose solution and glucose crystals are both in the range of  $0.5 \pm 0.1 \text{ Wm}^{-1}\text{K}^{-1}$  <sup>52,53</sup> while the reported value for ice are about 5-10 times higher depending on the crystal structure with the lower limit given by low-density amorphous ice to be  $0.6 \text{ Wm}^{-1}\text{K}^{-1}$  <sup>54</sup>. The increased tailing has to be attributed to a more direct influence of the ice <sup>15</sup>.

It is insightful to discuss and interpret the correct determination of the oxygen content. The interpretation of all signals stemming from pure bulk glucose as carbohydrates  $\text{C}_x\text{H}_z$  results in an extreme underestimation of oxygen (Table 3), which cannot be simply explained by the loss of neutral oxygen during the APT measurement.

Table 3: Calculated stoichiometry by different peak interpretations. The overlap of  $C_xH_y$  and  $C_xO_yH_z$  in certain areas were calculated (see supplementary part Table A2 marked in red).

	$C_x/H_y$	$C_xO_yH_z$	$C_xH_y/C_xO_yH_z$ 0.5/0.5	$C_xH_y/C_xO_yH_z$ 0.25/0.75	$C_xH_y/C_xO_yH_z$ 0.2/0.8	$C_xH_y/C_xO_yH_z$ 0.1/0.9
Oxygen loss	86.25%	0	50.29%	23%	15.66%	0.13%
Oxygen excess	0	18.31%	0	0	0	0
Hydrogen loss	0	32.90%	0	8.16%	12.33%	21.73%
Hydrogen excess	31.2%	0	8.65%	0	0	0

Instead, one may favour the interpretation of the mass signals as  $C_xO_yH_z$  molecules. In this case the amount of available oxygen is overestimated by 18.31 %. Thus, the reality seems to be a convolution between both possibilities. Further experiments are needed to resolve the sources of uncertainty.

### **Determination of compositional information for the aqueous solution**

In general, compositional information is of tremendous interest. Identification of compositions, and therefrom phases, diffusion coefficients, segregations factors are important cornerstones to identify processes and underlying mechanisms. In our case, the initial composition of the solution is well known. Since glucose and other sugars act as cryoprotectants and form glasses for highly concentrated solutions<sup>55</sup> it can be used to judge the peak identification and to identify open questions and problems to be addressed in the future.

A bijective identification of the retrieved mass information is difficult due to the already discussed overlapping mass signals and the large number of possible explanations. Various combinations of  $(H_2O)_nH$ ,  $C_xH_y$ ,  $C_xO_yH_z$  molecules would deliver reasonable explanations. Though a combination of all sources is highly likely and increases the complexity and leads to uncertainty and loss of local spatial resolution.

For stoichiometry calculations, all peaks are identified, and the resulting number of individual events counted. Different interpretation approaches are used, and the unfolding molecule was split into its atomic components carbon (C), oxygen (O) and hydrogen (H). In addition, a baseline correction was performed to obtain an accurate determination of the atomic ratios.

All obtained carbon atoms are assumed to be originating from glucose molecules. By dividing the total number of carbon atoms by a factor of six, the total amount of glucose molecules is calculated. In a next step the total number of glucose molecules is multiplied by six and subtracted from the total number of oxygen atoms. The remaining oxygen atoms are assumed to stem from water molecules. The saturated glucose solution should exhibit a mass fraction of glucose to water of 0.47, which corresponds to 21.27 water molecules per glucose molecule for a saturated solution. For the calculation of the stoichiometry

different peak interpretation approaches were used (Table A 3,4 supplementary part) and the results are listed in Table 4.

Interpreting the signals at  $m/q = 17, 18$  and  $19 \text{ u e}^{-1}$  as  $\text{HO}, \text{H}_2\text{O}$  and  $\text{H}_3\text{O}$ , respectively, and all other peaks as molecules by various combinations of  $\text{C}_x\text{H}_y$  results in a glucose to water mass-ratio of 519.94. The ratio deviates massively from the theoretical value by a factor of 1100.

Using our earlier measurements of pure water<sup>15</sup> as a footprint, all peaks visible in the water spectrum are attributed to water, while large molecules are assumed to be large glucose molecules. Signals at masses below  $91 \text{ u e}^{-1}$  which were not visible in the water measurements are assumed to be hydrocarbons with the formula  $\text{C}_x\text{H}_y$ . The mass-ratio of glucose to water is calculated to be 1.40. In this case, this corresponds to 7.17 water molecules per glucose molecule. The ratio still deviates from the theoretical value by a factor of 3. However, it shows that the ratio has improved by a factor of 366 with only the pure re-interpretation of the water molecules. Nevertheless, there is also an underestimation of oxygen, which is caused by the overlapping of peaks or by wrong interpretation of the carbon chains.

As already mentioned, various signals can also be explained by  $\text{C}_x\text{O}_y\text{H}_z$ . Using this interpretation only for signals not stemming from water leads to a ratio of 0.61, which deviates by a factor of only 1.3 from the theoretical value of 0.47. This corresponds to 16.36 water molecules for each glucose molecule.

Since we observe the evaporation of large and mostly intact glucose rings, it seems unlikely from previous considerations and DFT calculations, that the pyranose ring is cracked and fragments of  $\text{C}_x\text{O}_y\text{H}_z$  are observed. However, DFT calculation for the fragment with mass  $m/q = 91 \text{ u e}^{-1}$  reveal a chain like structure with unsaturated carbon atoms. The bond energy between a C-O bond<sup>56</sup> is slightly higher than that of a C-C bond<sup>56</sup>.  $\text{H}_x\text{O}$  molecules of water can possibly recombine with the adjacent glucose fragment and therefore  $\text{C}_x\text{O}_y\text{H}_z$  fragments would allow to explain the observed mass signals.

To achieve a result close to the correct stoichiometry, the results from bulk glucose (Fig. 5) are considered. Especially in the mass range from  $m/q = 27$  to  $36 \text{ u e}^{-1}$  and  $m/q = 41$  to  $48 \text{ u e}^{-1}$  significant overlap with water signals occurs. By assigning 33% of the signals to  $\text{C}_x\text{O}_y\text{H}_z$  fragments and 67% to water clusters the mass-ratio of 0.48 is achieved. This is very close to the theoretical value, but demonstrates the ambiguity of the correct determination of the stoichiometry at the moment. The excess of hydrogen is reduced by a factor of 1.3 if only the reinterpretation of the protonated water clusters was considered. However, the hydrogen content of still 25.15% is clearly too high, which is due to a wrong interpretation. Only with the assumption of  $\text{C}_x\text{O}_y\text{H}_z$  molecules and protonated water cluster the proportion is reduced by a factor of 5.2 and is about 4.28%. This overestimation of hydrogen can be attributed to residual hydrogen from the stainless-steel chamber and cannot be quantified exactly, but it is a very likely reason for the overestimation of the hydrogen content in APT measurements.

Correlative information sources have to be used to support the APT analysis.

Table 4: Calculated stoichiometry by different peak interpretation approaches.

Interpretation	mass ratio of glucose to water	water molecules per glucose molecule	Hydrogen (excess)
<b>Theoretical value</b>	<b>0.47</b>	<b>21.27</b>	<b>0 %</b>
(C <sub>x</sub> H <sub>y</sub> )	519.94	0.02	33.37 %
(C <sub>x</sub> H <sub>y</sub> ) + water peaks	1.40	7.17	25.15 %
(C <sub>x</sub> O <sub>y</sub> H <sub>z</sub> ) + water peaks	0.61	16.36	4.28 %
(C <sub>x</sub> O <sub>y</sub> H <sub>z</sub> ) + water peaks + overlap (C <sub>x</sub> O <sub>y</sub> H <sub>z</sub> )/H <sub>x</sub> O <sub>y</sub> - 25%/75 %+	0.46	21.68	4.60 %
(C <sub>x</sub> O <sub>y</sub> H <sub>z</sub> ) + water peaks + overlap (C <sub>x</sub> O <sub>y</sub> H <sub>z</sub> )/H <sub>x</sub> O <sub>y</sub> - 33%/67 %	0.48	20.81	4.60 %
(C <sub>x</sub> O <sub>y</sub> H <sub>z</sub> ) + water peaks + overlap (C <sub>x</sub> O <sub>y</sub> H <sub>z</sub> )/H <sub>x</sub> O <sub>y</sub> - 50%/50 %	0.52	19.10	4.49%
(C <sub>x</sub> O <sub>y</sub> H <sub>z</sub> ) + water peaks + overlap (C <sub>x</sub> O <sub>y</sub> H <sub>z</sub> )/H <sub>x</sub> O <sub>y</sub> - 67%/33 %	0.57	17.47	4.41 %
(C <sub>x</sub> O <sub>y</sub> H <sub>z</sub> ) + water peaks + overlap (C <sub>x</sub> O <sub>y</sub> H <sub>z</sub> )/H <sub>x</sub> O <sub>y</sub> - 75%/25 %	0.60	16.74	4.23%

### Three-dimensional structural analysis

The unique feature of APT is the reconstruction of the measured volume in 3D with a near atomic resolution. With the dominating evaporation of molecular species, the principal resolution limit is controlled by the size of the individual fragments. Desorption maps for selected molecules are depicted in Figure 6. While the distribution especially for larger protonated water clusters and glucose molecules is homogenous, local density changes for H<sub>3</sub>O<sup>+</sup> molecules are apparent. The occurrence of line features seems to be related to the laser incidence direction, as reported for pure water<sup>15</sup>.

In order to investigate the distribution of the solvated glucose molecules inside the water matrix the measurement data are reconstructed (see materials and methods section for more details) (Fig. 7). As majority component the signal from water and related water clusters is dominating. Small local density imperfections become visible which are related to the line structures in the desorption maps. Their origin has to be further investigated.

The distribution of HO, H<sub>2</sub>O and H<sub>3</sub>O molecules is probed by a cylinder, dimension of 20 x 100 nm, along the tip axis (Fig. 7a) and perpendicular to the axis (10 x 50 nm) (Fig. 7b). While the relative ratio of the water signals is relatively constant in tip direction, a change of relative abundance from H<sub>3</sub>O<sup>+</sup> to H<sub>2</sub>O<sup>+</sup> is recorded perpendicular to the tip axis. Since asymmetric with respect to the tip symmetry axis, again, the laser incidence direction can be made responsible. A suggested explanation would refer to the local heating of the sample due to the laser matter interaction leading to local change in tip curvature. As a

result, the effective field at the impact side of the laser is usually less than at the opposite. A lower field, appears to favour H<sub>2</sub>O molecules evaporating.

The glucose fragments show a homogeneous distribution in the 3D volume as observed in the desorption maps, which can be attributed to a very fast cooling of the sample, therefore no segregation zone is observed, caused by recrystallization of the water (Fig. 8).

By determining the radius of a sphere around an identified glucose molecule (masses  $m/q = 109 - 181 \text{ u e}^{-1}$ ) that includes a given number of nearest neighbouring (NN) glucose molecules, the distribution of the glucose inside the solution can be investigated. The number of included nearest neighbours' scales with the radius to the power 3.

$$NN_{\text{sphere}}(r) = \rho \cdot \frac{4}{3} \pi r^3 \quad (6)$$

Here  $\rho$  denotes the glucose density and is determined to  $0.5 \pm 0.1 \text{ molecules nm}^{-3}$ . As initially described, about 1.57 glucose molecules are expected per cubic nanometre. Taking the detection efficiency of 50% into account a partial fragmentation of a distinct number of molecules into smaller fragments, the value represents quite well the expected density distribution (Fig. 9).

In a saturated glucose solution, there are 21 water molecules available to solvate each glucose molecule. In order to visualize the hydrate shell around a solvated glucose molecule, two isosurfaces in a small volume of  $5 \times 5 \text{ nm}$ ,  $2 \times 3 \text{ nm}$ , and  $2 \times 2 \text{ nm}$  are plotted (Fig. 10 a-c). The position of evaporated glucose molecules is indicated by a second iso-surface. The surrounding water molecules/clusters are represented by the blue iso-surface with a water concentration around 67% (Fig. 10 a-c). The glucose molecules are surrounded by water molecules and homogeneously distributed inside the volume. The structural information of the individual molecules is of course not directly retrievable from the APT data. In Fig. 10c the DFT calculated structure is placed at the position of a detected glucose molecule implying a higher resolution than technically achieved. More advanced reconstruction algorithms combined with suitable simulation packages and DFT calculations might be able to derive the likely structure on an atomic scale from the purely molecular data sets, if of course other problematic sources of error are excluded. Any alteration of the sample structure by the freezing process has to be excluded. This is a necessary precondition like for all characterisation methods.

### Summary:

Equipped with the knowledge for pure water, we now have provided field evaporation measurements for a saturated glucose solution. Due to the low evaporation field of water the glucose molecules evaporated as large, partially intact, molecules. Smaller fragments overlap with the signals related to water. However, overlap and ambiguity of mass peaks leaves room for various approaches of chemical interpretation.

The APT data of pure bulk glucose differ strongly from the data of the glucose solution. The solid bulk material exhibits a higher fragmentation rate of the molecule rings as compared to the solution, probably caused by a high evaporation field of bulk glucose. Other causes may be different absorption coefficients and/or different binding conditions between bulk glucose and the solution. Nevertheless, with comparison of both measurements and the a-priori knowledge of the overall composition, a suitable interpretation of the mass peaks could be suggested.

A reasonable matching of the measured stoichiometry with the known one can be achieved by interpreting the overlap in the range  $m/q = 27-36 \text{ u e}^{-1}$  and  $m/q = 41-48 \text{ u e}^{-1}$  as a ratio of 33% of  $\text{C}_x\text{O}_y\text{H}_z$  to 67%  $(\text{H}_2\text{O})_{1-5}\text{H}^+$ . Resulting in a total mass fraction about 0.48, which agrees reasonably with the theoretical value of 0.47. However, there is also a certain overlap with many other peaks, which makes it exceedingly difficult to calculate them exactly, since the exact fractions are not yet known. A significant loss of oxygen or hydrogen cannot be excluded. The identification of glucose molecules within the matrix is only possible by the existence of nearly complete ionic molecules. These allow an unambiguous identification also locally in the volume space. The effective detection efficiency of the solute molecules results in only one molecule out of three identified with certainty.

What sounds like a disadvantage in comparison to the analysis of metals and semiconductors enables APT to retrieve distinguishable information's of solutes in aqueous solutions, if a sample modification by the freezing process can be excluded. A breakdown of the solutes into smaller molecular parts would render them invisible against the background of the matrix signals.

A better understanding of the measurement conditions is necessary with regard to the solutes, laser power, field strength, temperature. The change of molecular fragmentation in dependence of the local environment has to be understood. Furthermore, the influence of the evaporation of large clusters on the accuracy of the reconstruction is not known and must be examined and in the best case attributed by an improved reconstruction algorithm. Nevertheless, the possibility to detect certain molecules within an aqueous solution, opens the opportunity to inject suitable markers and biological molecules, to study their distribution in various reactions in 3D with sub-nanometre resolution.

## Materials And Methods

**Materials.** As sample material, glucose (EP, BP, JP, USP testing specifications, anhydrous, from Sigma Aldrich) was used. For the supersaturated glucose solution, water which was ionized and filtered through a Milli-Q system (Millipore) was used. By careful cleaning of all materials and the usage of pure constituents overlapping in the ToF-mass spectrum from impurities is avoided, since this would lead to an even more challenging peak analysis. To create a saturated glucose solution, glucose was added to water until the solubility limit was reached a sediment formed on the bottom of the test glass. Tungsten was chosen as a substrate material. To create a rough and reproducible surface we followed the approach described in <sup>15</sup>. For the Lift-Out process, tungsten wires were first etched in 2mol NaOH

solution by applying an AC voltage. Afterwards the tip was cut down to a flat post by using the Focused Ion Beam (FIB).

**Freezing Process:** As reported in previous work <sup>15</sup>, a solution droplet was dipped with a micro pipette on a precooled tungsten post, which is located within a liquid Nitrogen bath to create droplets on top (60-250  $\mu\text{m}$ ). Subsequently, the sample holder is transferred as fast as possible into the cooled body of the modified transfer shuttle VCT500 from Leica ( $T = -184\text{ }^\circ\text{C}$ ) and pumped to a pressure of  $6 \cdot 10^{-1}$  mbar. Shortly after, the shuttle is attached to the high vacuum-coater (Leica EM ACE600), to carry out a freeze etching process to remove ice crystals that were formed by the contact of the sample with air. By heating up the sample very precisely to a temperature of  $-90\text{ }^\circ\text{C}$  and a pressure of  $9 \cdot 10^{-7}$  mbar for 30 minutes, a sublimation process from solid ice to vaporous ice occurs, which is necessary for the controlled removal of condensed ice from the sample. In addition, the Leica EM ACE600 high vacuum coater is useful to improve the vacuum condition inside the shuttle ( $10^{-4}$  mbar), which is necessary to enable the transfer of the sample into the Focused-Ion-Beam (FIB) Microscope.

**Cryo FIB Preparation:** In order to prepare cryogenic samples in the FIB (FEI Scios) into a nano-shaped tips with an apex radius less than 100 nm, which is required for the APT method, the FIB has to be equipped with a custom made cryo-stage, which is cooled down to a temperature of  $-150\text{ }^\circ\text{C}$  by copper bands connected to a  $\text{N}_2$  Dewar attached to the microscope. The sample itself is transferred into the microscope using a dedicated VCT500 load lock. This load lock was mounted at the back side of the SEM to the port intentional designed for the STEM detector, which allows an easy sample transport into the cryo-stage. Cold surfaces act as a trap for surrounding gas or molecules. To avoid re-deposition of material onto the shaped sample, a cryo-shield was installed in the chamber additionally.

**Cryo APT specimen preparation.** SEM imaging was typically performed with low energy (5 kV 25 pA) to prevent melting of the sample by electron bombardment. For the milling process, the sample has been tilted to  $52^\circ$ , which aligns it vertically towards the ion beam. A circular ring pattern was used for azimuthal milling. Initial milling steps were performed at 30 kV acceleration voltage and an ion beam current of 50 nA until the tungsten substrate became visible again. After reaching a radius of 30  $\mu\text{m}$ , the beam current is gradually reduced with decreasing radius, down up to an inner ring diameter of 300 nm and a beam current of 0.1 pA. The thinning process is monitored by taking snapshots using the electron beam. The shaping process continues until a very sharp tip with a radius  $<100$  nm (Fig. 11 a-d) is obtained. The finished tip is then transferred back into the shuttle and can be attached to the APT. To our experience reproducible measurements with sufficient throughput require a high aspect ratio of the tip. A tip length of several tens of  $\mu\text{m}$  is desirable. The preparation way for a cryo-APT specimen is depicted in Figure 11 a-d.

**APT specimen preparation:** The glucose crystals were glued with a silver glue (RS pro silver conductive glue) on a SEM grid. For better conductivity and to minimize the charging effect, 50 nm copper was sputtered on top with the high vacuum-coater (Leica EM ACE600) by glow discharge sputtering. Deposition parameter were 60 mA current and an Ar pressure of  $2.0 \cdot 10^{-2}$  mbar and an effective sputter

rate of 0.15 nm/s. Afterwards a normal APT specimen Lift-Out method was used<sup>47</sup> (see Fig. 11 e-l) followed by a shaping process.

**APT Measurements:** Using a custom-made atom probe<sup>57</sup> equipped with NOPA to continuously change wavelength in the range between 350 and 900 nm, measurements were conducted with a laser wavelength of 355 nm. The pulse length was chosen to 250 fs. The spot size in the focus point accounts to a diameter of 50 microns. As detector a 120 mm diameter delay line detector with chevron MCP setup with an open area ratio (OAR) of 50% is used. The flight length accounts to 130 mm and the effective half angle of the field of view to 38°. The system is equipped with a custom made cryo-transfer port to accept a standard VCT500 from Leica for the transfer of cryogenic samples. The exchange of the sample in the buffer chamber uses a PEEK isolated storage position to avoid melting of the sample during transfer. The obtained datasets were analysed and reconstructed using the Scito<sup>58</sup> software package.

**Reconstruction:** The volume was reconstructed using a geometrical reconstruction algorithm<sup>59</sup> based on the original point projection method by Bas et al.<sup>60</sup>. SEM pictures were used to determine the initial radius of the tip. This radius and the taper angle were used to optimize the calculated evaporation-field curve. All necessary parameters, like field and image compression factors were determined in earlier experiments. For the z-axis reconstruction the respective densities of each molecular fragment were calculated to achieve the correct density values after splitting of the assumed molecule in its atomic constituents.

**DFT Calculation:** To explain the obtained fragments, we performed calculations with density functional theory (DFT) to study the stability of several ions that can explain the observed m/q ratios. All energies reported were obtained with the functional BP86<sup>61,62</sup> and the basis set def2-SVP<sup>63,64</sup> with D3 dispersion correction<sup>65</sup>. The calculations were performed with Turbomole<sup>66,67</sup> run through ChemShell<sup>68,69</sup>. Harmonic frequency calculations confirmed the structures as minima on the potential energy surface. The energies include the harmonic zero-point vibrational energy. All calculations were performed in the absence of an external electric field. Partial charges were analysed using natural bond orbitals (NBO)<sup>70</sup>.

The starting geometry for the calculation was an  $\alpha$ -D-glucopyranose molecule. The geometry at m/q = 181 u e<sup>-1</sup> is equivalent to a glucose molecule with an additional proton. The proton was attached to different atoms in the molecule in order to find the most likely binding site. The geometry with the lowest energy found the additional H to be bound between two oxygen atoms.

By adding a hydronium cation to glucose, the geometry for m/q = 199 u e<sup>-1</sup> is obtained. The hydronium can approach the glucose molecule from several different directions, thereby resulting in a multitude of possible molecular structures. The structure with the lowest energy is chosen as the most likely and serves as the starting geometry for the calculation of the next structure at m/q = 217 u e<sup>-1</sup> in which case H<sub>2</sub>O is added. The structure of glucose+(H<sub>2</sub>O)<sub>16</sub>H<sup>+</sup> is the result of an MD simulation with GFN-xTB<sup>71</sup>, followed by an energy minimization with BP86/def2-SVP.

The simulation of structures at mass-to-charge state ratios of  $m/q = 163 \text{ u e}^{-1}$  and below progressed in a similar fashion. One hydrogen and an OH group were removed from the glucose molecule. Several different possible combinations were investigated. Each structure with the lowest energy served as starting point for the structure with the next lower mass-to-charge state ratio.

## Declarations

### Acknowledgements:

The authors gratefully acknowledge funding by the German Research Foundation (Deutsche Forschungsgemeinschaft, DFG)-Project-ID 358283783 - SFB 1333.

### Contributions:

P.S conceived the idea of the research plan and the experiment. T.M.S and J.O produced the samples and performed the experiment with APT, supervised by P.S. Data Analysis was carried out by T.M.S, P.S and E.M.W. E.H and H.S helped with FIB sample processing. R.L programmed analysis tools for data evaluation. C.A.D and J.K carried out DFT calculations. B.G and G.S reviewed the manuscript. T.M.S and P.S wrote the paper with input from all authors.

### Competing interests:

The authors declare no competing interests.

## References

1. Gault, B., Moody, M. P., Cairney, J. M. & Ringer, S. P. *Atom Probe Microscopy*. (Springer).
2. Miller, M. & Forbes, R. *Atom-Probe Tomography - The Local Electrode Atom Probe* (Springer US, 2014).
3. Kelly, T. F. & Larson, D. J. Atom Probe Tomography 2012. *Annu. Rev. Mater. Res.* **42**, 1–31 (2012).
4. Lefebvre, Ulrikson, W., Vurpillot, F., Sauvage, X., *Atom Probe Tomography*. (Elsevier, Academic Press, 2016).
5. Schmitz, G. Nanoanalysis by Atom Probe Tomography. in *Nanotechnology* (Wiley-VCH Verlag GmbH & Co. KGaA, 2010). doi:10.1002/9783527628155.nanotech069.
6. Stender, P., Heil, T., Kohl, H. & Schmitz, G. Quantitative comparison of energy-filtering transmission electron microscopy and atom probe tomography. *Ultramicroscopy* **109**, (2009).
7. Rusitzka, K. A. K. *et al.* A near atomic-scale view at the composition of amyloid-beta fibrils by atom probe tomography. *Sci. Rep.* **8**, 17615 (2018).
8. Kelly, T. F., Nishikawa, O., Panitz, J. A. & Prosa, T. J. Prospects for Nanobiology with Atom-Probe Tomography. *MRS Bull.* **34**, 744–750 (2009).

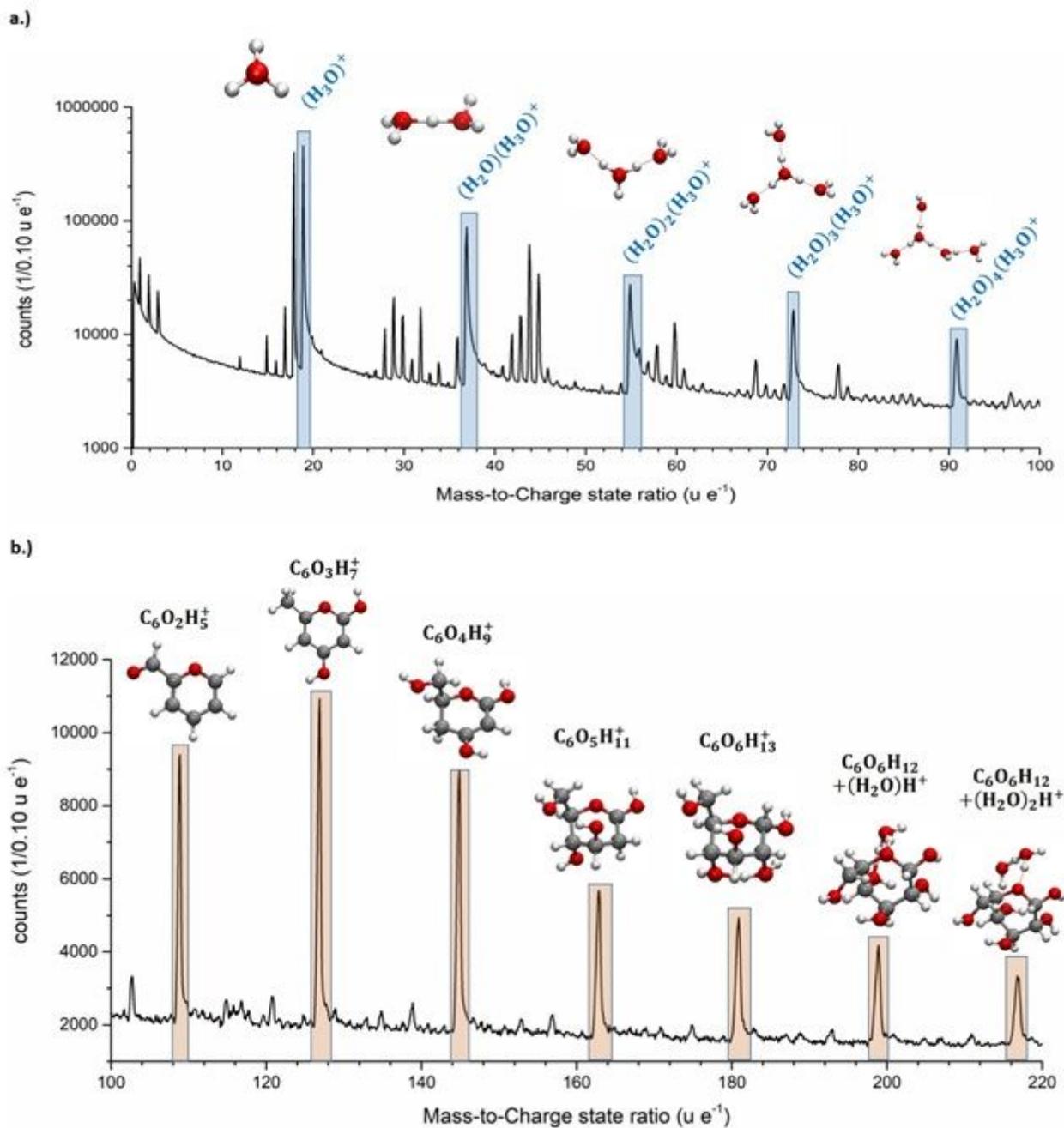
9. Adineh, V. R., Marceau, R. K. W., Velkov, T., Li, J. & Fu, J. Near-Atomic Three-Dimensional Mapping for Site-Specific Chemistry of 'Superbugs'. *Nano Lett.* **16**, 7113–7120 (2016).
10. Perea, D. E. *et al.* Tomographic mapping of the nanoscale water-filled pore structure in corroded borosilicate glass. doi:10.1038/s41529-020-0110-5.
11. Schreiber, D. K., Perea, D. E., Ryan, J. V., Evans, J. E. & Vienna, J. D. A method for site-specific and cryogenic specimen fabrication of liquid/solid interfaces for atom probe tomography. *Ultramicroscopy* **194**, 89–99 (2018).
12. Qiu, S. *et al.* Three-Dimensional Chemical Mapping of a Single Protein in the Hydrated State with Atom Probe Tomography. *Anal. Chem.* **92**, 5168–5177 (2020).
13. Qiu, S. *et al.* Graphene encapsulation enabled high-throughput atom probe tomography of liquid specimens. *Ultramicroscopy* **216**, (2020).
14. Qiu, S. *et al.* Direct Imaging of Liquid-Nanoparticle Interfaces with Atom Probe Tomography. *J. Phys. Chem. C* **124**, 19389–19395 (2020).
15. Schwarz, T. M. *et al.* Field evaporation and atom probe tomography of pure water tips. *Sci. Rep.* 1–14 (2020) doi:10.1038/s41598-020-77130-x.
16. El-Zoka, A. A. *et al.* Enabling near-atomic-scale analysis of frozen water. (2020).
17. Stintz, A. & Panitz, J. A. Cluster ion formation in isothermal ramped field-desorption of amorphous water ice from metal surfaces. *Surf. Sci.* **296**, 75–86 (1993).
18. Tsong, T. T. & Liou, Y. Cluster-ion formation in pulsed-laser-stimulated field desorption of condensed materials. *Phys. Rev. B* **32**, 4340–4357 (1985).
19. Beckey, H. D. Massenspektrometrische untersuchungen über ionen-molekülreaktionen und über die assoziation des wassers mit hilfe einer feldemissions-ionenquelle. *Zeitschrift für Naturforsch. - Sect. A J. Phys. Sci.* **15**, 822–827 (1960).
20. Karahka, M. & Kreuzer, H. J. Water whiskers in high electric fields. in *Physical Chemistry Chemical Physics* vol. 13 11027–11033 (The Royal Society of Chemistry, 2011).
21. Dirks, J. The interaction of synchrotron light with water adsorbed on a Ag-field emitter in the presence of a high electric field. *J. Vac. Sci. Technol. B Microelectron. Nanom. Struct.* **10**, 231 (1992).
22. Marko, M., Hsieh, C., Moberlychan, W., Mannella, C. A. & Frank, J. Focused ion beam milling of vitreous water: Prospects for an alternative to cryo-ultramicrotomy of frozen-hydrated biological samples. *J. Microsc.* **222**, 42–47 (2006).
23. Medeiros, J. M. *et al.* Robust workflow and instrumentation for cryo-focused ion beam milling of samples for electron cryotomography. *Ultramicroscopy* **190**, 1–11 (2018).
24. Stoffers, A., Oberdorfer, C. & Schmitz, G. Controlled field evaporation of fluorinated self-assembled monolayers. *Langmuir* **28**, 56–59 (2012).
25. Gruber, M., Oberdorfer, C., Stender, P. & Schmitz, G. Laser-assisted atom probe analysis of sol-gel silica layers. *Ultramicroscopy* **109**, (2009).

26. Perea, D. E. *et al.* Atom probe tomographic mapping directly reveals the atomic distribution of phosphorus in resin embedded ferritin. *Sci. Rep.* **6**, 1–9 (2016).
27. Dirks, J. The interaction of synchrotron light with water adsorbed on a Ag-field emitter in the presence of a high electric field. *J. Vac. Sci. Technol. B Microelectron. Nanom. Struct.* **10**, 231 (1992).
28. Anway, A. R. Field ionization of water. *J. Chem. Phys.* **50**, 2012–2021 (1969).
29. Karahka, M., Xia, Y. & Kreuzer, H. J. The mystery of missing species in atom probe tomography of composite materials. *Appl. Phys. Lett.* **107**, (2015).
30. Silaeva, E. P., Karahka, M. & Kreuzer, H. J. Atom Probe Tomography and field evaporation of insulators and semiconductors: Theoretical issues. *Current Opinion in Solid State and Materials Science* vol. 17 211–216 (2013).
31. Gault, B. *et al.* Behavior of molecules and molecular ions near a field emitter. *New J. Phys.* **18**, 033031 (2016).
32. Yao, L., Gault, B., Cairney, J. M. & Ringer, S. P. On the multiplicity of field evaporation events in atom probe: A new dimension to the analysis of mass spectra. *Philos. Mag. Lett.* **90**, 121–129 (2010).
33. Peng, Z. *et al.* On the detection of multiple events in atom probe tomography. *Ultramicroscopy* **189**, 54–60 (2018).
34. Thuvander, M. *et al.* Reduction of multiple hits in atom probe tomography Reduction of multiple hits in atom probe tomography. *Ultramicroscopy* 81–85 (2013) doi:10.1016/j.ultramic.2012.12.005.
35. Meisenkothen, F., Steel, E. B., Prosa, T. J., Henry, K. T. & Prakash Kolli, R. Effects of detector dead-time on quantitative analyses involving boron and multi-hit detection events in atom probe tomography. *Ultramicroscopy* **159**, 101–111 (2015).
36. Saxey, D. W. Correlated ion analysis and the interpretation of atom probe mass spectra. *Ultramicroscopy* **111**, 473–479 (2011).
37. Hans, M. & Schneider, J. M. Electric field strength-dependent accuracy of TiAlN thin film composition measurements by laser-assisted atom probe tomography. *New J. Phys* **22**, 33036 (2020).
38. Blum, I. *et al.* Dissociation Dynamics of Molecular Ions in High dc Electric Field. *J. Phys. Chem. A* **120**, 3654–3662 (2016).
39. Müller, M., Saxey, D. W., Smith, G. D. W. & Gault, B. Some aspects of the field evaporation behaviour of GaSb. *Ultramicroscopy* **111**, 487–492 (2011).
40. Müller, M., Gault, B., Smith, G. D. W. & Grovenor, C. R. M. Accuracy of pulsed laser atom probe tomography for compound semiconductor analysis. in *Journal of Physics: Conference Series* vol. 326 012031 (Institute of Physics Publishing, 2011).
41. Mancini, L. *et al.* Composition of wide bandgap semiconductor materials and nanostructures measured by atom probe tomography and its dependence on the surface electric field. *J. Phys. Chem. C* **118**, 24136–24151 (2014).
42. Watson, J. D. & Crick, F. H. C. Molecular structure of nucleic acids: A structure for deoxyribose nucleic acid. *Nature* **248**, 765 (1974).

43. Robyt, J. F. *Essentials of Carbohydrate Chemistry*. (Springer, 1998).
44. Timp, W. & Timp, G. Beyond mass spectrometry, the next step in proteomics. *Sci. Adv.* **6**, eaax8978 (2020).
45. Tsong, T.T., *Atom-Probe Field Ion Microscopy: Field Ion Emission, and Surfaces and Interfaces at atomic resolution* Tien T. Tsong (Cambridge University Press) 1990.
46. A Schmidt, V. W., Müller, E. W., auch Zitat, vgl, Inghram R Gomer, M. G. & Chem Phys, J. *Massenspektrometrische Untersuchung der Feldionisation von Wasserdampf an Spitzen aus Wolfram, Platin und Iridium 1 ' 2. Z. Naturforschg* vol. 19 (1964).
47. Prosa, T. J. & Larson, D. J. Modern Focused-Ion-Beam-Based Site-Specific Specimen Preparation for Atom Probe Tomography. in *Microscopy and Microanalysis* vol. 23 194–209 (Cambridge University Press, 2017).
48. Nishikawa, O., Taniguchi, M. & Ikai, A. Atomic Level Analysis of Amino Acids by the Scanning Atom Probe. *MRS Proc.* **1231**, 1231-NN06-09 (2009).
49. Nishikawa, O. & Taniguchi, M. Atomic level analysis of carbon fibers by the scanning atom probe. *Surf. Interface Anal.* **46**, 1231–1235 (2014).
50. Nishikawa, O. & Taniguchi, M. Quantitative evaluation of carbon nanotubes by the scanning atom probe. *J. Vac. Sci. Technol. A Vacuum, Surfaces, Film.* **27**, 1076–1079 (2009).
51. Nishikawa, O., Taniguchi, M. & Ikai, A. Atomic level analysis of biomolecules by the scanning atom probe. *Appl. Surf. Sci.* **256**, 1210–1213 (2009).
52. Werner, M., Baars, A., Werner, F., Eder, C. & Delgado, A. Thermal conductivity of aqueous sugar solutions under high pressure. *Int. J. Thermophys.* **28**, 1161–1180 (2007).
53. Krokida, M. K., Panagiotou, N. M., Maroulis, Z. B. & Saravacos, G. D. THERMAL CONDUCTIVITY: LITERATURE DATA COMPILATION FOR FOODSTUFFS. *Int. J. Food Prop.* **4**, 111–137 (2001).
54. Andersson, O. & Suga, H. Thermal conductivity of low-density amorphous ice. *Solid State Commun.* **91**, 985–988 (1994).
55. Thanatuksorn, P., Kajiwara, K., Murase, N. & Franks, F. Freeze-thaw behaviour of aqueous glucose solutions - The crystallisation of cubic ice. *Phys. Chem. Chem. Phys.* **10**, 5452–5458 (2008).
56. Olaf Köhl. *Organische Chemie für Biochemiker, Lebenswissenschaftler, Mediziner, Pharmazeuten*. (Wiley-VCH, 2012).
57. Schlesiger, R. *et al.* Design of a laser-assisted tomographic atom probe at Münster University. *Rev. Sci. Instrum.* **81**, 043703 (2010).
58. Stender, P., Balla, I. Scito - APT Reconstruction Software., [www.inspico.eu](http://www.inspico.eu)
59. Jeske, T. & Schmitz, G. Nanoscale analysis of the early interreaction stages in Al/Ni. *Scr. Mater.* **45**, 555–560 (2001).
60. Bas, P., Bostel, A., Deconihout, B. & Blavette, D. A general protocol for the reconstruction of 3D atom probe data. *Appl. Surf. Sci.* **87–88**, 298–304 (1995).

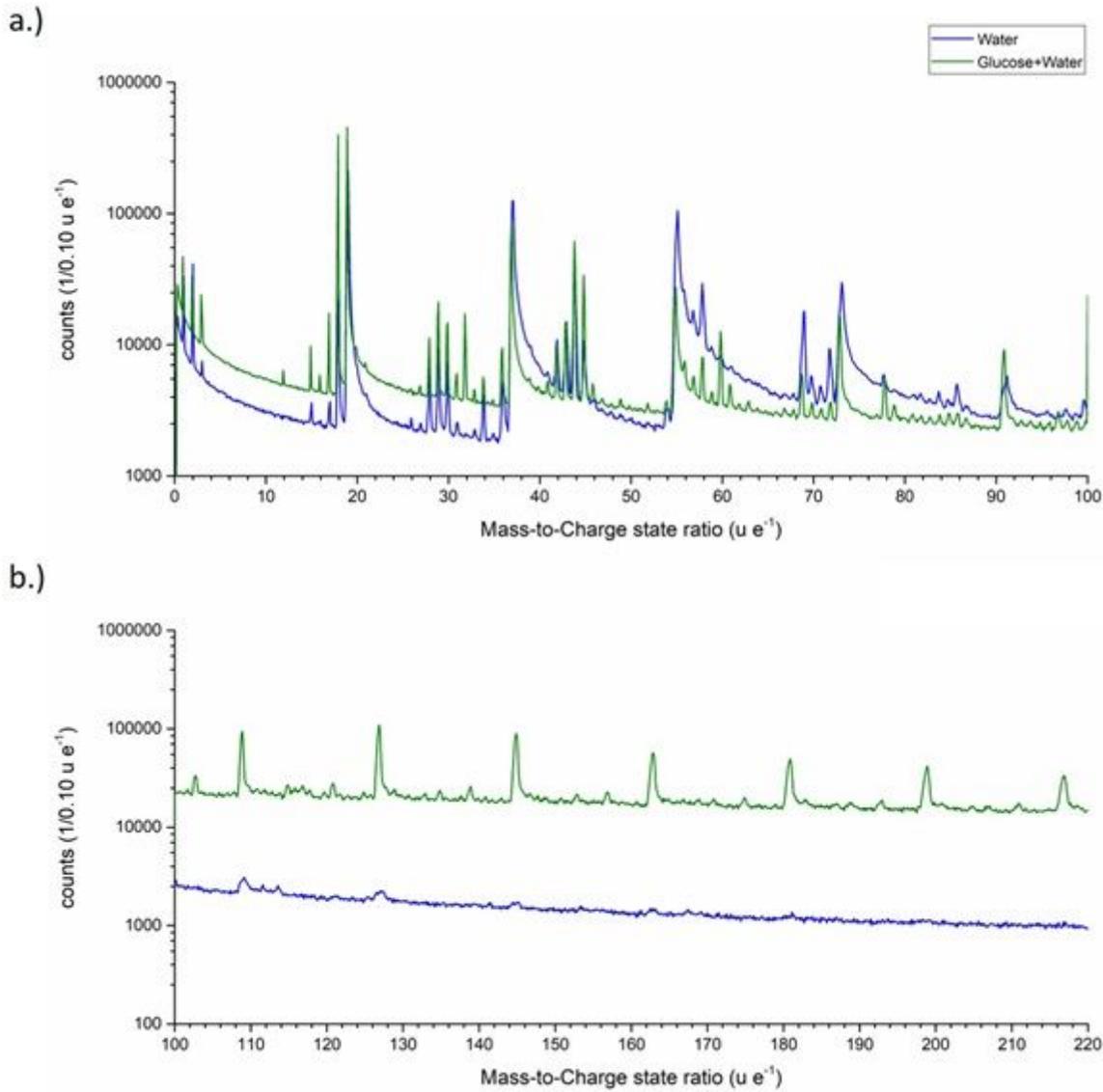
61. Becke, A. D. Density-functional exchange-energy approximation with correct asymptotic behavior. *Phys. Rev. A* **38**, 3098–3100 (1988).
62. Perdew, J. P. Density-functional approximation for the correlation energy of the inhomogeneous electron gas. *Phys. Rev. B* **33**, 8822–8824 (1986).
63. Eichkorn, K., Weigend, F., Treutler, O. & Ahlrichs, R. Auxiliary basis sets for main row atoms and transition metals and their use to approximate Coulomb potentials. *Theor. Chem. Acc.* **97**, 119–124 (1997).
64. Weigend, F. & Ahlrichs, R. Balanced basis sets of split valence, triple zeta valence and quadruple zeta valence quality for H to Rn: Design and assessment of accuracy. *Phys. Chem. Chem. Phys.* **7**, 3297–3305 (2005).
65. Grimme, S., Antony, J., Ehrlich, S. & Krieg, H. A consistent and accurate ab initio parametrization of density functional dispersion correction (DFT-D) for the 94 elements H-Pu. *J. Chem. Phys.* **132**, 134101 (2010).
66. Ahlrichs, R., Bär, M., Häser, M., Horn, H. & Kölmel, C. Electronic structure calculations on workstation computers: The program system turbomole. *Chem. Phys. Lett.* **162**, 165–169 (1989).
67. University of Karlsruhe, and, Forschungszentrum Karlsruhe, G. Turbomole.
68. Sherwood, P. *et al.* QUASI: A general purpose implementation of the QM/MM approach and its application to problems in catalysis. *J. Mol. Struct. THEOCHEM* **632**, 1–28 (2003).
69. Metz, S., Kästner, J., Sokol, A. A., Keal, T. W. & Sherwood, P. ChemShell-a modular software package for QM/MM simulations. *Wiley Interdiscip. Rev. Comput. Mol. Sci.* **4**, 101–110 (2014).
70. Reed, A. E., Weinstock, R. B. & Weinhold, F. Natural population analysis. *J. Chem. Phys.* **83**, 735–746 (1985).
71. Bannwarth, C., Ehlert, S. & Grimme, S. GFN2-xTB - An Accurate and Broadly Parametrized Self-Consistent Tight-Binding Quantum Chemical Method with Multipole Electrostatics and Density-Dependent Dispersion Contributions. *J. Chem. Theory Comput.* **15**, 1652–1671 (2019).

## Figures



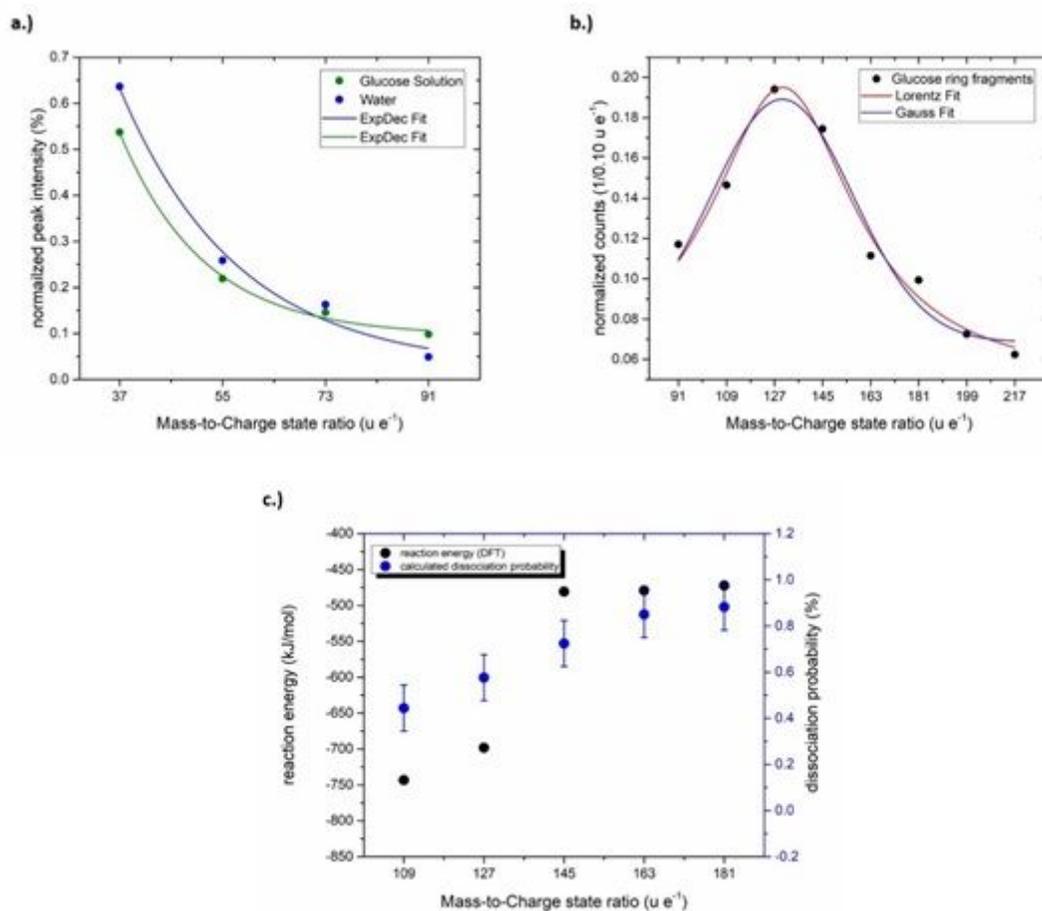
**Figure 1**

Mass spectrum of a supersaturated glucose solution with a mass-to-charge state ratio from 0-100  $u e^{-1}$  as logarithmic plot. The protonated water molecules/clusters are highlighted and the respective molecular structure of  $(H_2O)_nH^+$  with  $n = 1-5$  are shown. b.) Mass spectrum of a supersaturated glucose solution with a mass-to-charge state ratio from 100-220  $u e^{-1}$  as linear plot. The peaks stemming from glucose from  $m/q = 91$  up to 217  $u e^{-1}$  are marked and the respective molecular structures are shown.



**Figure 2**

Comparison of the mass spectra of pure water and a saturated glucose solution. In a.) from the mass-to-charge state ratio of  $m/q = 0-100 \text{ u e}^{-1}$  and in b.) from  $m/q = 100-220 \text{ u e}^{-1}$  as logarithmic plot. In b.) the mass spectrum of the supersaturated glucose solution was shifted upwards by a factor of 10 to make both spectra more visible.



**Figure 3**

In a.) the relative peak intensity of the water molecules  $(H_2O)_nH^+$  with  $n = 1-5$  from a pure water measurement and the glucose solution, in b.) the peak distribution of the peaks of the glucose ring from  $m/q = 91$  up to  $217 u e^{-1}$  and in c.) the calculated probabilities and the reaction energy calculated by DFT to remove one  $HO+H$  from a glucose ring-/fragment are plotted against the respective mass-to-charge state ratio.

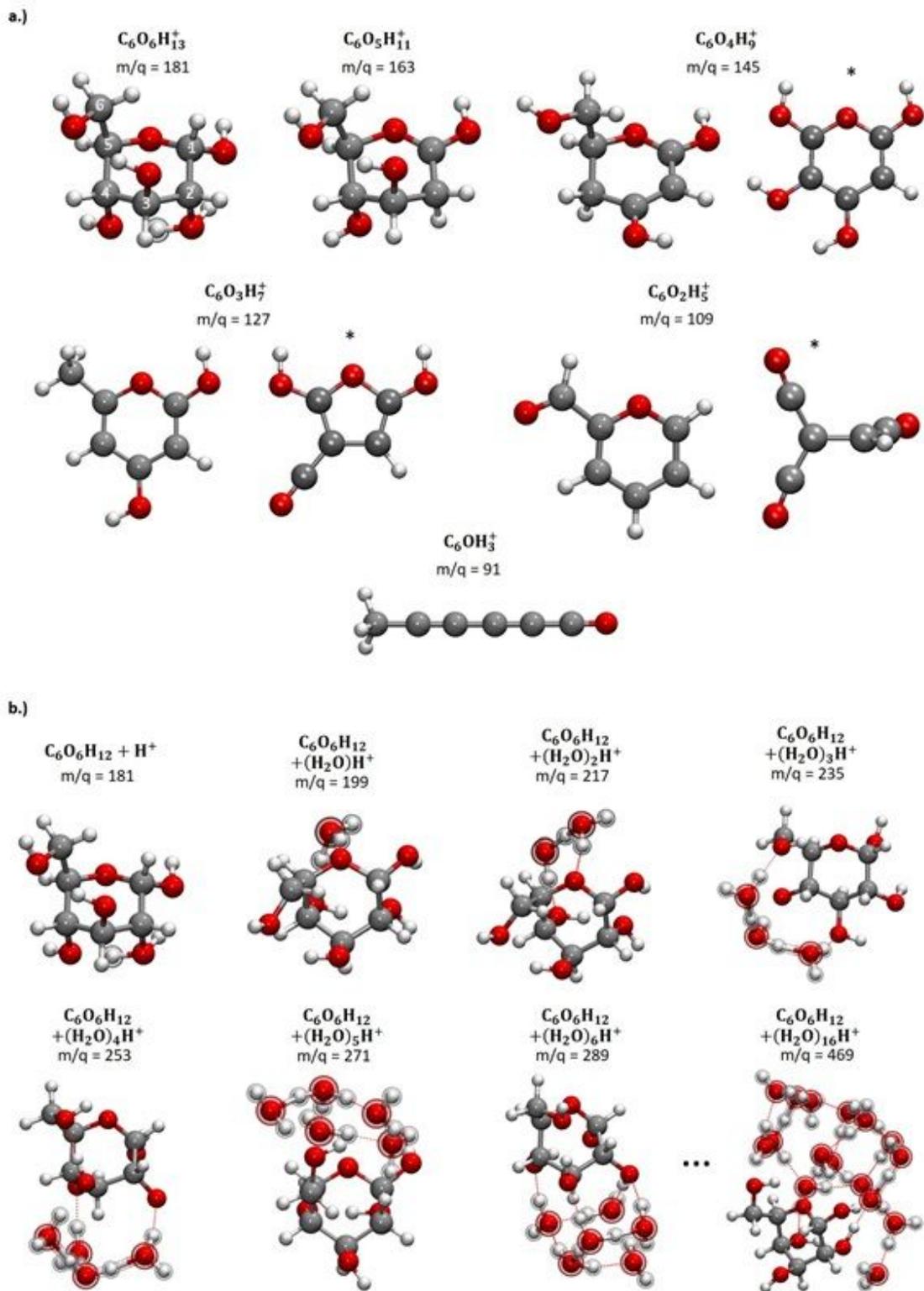
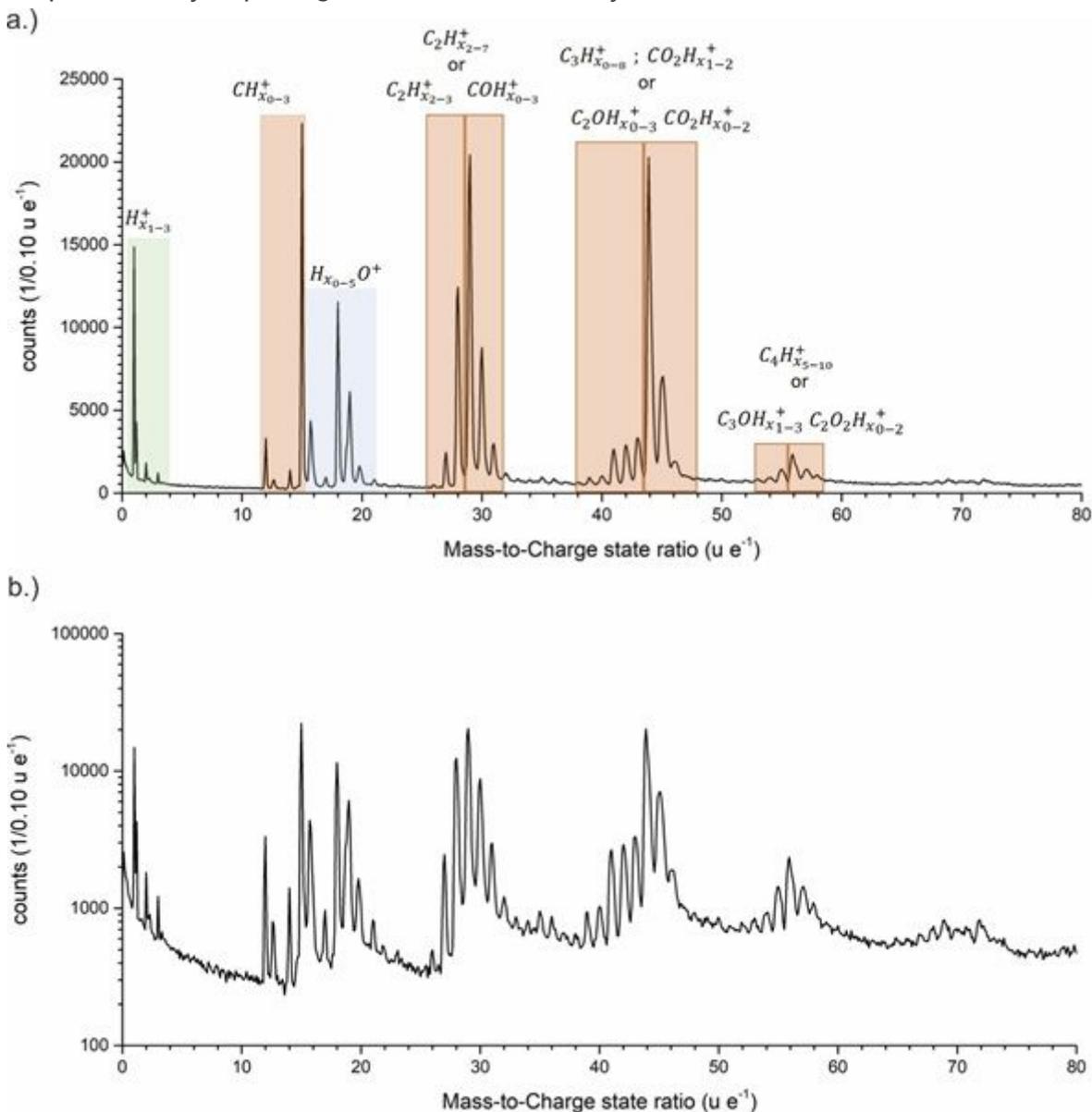


Figure 4

a.) Molecular structures corresponding to the mass-to-charge state ratios of  $m/q = 181 - n \cdot 18 \text{ u e}^{-1}$  with  $n = 0$  to 5 until  $m/q = 91 \text{ u e}^{-1}$ . All structures shown here represent the isomers with the lowest energy. Alternative molecules at  $m/q = 145$ , 127 and 109  $\text{u e}^{-1}$  are marked with (\*). However, they are deemed unlikely, because they would require the simultaneous removal of one carbon along with several H atoms.

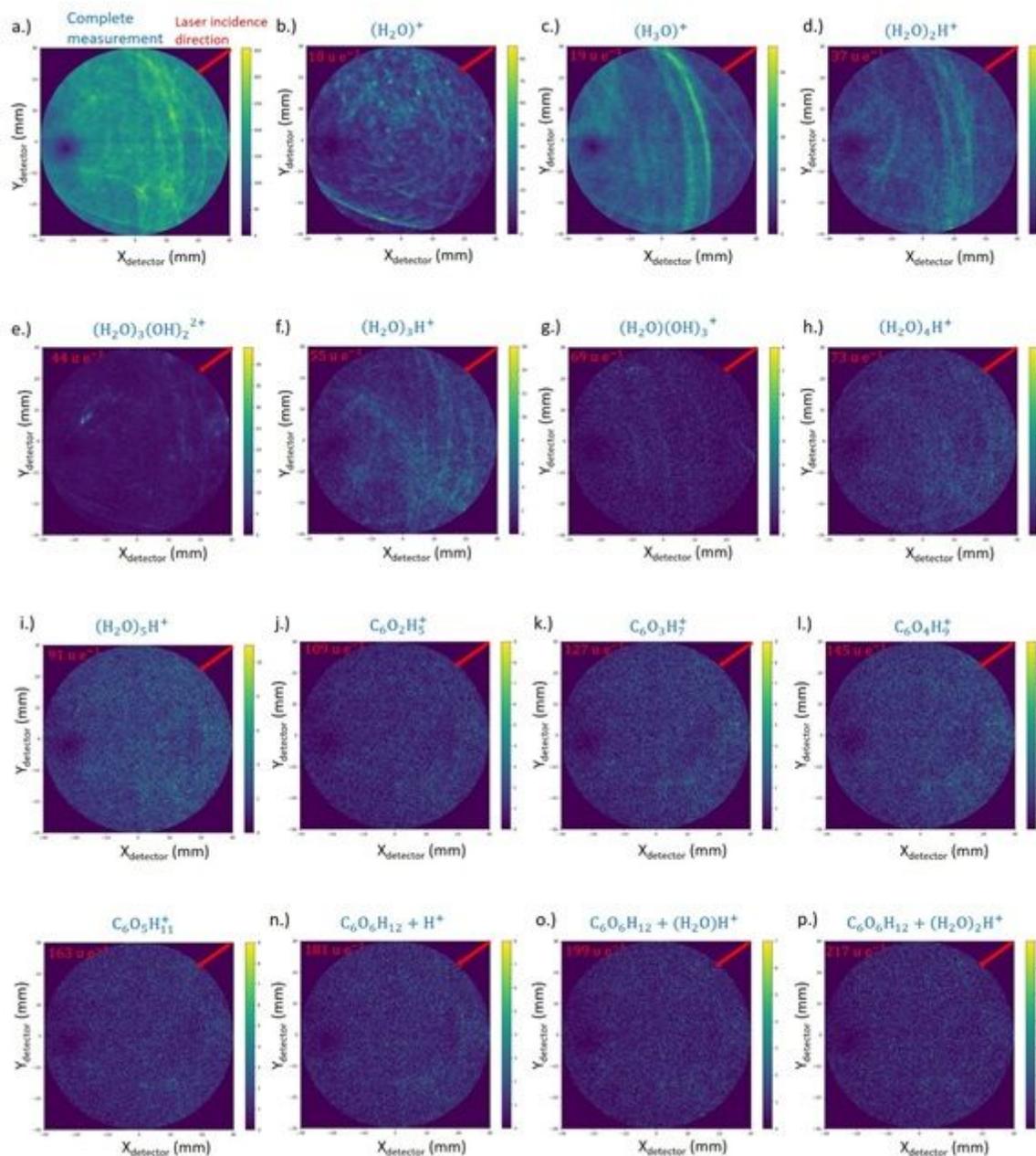
b.) Molecular clusters formed by adding cations to glucose: left to right: glucose molecule ( $\alpha$ -D-

glucopyranose) with addition of  $H^+$ ,  $H_3O^+$  and  $H_5O_2^+$  until  $(H_2O)_6H^+$ , which form water clusters bound to glucose via hydrogen bonds (dotted lines). The structure shown for  $C_6O_6H_{12} + (H_2O)_{16}H^+$  is the result of an MD simulation with GFN-xTB, followed by an energy minimization with BP86/def2-SVP. It is obvious that the glucose molecule locates on the surface of a water cluster. The additional atoms are emphasized by depicting them in a different style.



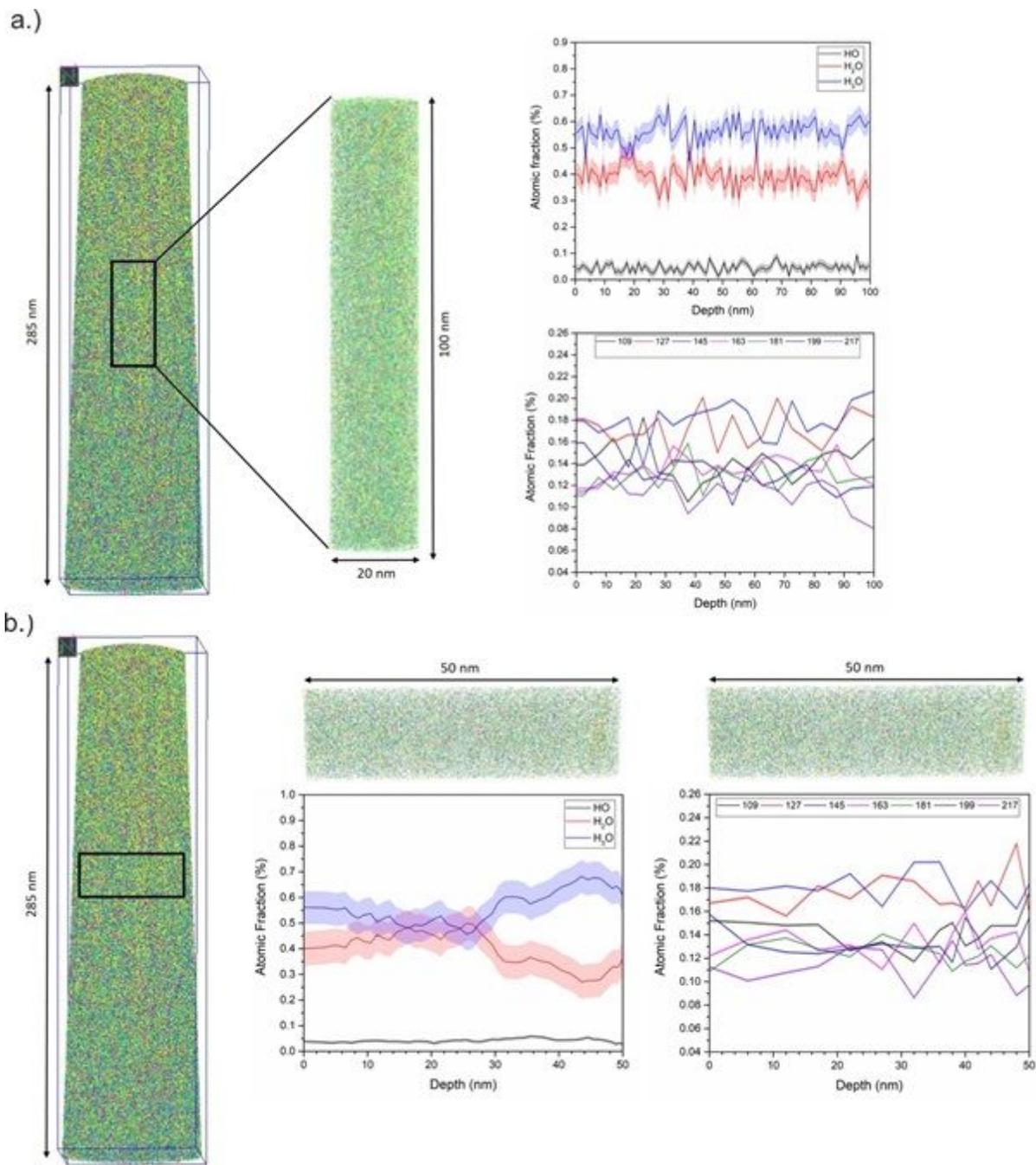
**Figure 5**

The mass spectra of the glucose bulk specimen. In a.) from the mass-to-charge state ratio of  $m/q = 0-80$  u e<sup>-1</sup> as linear in b.) from  $m/q = 0-80$  u e<sup>-1</sup> as logarithmic plot.



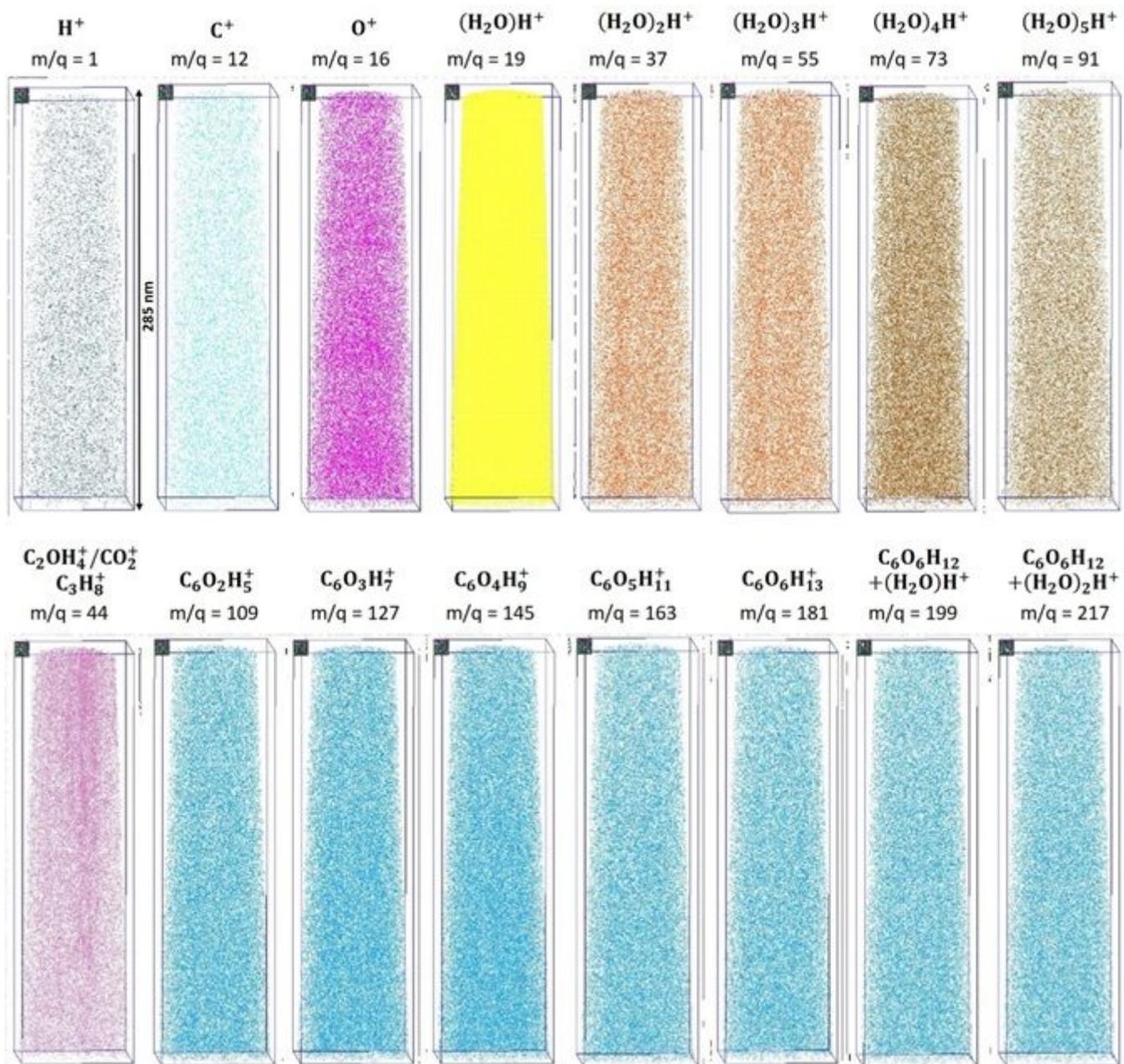
**Figure 6**

Desorption maps; a.) all events b-o.) desorption maps of different protonated water and glucose fragments ordered by increasing masses and size. Laser incidence direction is marked with a red arrow.



**Figure 7**

a.) Representative 3D reconstruction of a measured specimen with a total length of 285 nm. A cylinder with a dimension of 50 x 50 x 100 nm was selected vertical and in b.) horizontal to the tip from the total volume in order to analyse the composition profile of the H<sub>x</sub>O molecules with x = 1-3 and the glucose fragments from m/q = 91 until 217 u e-1.



**Figure 8**

Individual maps of the reconstructed volume for different molecules are shown.

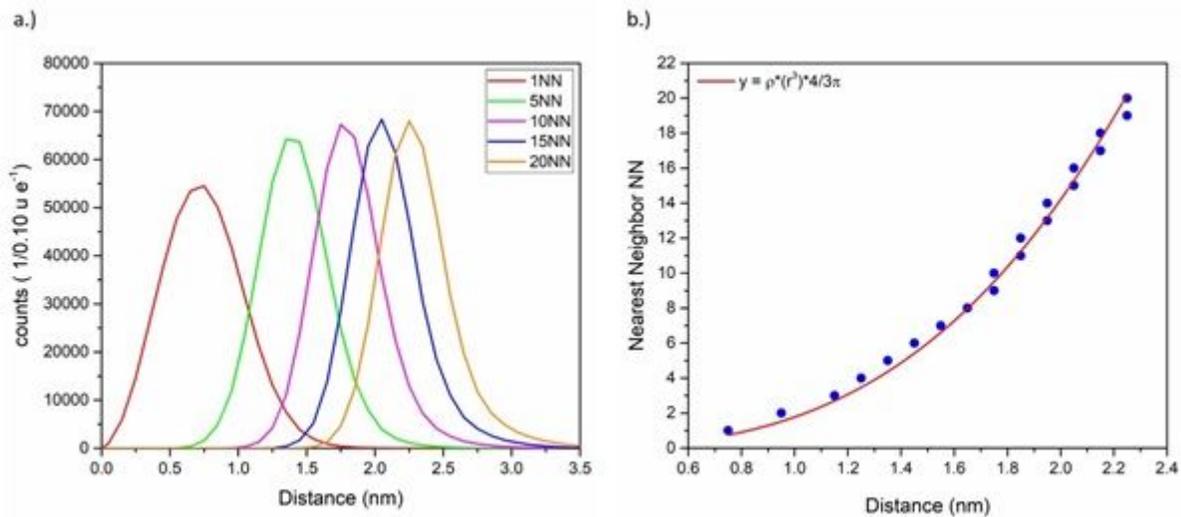
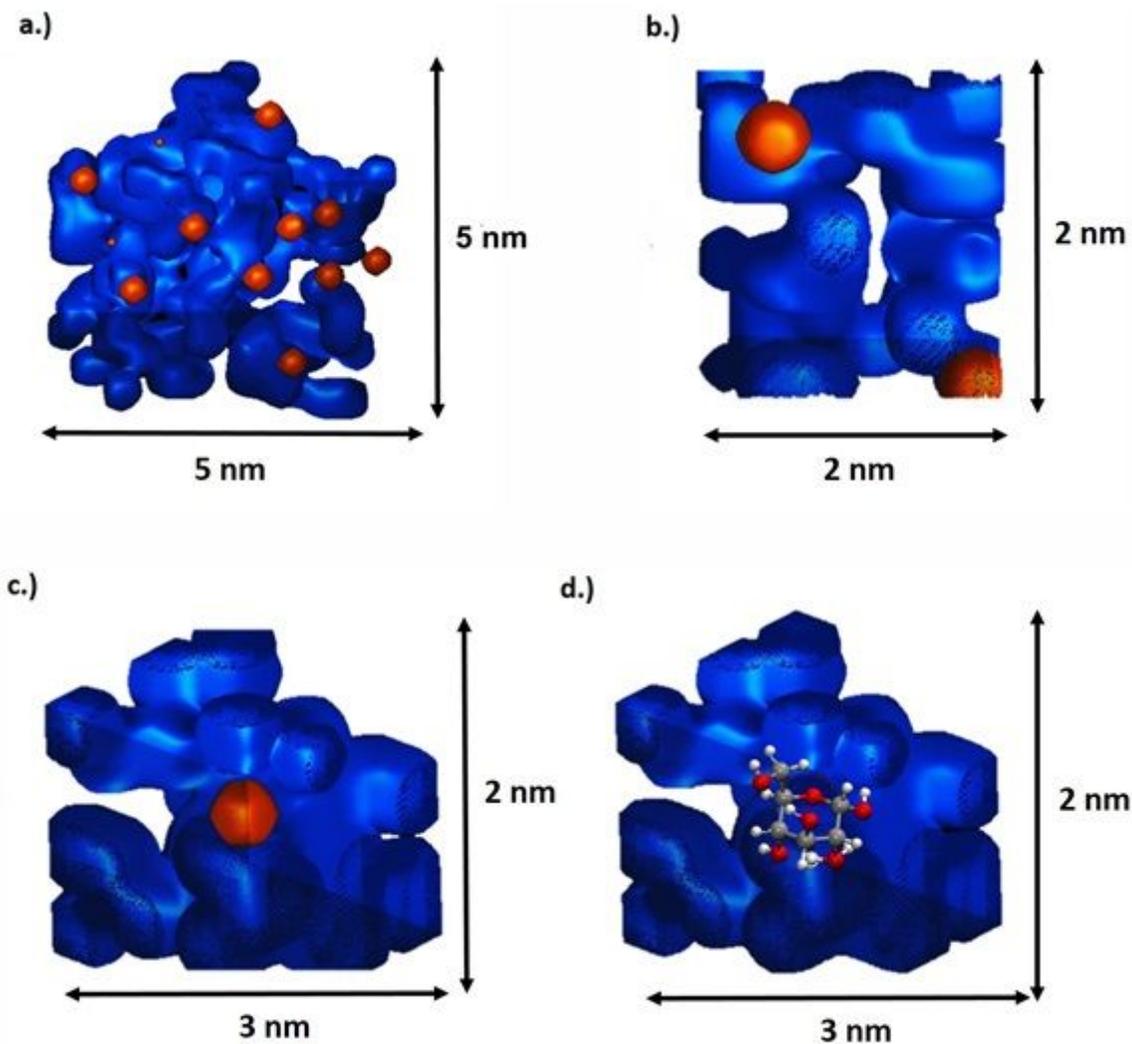


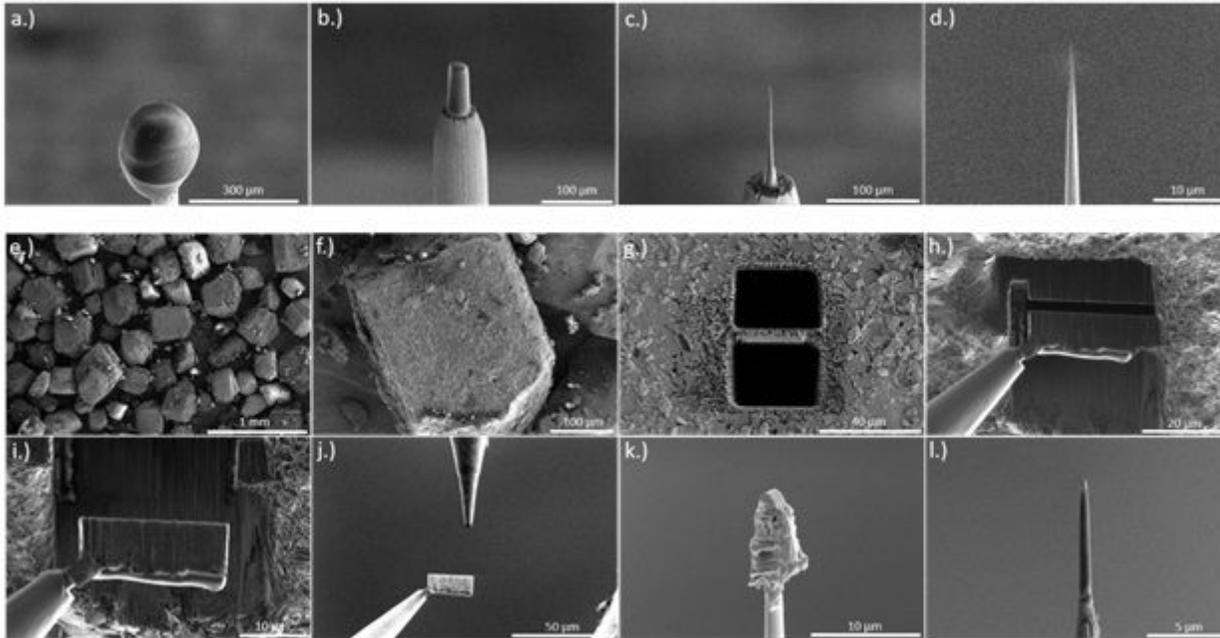
Figure 9

a.) Radius distribution to include the NN other glucose molecules around a glucose molecule. b.) Increase of the mean radius with number of nearest neighbours.



**Figure 10**

Isosurfaces of water in blue 67% and the complete glucose molecule at  $m/q = 181 \text{ u e}^{-1}$  in orange are shown to visualize the hydrate shell of the solvent glucose molecules.



**Figure 11**

FIB preparation – a.) Frozen droplet of a saturated glucose solution on top of a tungsten post. b-d.) Milling process with an annular pattern with decreasing inner diameter until a tip radius  $<100 \text{ nm}$  (d.) occurs. From e.) to l.) the lift-out procedure from a glucose bulk sample is shown, followed by an annular milling process to the final obtained tip (l.)

## Supplementary Files

This is a list of supplementary files associated with this preprint. Click to download.

- [SupplementaryPartvers.2.0final.pdf](#)