BerILL: The ultimate humidity chamber for neutron scattering

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Abstract. We present the design and the performances of a precision humidity chamber for neutron diffraction which is the result of a very fruitful collaboration between the Helmholtz-Zentrum Berlin and the Institut Laue-Langevin. The sample temperature can be set between 10 and 85°C. The relative humidity is controlled from 10 to 99.9% ± 0.01% by controlling the temperatures of the sample and a water bath at the millidegree Celsius scale. Sample exchanges are extremely easy and the operation only requires entering two parameters: sample temperature and relative humidity. The performances are confirmed by neutron diffraction measurements performed on DMPC and DOPC.

Keywords: Neutron scattering, soft matter, sample environment, humidity

1. Introduction

Neutrons are well-suited for studying biological and soft-matter samples. They are very sensitive to light elements, such as hydrogen, carbon, oxygen and nitrogen, which cover the majority of atoms found in most organic systems. Interacting with nuclei, they also have the advantage of differentiating isotopes of these atoms. The isotope effect between H and D is often exploited in diffraction as well as in incoherent scattering experiments, as the coherent scattering length changes from $-3.739 \text{ fm}$ for H to $+6.671 \text{ fm}$ for D, and the incoherent cross section from $80.26 \times 10^{-24} \text{ cm}^2$ to $2.05 \times 10^{-24} \text{ cm}^2$, respectively [20].

The characteristics and behaviours of samples investigated in biological or soft-matter neutron scattering experiments are strongly coupled to the temperature and relative humidity of the surroundings. Accurate observation and control of these parameters are crucial to the success of the experiments. The currently available humidity and temperature control chambers for neutron scattering allow experimentalists to set the relative humidity at the sample by using techniques such as saturated salt solutions [4,9,19], saturated/dry gas flow and partial vapour pressure through temperature of a water reservoir [8,14].
In recent years the performance of these experimental sample environments has improved. But due to the un-
reliability of humidity sensors and the presence of thermal gradients resulting in unwanted condensation, these
systems have been restrictive or finicky. In particular, it has been difficult to achieve relative humidity above 95%
while this near-saturation regime is of special interest to the field of biological science. Biological systems are gen-
erally warm and moist, and we can get insight of their in-situ behaviour only by creating experimental conditions
which closely match their natural conditions.

Many lipids spontaneously form bilayers, whose structure can be determined using neutron diffraction. The
structure of a bilayer is related to the bilayer’s phase. Of particular interest is $L_\alpha$ phase, which is biologically
relevant. The $L_\alpha$ phase can be characterised by rapid translational diffusion and trans-gauche isomerisation of
the hydrophobic tails preventing any long-range order [10]. The phase of a lipid sample depends on the temperature $T$
and relative humidity RH surrounding the sample:

$$ \text{RH}[%] = \frac{p_{\text{partial}}}{p_{\text{saturation}}(T)} \times 100 $$

where $p_{\text{partial}}$ is the partial water vapour pressure and $p_{\text{saturation}}$ is the saturation vapour pressure of water in ther-
modynamic equilibrium with the condensed liquid. The relative humidity can be induced by controlling $p_{\text{partial}}$
through precise control of temperature. This effect can be achieved by cooling pure water in a reservoir and heat-
ing the sample. By lowering the temperature of the water reservoir, the partial pressure $p_{\text{partial}}$ of the water vapour
in the cell decreases according to the equilibrium vapour pressure curve of the liquid, which is determined directly
from the bath temperature using the Antoine equation [1]:

$$ \log(p) = A - \frac{B}{C + T} $$

where $p$ is the vapour pressure in bar, $T$ is the temperature in °C, and $A$, $B$ and $C$ are component-specific constants
of the system. For light water (H$_2$O) and heavy water (D$_2$O), these coefficients are provided in Table 1(a) and (b)
from [3,16]. And from equations (1) and (2) above, we deduce the relative humidity:

$$ \text{RH}[%] = 100 \times \frac{10^B/(C+T_{\text{sample}})-B/(C+T_{\text{bath}})}{A} $$

This method allows to explore a continuous range of relative humidities with the possibility of fast changes
in humidity using temperature controllers. Although the change of humidity cannot be as quick as in a gas-flow
system, this technique is significantly more accurate. The temperature control technique also solves the problem
of inaccurate humidity sensors above 95% RH by relying on thermometers with high accuracy.

The drawback of this method is the requirement for an environment with extreme temperature homogeneity in
the water bath as well as in the sample surrounding area (±0.03°C to get ±0.1% RH at 95% RH and 30°C). Also,
any cold spot (with respect to the water bath) in the sample chamber creates a condensation point and in principle
a new water reservoir at an uncontrolled temperature. By effectively insulating the humidity chamber, the influence
of unwanted temperature gradients and cold spots can be suppressed.

<table>
<thead>
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<th>Temperature [°C]</th>
<th>A</th>
<th>B</th>
<th>C</th>
</tr>
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<td>5.40221</td>
<td>1838.675</td>
<td>241.413</td>
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<tr>
<td>31–60</td>
<td>5.20389</td>
<td>1733.926</td>
<td>233.665</td>
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<tr>
<td>61–90</td>
<td>5.0768</td>
<td>1659.793</td>
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<tr>
<td>71–100</td>
<td>5.08354</td>
<td>1663.125</td>
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<table>
<thead>
<tr>
<th>Temperature [°C]</th>
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<tr>
<td>106–300</td>
<td>5.15107</td>
<td>1700.073</td>
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Table 1
Antoine’s constants for H$_2$O and D$_2$O extracted from [3,16]
In the following sections, we present the design and performance of the new BerILL humidity chamber whose name comes from the collaboration established between the Helmholtz-Zentrum Berlin (HZB) and the Institut Laue-Langevin (ILL). BerILL is designed for neutron scattering (preferably diffraction) and exploits the temperature control technique to induce humidity on a sample with excellent accuracy and ease of control (the user only needs to request a relative humidity and sample temperature). Performances are then demonstrated with the well defined DMPC lipid bilayer [15].

2. Design

The chamber has been designed to be compatible with commonly used neutron diffraction instruments which have a large user base in biology (e.g. V1 at the HZB [11] and D16 at the ILL [6]), but with the possibility to adapt it to reflectometry, SANS, backscattering or time of flight instruments in future versions. The design has been realised with a particular focus on thermal insulation from external heat sources and ease of use. So we started from a simple double-wall geometry allowing to vacuum insulate the sample space. The dimensions were optimised in an iterative process to determine the geometry which resulted in the most stable, thermal gradient free design. Each step in this iterative design process was scrutinised using finite element simulations of thermal equilibrium and convection with the COMSOL software package [5]. Figure 1 shows how the heater placed below the goniometer allows to reduce sensibly the temperature gradient in the sample region. The distance between the water bath and the goniometer, i.e. the height of the post, was also optimised.

The humidity chamber shown in Fig. 2 is a double walled cylinder construction with a vacuum insulating the inner chamber. The outer and inner chambers are both split in two parts, upper and lower, to ease the access to the sample space and facilitate the operations.

![Fig. 1. Finite element simulations of the design at room temperature and 40% RH.](image-url)

(a) Opened COMSOL model of the chamber showing the goniometer mounted on a base in copper, itself mounted on a post in PEEK above the water bath.
(b) Temperature equilibration simulated at 40% RH without heating the base of the goniometer.
(c) Temperature equilibration simulated at 40% RH while heating the base of the goniometer. The temperature around the sample is perfectly constant.

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Fig. 2. CAD design of the humidity chamber showing internal parts. The temperature of the water bath is controlled with a thermostated bath connected to the chamber base. The sample, mounted on a goniometer insulated from the water bath, is thermalized with a second thermostated bath connected at the top of the chamber.

The lower part, which is fixed to the sample table of a neutron diffractometer, contains the water bath and several access ports to evacuate the chamber, read the temperature of the water, read the temperature and relative humidity around the sample, read the pressure in the vacuum space and control the temperature at the base of the goniometer. The water is poured in a copper bath decoupled thermally from the sample space by using polyether ether ketone (PEEK) which has excellent mechanical and chemical resistance properties. The temperature of the bath is controlled with a first thermostatic bath. The outer vessel through which the thermostatic bath is connected is made from stainless steel to reduce the heat conduction with the upper part. The in and out connections to the bath are made through 0.08 mm thick bellows accepting thermal expansions and the mechanical mount is ensured with 3 pillars made from PEEK.

The upper part of the humidity chamber is removable to provide a quick access to the sample: the inner and outer chambers are permanently attached to the top flange and the whole upper part can therefore be removed in one go. The upper part controls the sample temperature with a thermostatic bath between 10 to 90°C with highest homogeneity across the sample area. Because simulations revealed a potentially very slightly lower temperature at the bottom of the sample (a few 1/100°C), a resistive heater and a thermometer are introduced in the copper base plate onto which a standard Huber manual goniometer is fixed [13]. According to the simulations (Fig. 1), this should allow to control the temperature in the sample volume with an accuracy better than ±0.002°C, corresponding to a theoretical stability in the relative humidity of ±0.01%.

The lower and upper parts need to be thermally insulated as well as possible while simultaneously allowing a fast gas exchange for short humidity equilibration times. Therefore, from COMSOL simulations, the 80 mm bath surface and the cross-section for gas exchange have been increased as much as possible with respect to the thermal isolation performance needed to avoid unwanted gradients and resulting inhomogeneities in the humidity across the sample area. The sample was also placed at an optimum distance from the bath with a post also made from PEEK.

The effective sample volume is a 20 mm × 40 mm cylinder. The total horizontal and vertical angular accesses are respectively 300° and 35°. The sample alignment can be done in place optically with the goniometer and a laser. Closing the chamber is performed by lowering the upper part of the cell on the bottom part containing the
sample and the water bath. A gliding seal seals the inner chamber against the insulating vacuum and the closing
procedure is simplified by two guiding posts which prevent contact between the upper part of the chamber and the
sample and sensors. If necessary, the entire cell can be stored away from the instrument during stabilisation and
reassembled in the same mechanical position.

All parts inside the sample space including sensors and wiring are protected from corrosion by coatings from
PTFE (wires, sensors), gold plating (Cu parts) and anodisation (Al parts) where necessary. Temperature, pressure
and humidity sensors are mounted near the sample in the 60° dark angle of the humidity chamber. The wires
penetrate the vacuum space of the chamber in the lower part and reach the sample space through the insulating
post onto which the goniometer is placed. Leak tight feedthroughs made in the copper base allow wires to enter
the sample space in the dark angle of the cell.

3. Construction

3.1. Mechanical components

Figures 3 and 4 present respectively drawings and photos of the cell. The sectional view A-A shows the internal
parts and sectional views B-B and C-C present in detail the circuitry of the fluids delivered by the two thermostated
baths. The overall dimensions without support are $180 \times 480$ mm.

The inner chamber hosting the sample and driving the fluid from top to bottom and then to the top is machined
from a single rod of hard aluminium EN AW 5754 (Fig. 4(a)). As for the lower part of the cell, the in and out
fluid connections are performed through 0.08 mm thick bellows accepting thermal expansions and the mechanical
mount of the inner chamber is ensured with 3 pillars made from PEEK. To reduce the neutron background and
keep a good neutron transmission, the thickness of the wall is reduced to 0.5 mm at the beam height. The top and
bottom ends of this central part are tightly closed with respectively stainless steel and PEEK flanges on O-rings.
The use of stainless steel decouples thermally the inner chamber from the outer vessel. The PEEK flange which
will be introduced into the top of the water bath when closing the chamber is mechanically strong enough to ensure
a good tightness and prevent thermal conduction with the water bath.

On Fig. 4(a), the small central top flange is closed with a tap but was later used for installing a Jaeger connector
allowing to read temperature sensors glued with GE varnish to the top and bottom ends of the inner chamber where
the fluid circulates. The insert shows details of the fluid circuitry before assembly. A 30 $\mu$m foil of aluminium was
then wrapped around the central part to screen the radiation of the outer vessel and further reduce the gradient of
temperature.

The lower part of the humidity chamber (Fig. 4(b)-left) is a little trickier to assemble because of the installation
of the wiring in a tiny space. All wires, except those of the thermometer of the bath, are inserted in the central
post. Six wires are used to power the heater and read the temperature inside the copper base. The others are used
to measure the sample environmental conditions ($p, T$ and RH) and exit the base through a capillary filled with
GE varnish. As shown on Fig. 4(b), these sensors are attached to a glass fibre rod parallel to the Silicon wafer
on which the sample is deposited. The vacuum valve, the Jaeger connectors and the guiding posts are all located
in the dark angle of the cell. After assembly, the whole chamber was successfully helium leak tested down to
2.4 $\times 10^{-9}$ mbar · l/s. The right of Fig. 4(b) shows the upper part fully assembled and ready to close the chamber.

3.2. Humidity control

To reach the required temperature precision, we employ precision thermistor resistances Omega TH-44007-40-T
(5 k$\Omega$ @ 25°C) [18] recalibrated to reach an absolute precision of ±0.025°C on the range 0 to 90°C (i.e. $\approx$ ±0.1%
RH). As shown in Fig. 5(a), these sensors are much more sensitive than Pt100 thermometers, typically by factors
of $\approx$700 at 20°C, $\approx$430 at 30°C and $\approx$270 at 40°C. According to the supplier, they are also chemically stable and
not significantly affected by ageing or exposure to strong fields of hard nuclear radiation.
To observe the equilibration times of the relative humidity in the cell, we use an Honeywell HIH-4000 humidity sensor of ±3.5% accuracy and interchangeability of ±5% up to 59% and ±8% from 60 to 100% [12]. This sensor is a laser trimmed, thermoset polymer capacitive sensing element with on-chip integrated signal conditioning. According to the supplier, the sensing element’s multilayer construction provides excellent resistance to most application hazards such as wetting, dust, dirt, oils and common environmental chemicals. The operation environment of this sensor covers the temperature and humidity produced by the chamber but the supplier does not recommend its use at high temperatures and humidities as shown in Fig. 5(b). It is also light sensitive and an extended exposure to >90% RH causes a reversible shift of 3% RH.

As shown in Fig. 6, this sensor brings heat locally, to a point where it creates a temperature gradient $\Delta T \approx 0.27^\circ C$ in the sample space leading to a potential humidity shift of $\approx 1.5\%$. The humidity being controlled and determined by the sample and bath temperatures, we decided to power the sensor only when appropriate. The
Fig. 4. Photos of humidity cells built during this project. Inner chambers as the one shown in photo (a) are mounted inside the upper part of the cells shown at the right of photo (b). The inset in (b) shows the detailed assembly of the sensors placed in the dark angle.

(a) Photo of the assembly of the inner chamber. The insert shows details of the circuitry before assembly. (b) Photos of the lower and upper parts of the humidity chamber. The connectors, the vacuum port and the guiding posts are gathered in the dark angle of the cell.

Fig. 5. The relative humidity in the chamber is precisely controlled with temperature sensors and its equilibration is determined with a humidity sensor.

(a) Calibration curve of one of the precision thermistor resistances installed in the chamber compared to the curve of a Pt100. (b) Operating conditions of the humidity sensor recommended by the supplier.

Fig. 5. The relative humidity in the chamber is precisely controlled with temperature sensors and its equilibration is determined with a humidity sensor.
Fig. 6. The heat brought by the humidity sensor generates a gradient of temperature across the sample of ≈0.27°C at 30°C. This leads to a shift of the relative humidity by ≈0.9% at that temperature. We therefore decided to power the sensor continuously only above 50°C to prevent the saturation of the sensor and when the heating effect becomes less important. We can see at the right of the graph that the temperature gradient is less than 0.005°C when the humidity sensor is not powered.

Fig. 7. Evolution of the relative humidity at 30°C with (left) and without (right) employing the overshoot technique. The humidity sensor is shifted by several % as expected from the sensor performance specifications. The continuous blue lines correspond to values measured by the humidity sensor and the dotted red lines are deduced from equation (3) and temperature sensor readings. The curves on the left show that the equilibration time increases with the humidity from about 2 to 6 hours. On the right, the temperature setpoints are overshot by 10% RH at the beginning of each step and set to wanted values when the sensor reaches that value. As can be seen in the inset of Fig. 7, the equilibration times are strongly reduced and range between 20 minutes and 1 hour, i.e. about 6 times faster.
As expected, we also observed a slight difference between the sample temperature and temperature set into the thermostated bath controller (we use 5 m long pipes on the experimental area). To reduce it, we tried to exploit the internal PID of the thermostatic bath by connecting a sample thermometer to the controller of the thermostatic bath. This did not improve the situation because of the concurrent inertia of the two systems. We therefore decided to let the user correct the sample temperature setpoint of the thermostatic bath if needed.

Finally, to check the pressure in the cell, we measure the absolute pressure within ±30 mbar with a Freescale Semiconductor MPX4200A series sensor which integrates on-chip, bipolar op amp circuitry and thin film resistor networks and provides a high level analog output signal and temperature compensation.

4. Performance

In order to quantify the absolute precision of the humidity chamber, we have measured the $d$-spacing of DMPC (1,2-dimyristoyl-sn-glycero-3-phosphocholine) and DOPC (1,2-dioleoyl-sn-glycero-3-phosphocholine) lipid molecules with the neutron diffractometers D16 (ILL) and V1 (HZB) at different temperatures and relative humidities [6,11]. During the first tests, with the beam time available, a maximum $d$-spacing of only 52.3 Å could be achieved at 30°C with DMPC, which corresponds to a maximum relative humidity of ≈99% according to [15].

To overcome the slowness with which these lipids absorb water, we mounted the sample on an aluminium block in thermal contact with the cold side of Peltier elements to speed up changes of the environmental conditions (in place of the goniometer). The sample holder and details of the experiments are described in the PhD manuscript of one of the authors [2]. The measurements were conducted on V1 (HZB) with a DOPC bilayer sample which begins drastic swelling at a lower humidity than DMPC. As can be seen in Fig. 8, we controlled the relative humidity up to 99.9% but a curve splitting occurred above 59 Å indicating the presence of two domains in the sample. This may come from the operation of the two Peltier elements: the sample is cooler by a few 0.001°C on one side, causing a phase separation.

We then performed a new series of measurements on D16 (ILL). Figure 9 presents the evolution of the $d$-spacing of DMPC with the relative humidity at a constant temperature of 30°C. The red curve corresponds to the humidity deduced from the control of the temperature of the water bath and the blue curve to the value measured with the humidity sensor. The sensor always measures larger values but remains within the tolerance provided by the supplier. During these tests, the DMPC was in the gel ($L_{β}$) phase [21] and the distance increased slowly as expected.

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**Fig. 8.** Bragg reflections from a DOPC lipid bilayer at 20°C (right). Calculated $d$-spacings of each measurement are shown on the left up to 99.9% RH. Above a $d$-spacing of 59 Å splitting occurs, which may indicate two domains in the sample.
As shown in Fig. 10, we then requested a relative humidity of 97% by controlling the temperature of the water bath and measured \( d = 51.8 \, \text{Å} \). We opened the chamber and filled the water bath with a saturated salt of potassium sulfate to define the same environment of 97.00 ± 0.40% [17]. In the morning, the neutron source unpredictably stopped before the DMPC reached the equilibrium but we could fit the evolution of the \( d \)-spacing with an exponential relaxation function and deduced a value of 51.9 Å. The small difference of 0.1 Å corresponds to a gap of \( \approx 0.1\% \) RH, which is of the order of the uncertainty with which the relative humidity is obtained with the saturated salt.

The ultimate test consisted in measuring the \( d \)-spacing at the highest possible relative humidity. We tuned the chamber to 100% RH and stored the cell aside for one day until we remeasure the \( d \)-spacing on D16. We fitted a distance of 57.1 Å which corresponds to a relative humidity of 99.9% according to [15]. The \( d \)-spacings of DMPC
5. Conclusion

We have designed a precision humidity chamber for neutron diffraction allowing to maintain a $20 \text{ mm} \times 40 \text{ mm}^3$ sample at any temperature between 10 and 85°C in a relative humidity between 10 to 100% (this range is obviously reduced at the lowest temperatures). This relative humidity is created with a water reservoir (H$_2$O or D$_2$O) regulated at a temperature deduced from Antoine’s equation. The sample space is thermally decoupled from the water bath and the experimental environment. This allows the control of relative humidity up to 99.9% with an accuracy of $\pm 0.01\%$ RH.

Today, one humidity chamber is available at the HZB and three identical chambers are available at the ILL on D16 where $d$-spacings can be measured after short or long equilibration times. Sample alignment can be performed off-line with a laser mounted on an optical bench, and equilibration can be started with no need to access the neutron beam. Future modified designs of this chamber are foreseen for other neutron techniques, in particular small angle scattering and reflectometry.

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