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The suitability of Tedlar bags for breath sampling in medical diagnostic research

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Abstract

Tedlar bags are tested for their suitability for breath sampling for medical diagnostic purposes. Proton-transfer reaction-mass spectrometry was used to monitor the changes in composition of various mixtures contained in custom-made black-layered Tedlar bags. Characteristic ions at \( m/z \) 88 and 95 amu reflect considerable pollution from the bag material. The pollutant found on \( m/z \) 88 amu is most probably \( N,N \)-dimethylacetamide, a latent solvent used in the production of Tedlar film. Gas composition losses during filling were found to range from 5 to 47%, depending on the compound. Once stored, the half-lives of methanol, acetaldehyde, acetone, isoprene, benzene, toluene and styrene were estimated between 5 and 13 days. Losses from breath samples (52 h after filling) were found to be less than 10%. No observable decrease was found for ethylene over 3 days, using laser-based photoacoustic detection. For the use of Tedlar bags, a standardized protocol is advised, where the time point of analysis is fixed for all samples and should be kept as close as possible to the time of sampling.

Keywords: PTR-MS, breath analysis, Tedlar bags, VOCs

1. Introduction

plasma diagnostics, investigation of engines or automobile exhaust measurements or headspace
publications have shown the diagnostic potential of relatively new, advanced techniques.
These include gas chromatography–mass spectrometry (GC–MS) (Phillips and Greenberg
1991), selected ion flow tube-mass spectrometry (SIFT-MS) (Wang et al 2004, Davies
et al 2001) and proton transfer reaction-mass spectrometry (PTR-MS) (Lindinger et al 1998a,
Prazeller et al 1998, Critchley et al 2004). The latter two techniques are very promising in
breath analysis and fast diagnostics, since they combine on-line measurement capacity with
non-invasive, sensitive and multi-component detection of VOCs.

It is not always possible to bring the gas emission source of interest and the gas detection
system together. In such cases, the gas sample under analysis needs to be stored in a container
and transported to the laboratory. For this, several methods are used, including canisters,
cold trapping, adsorbing agents and Teflon and Tedlar bags. These containers should be
easy to operate, easy to handle and be able to store a sample for a prolonged period of time.
Costs, re-usability, durability, size and versatility are other important issues when choosing a
sample holder. Adsorbing agents like Tenax are compound specific, whereas canisters have
the disadvantage that they need to be evacuated before sampling, making them inapplicable to
breath measurement. Teflon bags are relatively expensive and fragile, making them difficult
to be re-used. Here, as an alternative to these widely used methods we have investigated the
use of Tedlar bags for the sampling of breath from patients and healthy volunteers.

Tedlar bags have been used previously by Nielsen and co-workers (Nielsen 2002), who
have measured nine volatile sulphur compounds from biogas-production plants and reported
no detectable losses over a period of 20 h. Only water was found to permeate through the
walls until the relative humidity inside reached ambient air values. Sulyok et al (Sulyok
2001) observed a 90% recovery rate of sulphuric compounds after 1 week of containment
for clear layered Tedlar bags, but found significantly lower recovery rates for black-layered
bags. They assumed the losses to be due to uncontrolled adsorption to the carbon-filled black
layer. Lau and co-workers (Lau 1989) reported half-lives of six sulphur gases ranging from
0.02 days to 100 days. For acetone, a significant decrease after the first 6 h was found, using
standard Tedlar bags (Deng 2004). The same result has been obtained for isoprene levels
(Hyspler 2000). N-hexene, p-xylene and 1-propanol were stable for at least 30 min (Myung
1997).

Groves et al (Groves and Zellers 1996) studied the significance of water content in the
sampled air for the stability of methanol, acetone, 2-butanone, m-xylene, trichloroethane and
perchlorethylene. They spiked the bags already containing the sampled air with water to
supersaturation to invoke condensation in the bag. Significant differences between wet and
dry samples were only found for methanol, acetone and 2-butanone. However, the relative
humidity needed to induce this difference was about three times higher than the humidity
inside the bag expected from breath (Groves and Zellers 1996). Only methanol was slightly
affected at breath humidity levels.

These studies mostly used transparent Tedlar bags and focused on isolated compounds
or a complex of several sulphur compounds. It is not clear how this can be translated to
breath samples and to black-layered Tedlar bags. To our knowledge, the storage capabilities
of these bags for most breath VOCs have not been tested. Here, we test the changes in gas
composition of breath and controlled mixtures during storage in black-layered Tedlar bags.
PTR-MS, of which the potential for breath measurements was shown before (Jordan et al 1995,
Karl et al 2001, Lechner et al 2005), was used for the analysis of more complex mixtures,
including breath, while photoacoustic spectroscopy was used to monitor the behaviour of
ethylene (Harren and Reuss 1997, Harren et al 1999). The contents of bags with several
different mixtures were analysed and compared at several instances between 4 and 250 h after filling.

2. Materials and methods

2.1. Instrumentation

The proton transfer reaction–mass spectrometer (PTR-MS) used for this work is analogous to that described in Lindinger et al (1998b) (figure 1). A detailed description of the home-built system can be found in Boamfa et al (2004) and Steeghs et al (2006). The working principles of PTR-MS have been given in detail elsewhere (Lindinger et al 1998b, Boamfa et al 2004, De Gouw et al 2003). Therefore, only a brief description is given here.

The instrument consists of four parts: an ion source where H₃O⁺ ions are produced, a drift tube section, a separately pumped transition chamber and an ion detection section containing a quadrupole mass spectrometer and a secondary electron multiplier. In the drift tube, the trace gases from the sample gas are ionized by proton-transfer reactions with H₃O⁺ ions:

\[ \text{H}_3\text{O}^+ + \text{R} \xrightarrow{k} \text{RH}^+ + \text{H}_2\text{O} \]

where \( k \) is the reaction rate constant, usually close to or equal to the collision rate constant. This reaction only takes place when the proton affinity (PA) of the trace compound R is higher than that of water (166.5 kcal mol⁻¹ = 7.16 eV/molecule). A major advantage of using H₃O⁺ as the reagent ion is that the PA of water is higher than the PA of the normal constituents of air (cf N₂, O₂, CO₂, CH₄, CO, NO, Ar) and that most of the typical organic compounds are ionized by the proton-transfer (PT) reaction, since their PA are in the range between 7 and 9 eV. The reaction rate can be measured or calculated and is known for many of the PT reactions of interest (http://Webbook.Nist.Gov/Chemistry/, Lindinger et al (1998b)). Since the excess energy of the reaction is low, it results in only one or two characteristic ions per neutral molecule. For some molecular groups dissociation can occur to form one or two fragments of significant intensity (e.g. alcohols can split off a water molecule, which results in a fragment ion at molecular mass minus 17). Due to this soft-ionization the matrix of signals is less complicated than with other mass spectrometry techniques. One disadvantage of the technique is that it does not provide positive identification of the compounds monitored (Steeghs et al 2004).
Ethylene, a well-known indicator for lipid peroxidation (Kneepkens et al 1994) was measured using laser-based photoacoustic detection, as described in Harren and co-workers (Harren et al 1999, Harren and Reuss 1997). Briefly, the infrared laser is tuned to a wavelength where the gas under investigation has a high absorption coefficient. By modulating the intensity of the laser light, a periodic heating of the sample gas is caused. This results in pressure variations inside the sample cell of the same frequency as the laser intensity modulation. By carefully choosing the modulation period, an acoustic wave is created inside the closed sample cell, which is detected with a microphone. The amplitude of the acoustic wave is directly proportional to the laser intensity and to the concentration of the absorbing compound. By monitoring the laser power and the intensity of the sound wave, the concentration of the compound under investigation can be determined.

For both systems, calibration of concentrations was performed using certified mixtures. For the PTR-MS system we used range of dilutions of a mixture containing 600 ppbv of methanol, 800 ppbv of acetaldehyde, 900 ppbv of acetone and 1000 ppbv of isoprene, benzene, toluene and styrene in nitrogen. For the photoacoustic system the same procedure was used with a mixture containing 1 ppmv of ethylene in air.

2.2. Tedlar bags

We used 1.0 l black-layered Tedlar bags (SKC Limited, UK), especially designed for single-breath sampling experiments. The inner bag is made of a clear thin film of Tedlar and is covered by an outer layer of black Tedlar. This outer layer is blackened by adding carbon to the film to avoid UV-induced breakdown of any of the compounds possibly contained in the bag. Every bag has two Teflon hose/valve septum fittings, which simplifies the automated analysis and cleaning of the bags before re-use.

2.3. Experiments

All bags were flushed with synthetic air (mixture of purified nitrogen and 20 ± 1% purified oxygen; <3 ppm H2O, <0.5 ppm CnHm; Air Liquide BV, Eindhoven, The Netherlands) with a flow of ~25 l h⁻¹ for more than 2 h before use. Tedlar bags were filled with various mixtures of air using mass flow controllers (Brooks Instruments, 850S) to compare the composition of the contents at several points in time within a period of 10 days. Since the drift tube of the PTR-MS is operated at a pressure of around 2.25 mbar, no flow controllers or pumps were needed to measure contents of the bags. The sampling lines to and from the bags as well as the in-stream PTR-MS inlet were made of Teflon (PTFE and PFA, Polyfluor Plastics, The Netherlands) to minimize losses due to sticking of compounds to the walls. The calibrated mixture through the flow controller is kept constant at 0.2 l h⁻¹ at all times, so equilibrium situation is guaranteed and no losses occur here either.

We investigated whether any artefacts (increase of concentrations of known or unknown compounds) could be observed and whether there were losses to or through the bag wall. The influence of humidity on the stability of the contents was tested by varying the humidity of the sampled mixtures. Some bags were heated (T ~ 60 °C) to see if the temperature had an effect on the concentrations found from the bag contents.

2.3.1. Pollution of contents. A total of eight bags were filled with pure nitrogen (4.5 purity level) or synthetic air. These gas mixtures contained no detectable concentrations of hydrocarbons (at the (sub-) ppbv level) before filling. The contents of these bags were monitored to check for increase in concentrations from polluting compounds, either by
permeation through the walls of the bags, or by production from the bag material. These and all other measurements were performed at laboratory temperature, which is controlled at 22 °C, unless stated otherwise. Bags were always stored at lab temperature.

2.3.2. Loss of contents. The stability of a controlled mixing ratio of hydrocarbons was checked by filling 12 bags with a certified mixture (600 ppbv of methanol, 800 ppbv of acetaldehyde, 900 ppbv of acetone, 1000 ppbv of isoprene, benzene, toluene and styrene, Scott Specialty Gases, USA), diluted to a mixing ratio of 1 in 5 with nitrogen.

Also, a total of 24 bags were filled with the breath of 9 healthy volunteers. The contents of 10 of these bags were monitored 6 times in a period of 250 h; the other 14 bags were analysed 5 times within 72 h. It should be noted that condensation of water contained in the exhaled breath occurs in the sampling device before the bag, so no water droplets are formed inside the bag.

To test the suitability of these Tedlar bags for ethylene, 3 bags were filled with room air, containing low amounts of ethylene (below 1 ppbv), another 3 bags were filled with breath and 3 bags were filled with varying higher amounts of ethylene (65–100 ppbv). The concentrations in the bags at 1 h after filling were compared to the concentrations at 72 h after filling.

Possible losses could occur due to permeation through the walls or due to adhesion to the walls. To test whether compounds permeate through the wall, three bags were filled with a calibrated mixture (containing methanol at 600 ppbv, acetaldehyde at 800 ppbv, acetone at 900 ppbv and isoprene, benzene, toluene and styrene at 1000 ppbv). These bags were placed in a larger glass cuvette. The glass cuvette was flushed with nitrogen (4.5 purity level) gas for 10 min via an inlet and outlet before sealing, to make sure that no detectable hydrocarbon levels surrounded the bags at the moment the experiment started. After 2, 3 and 7 days, the contents of the glass cuvette were measured by PTR-MS and compared to nitrogen values and lab air values. After 7 days the contents of the bags were also analysed.

2.3.3. Relative humidity. Relative humidity was increased to 80% for 6 of the 12 bags by mixing the calibrated mixture with humidified (100% RH) nitrogen gas. The latter was done by bubbling the nitrogen gas through a water reservoir, resulting in a RH of 80% for the calibrated mixture. These experiments were performed to see if the water vapour in the bag affects the lifetime of compounds inside. The relative humidity is monitored by PTR-MS via the amount of cluster formation at mass 37 amu, where the water clusters H_3O^+ · H_2O can be observed.

All bags were measured several times over a prolonged period of time. Due to the limited volume of the bag (1 litre), we could only sample 5 or 6 times. Each time the bag contents were analysed, 8 mass scans from 20 amu to 150 amu were made (1 amu s⁻¹ scan rate) and averaged. The bags were stored and measured at laboratory temperature, which is set to 22 °C.

3. Results and discussion

3.1. Pollution of contents

Several bags were filled with clean air or pure nitrogen. These bags displayed significant amounts of pollution. Figure 2 (left panel) shows pollution reflected by characteristic ions at masses 88 and 95 amu where significant increases were found. Every single bag used in this study showed an increase of these two masses. The compound found at mass 88 is most
probably \(N,N\)-dimethylacetamide (\(\text{C}_7\text{H}_9\text{NO}, \text{MW} = 87 \text{ amu}, \text{PA} 217.0 \text{ kcal mol}^{-1}\)) and was found to originate from the Tedlar film (Chase 2001, Du Pont 1964). The Tedlar bag is made out of polyvinyl fluoride film, which is produced from polyvinyl fluoride powder, which is dissolved in a ‘latent solvent’ at temperatures above 100 °C, so the particles are allowed to coalesce into a film before the solvent evaporates (Chase 2001). Several candidates for this latent solvent are listed, among which are \(N,N\)-dimethylacetamide and related compounds (Du Pont 1964). After the drying process the total amount of latent solvent left in the material is assured to be less than 0.5% (Du Pont 1964). The concentrations of \(N,N\)-dimethylacetamide ranged from 40 to 185 ppbv (this is a lower limit, calculated using \(k = 2 \times 10^{-9} \text{ cm}^2 \text{ s},\) assuming no fragmentation) at mass 88 after 4 h and increased in time. Clean air and nitrogen contained no measurable quantities of \(N,N\)-dimethylacetamide (\(<0.5 \text{ ppbv}\)). The manufacturer of the bags assured that besides \(N,N\)-dimethylacetamide, no other solvent or chemical was used to make the bags. Our results, however, show that a second pollutant is present at mass 95. Independent GC-MS analysis has identified the compound at mass 88 amu to be indeed \(N,N\)-dimethylacetamide and the compound at mass 95 amu to be phenol (Amann, private communication, Di Francesco, private communication). Heating of the bags increased the signals on both mass 88 and 95 amu significantly (\(\sim170\%\) and 90% respectively, results not shown).

Some other masses also show a slight increase in ion intensity. This much lower degree of pollution was only observed for those bags containing synthetic air or nitrogen (figure 2, right panel). Mass 59 was found to be increased to about 5 ppbv after 250 h. It should be noted that these values are still less than 10% of the values expected in human breath and the influence will therefore be within the system’s uncertainty of the determination of breath concentrations.

Figure 2. Pollution of synthetic air and nitrogen gas (4.5 purity). Left panel: increase in masses 88 and 95 due to pollution from the Tedlar material (background values are zero). The large error bars are caused by large differences in amounts of pollution coming from the respective bags. Right panel: increase in masses 33, 45 and 59 (corresponding to expected compounds methanol, acetaldehyde and acetone, respectively) concentrations (background values subtracted). The large errors are caused by the low concentrations of the compounds measured and the variation between different bags.
3.2. Loss of contents

Losses of compounds contained in a Tedlar bag were monitored by sampling controlled amounts of calibrated mixture and breath. Figure 3 shows the evolution of the concentrations of the various compounds from the calibrated mixture. The values at 0 h are the mixed flows used to fill the bags.

There is a considerable loss ranging from 5% up to 47% at 4 h after filling. We consider these losses to occur during filling, presumably due to sticking to the septum in the inlet valves, since over time we find only a gradual decrease with respect to those values measured after 4 h. From the seven compounds, styrene is the only compound found to decrease for more than 15% (30 ± 7%) between 4 h and 52 h. From the decline of their concentration between 4 h and 72 h after gas sampling, a half-life for the respective compounds can be calculated. These half-lives are 8.1 ± 0.6 days for methanol, 6.5 ± 0.3 days for acetaldehyde, 8.4 ± 0.6 days for acetone, 13 ± 0.8 days for isoprene, 8.2 ± 0.8 days for benzene, 8.4 ± 0.7 days for toluene and 5.9 ± 0.3 days for styrene.

Other authors (cf Sulyok (2001)) have found lower recovery rates in black-layered Tedlar bags, as compared with clear Tedlar bags. In this study we aimed at characterizing black-layered bags and no attempt was made to compare both black-layered and clear Tedlar bags.

Low ethylene levels in the bags displayed no measurable decrease over 3 days. These bags include the ones filled with breath, where ethylene levels are usually around 1 ppbv. Higher ethylene concentrations in the bags decreased with less than 5% after 3 days.

Water vapour content decreases drastically by 80% within about 24 h (figure 3) and is then only slowly reduced to about 12.5% of its initial value after 250 h. Heating the bag does not regain this loss of water. This establishes the findings of Groves and co-workers (Groves and Zellers 1996), who suggest water to permeate through the walls of the bag.

In bags filled with breath, the decrease of sampled gas concentrations in time was found to be somewhat lower (table 1). From breath (tentatively identified) methanol, acetaldehyde,
Table 1. Relative average ion intensity values found at several masses from the breath of ten volunteers at different time points. The data at 2 h indicate average (±sd) count rates, where the values at later time points are expressed in percentages of the values at 2 h. P-value indicates the significance of a change in ion intensity between 2 h and 240 h. The spread in the 2 h values indicates the inter-personal variation in the concentration of the compounds.

<table>
<thead>
<tr>
<th>Mass (compound)</th>
<th>2 h signal intensity</th>
<th>24 h (% ± %)</th>
<th>72 h (% ± %)</th>
<th>144 h (% ± %)</th>
<th>240 h (% ± %)</th>
<th>P value of changeb</th>
</tr>
</thead>
<tbody>
<tr>
<td>(amu)</td>
<td>(ncps ± sd)</td>
<td>2.6 x 10^5</td>
<td>0.9 x 10^5</td>
<td>0.7 x 10^5</td>
<td>0.4 x 10^5</td>
<td>0.004b</td>
</tr>
<tr>
<td>31 (methanol)</td>
<td>14 ± 10</td>
<td>2.6 x 10^5</td>
<td>0.9 x 10^5</td>
<td>0.7 x 10^5</td>
<td>0.4 x 10^5</td>
<td>0.004b</td>
</tr>
<tr>
<td>41</td>
<td>87 ± 50</td>
<td>3.7 x 10^5</td>
<td>1.7 x 10^5</td>
<td>1.5 x 10^5</td>
<td>1.0 x 10^5</td>
<td>0.001b</td>
</tr>
<tr>
<td>43 (acetaldehyde)</td>
<td>2.8 x 10^5</td>
<td>1.2 x 10^5</td>
<td>1.7 x 10^5</td>
<td>1.5 x 10^5</td>
<td>1.0 x 10^5</td>
<td>0.761</td>
</tr>
<tr>
<td>47 (ethanol)</td>
<td>1.6 x 10^5</td>
<td>1.2 x 10^5</td>
<td>0.9 x 10^5</td>
<td>1.0 x 10^5</td>
<td>1.0 x 10^5</td>
<td>0.072b</td>
</tr>
<tr>
<td>59 (acetone)</td>
<td>3.7 x 10^5</td>
<td>1.3 x 10^5</td>
<td>1.0 x 10^5</td>
<td>1.4 x 10^5</td>
<td>1.0 x 10^5</td>
<td>0.851</td>
</tr>
<tr>
<td>61</td>
<td>1.7 x 10^5</td>
<td>0.9 x 10^5</td>
<td>1.2 x 10^5</td>
<td>1.1 x 10^5</td>
<td>1.3 x 10^5</td>
<td>0.450</td>
</tr>
<tr>
<td>63</td>
<td>45 ± 23</td>
<td>4.2 x 10^5</td>
<td>3.8 x 10^5</td>
<td>3.6 x 10^5</td>
<td>2.6 x 10^5</td>
<td>0.0018</td>
</tr>
<tr>
<td>69 (isoprene)</td>
<td>4.6 x 10^5</td>
<td>1.2 x 10^5</td>
<td>1.2 x 10^5</td>
<td>1.3 x 10^5</td>
<td>0.9 x 10^5</td>
<td>0.380</td>
</tr>
<tr>
<td>71 (N,N-dimethylacetamide)</td>
<td>6.8 x 10^5</td>
<td>0.8 x 10^5</td>
<td>2.5 x 10^5</td>
<td>1.1 x 10^5</td>
<td>2.9 x 10^5</td>
<td>0.007b</td>
</tr>
<tr>
<td>93</td>
<td>22 ± 14</td>
<td>0.6 x 10^5</td>
<td>0.6 x 10^5</td>
<td>1.2 x 10^5</td>
<td>1.5 x 10^5</td>
<td>0.410</td>
</tr>
<tr>
<td>95</td>
<td>2.5 x 10^5</td>
<td>1.3 x 10^5</td>
<td>2.9 x 10^5</td>
<td>2.1 x 10^5</td>
<td>3.3 x 10^5</td>
<td>0.000b</td>
</tr>
</tbody>
</table>

a Identification is tentative.
b P value of student’s t-test for equality of means.
c P values smaller than 0.1 are considered significant.
The suitability of Tedlar bags for breath sampling in medical diagnostic research

Figure 4. Concentrations of six different compounds in the glass cuvette enclosing three Tedlar bags filled with calibrated mixture. As the concentrations of compounds in the bags decrease (see figure 3), a corresponding increase is found in the surrounding airspace in the cuvette. Temperature was the same for all time points (22 °C).

acetone and isoprene can be compared with the certified mixture measurements. Of these four compounds, only methanol is found to decrease significantly. Besides methanol, only ethanol is found to also decrease significantly. On the other hand masses 41, 88, 95 increase significantly. Deviations from the average values are increasing over time.

Heating of the bags resulted in an increase of several compounds, although the effect for most compounds was in the range of 2–10% of the value for the unheated bag. This indicates that only a small proportion of the volatile contents is adsorbed to the wall and that most of it probably diffused through. Masses 88 and 95, which are due to the bag material itself, displayed an increase of about 170% and 88% with respect to the values of the unheated bag.

A separate experiment was performed to test the diffusion through the bag wall. Figure 4 shows the concentrations of six compounds in the glass cuvette enclosing three Tedlar bags on four different moments. Methanol, acetaldehyde and acetone clearly increase over time, whereas benzene and toluene show no significant increase. The gain in concentration in the cuvette corresponds well with the decreases found from the bags for acetaldehyde and acetone (within measurement uncertainty). For methanol, only 20% of the losses from the bag content are found back in the cuvette. As can be seen from the trends in figure 4 the loss rate decreases when the difference in concentration between the inside and outside of the bags decreases, which can be expected from a diffusion phenomenon.

3.3. Relative humidity

The influence of relative humidity on the concentrations in a bag was tested by increasing the relative humidity to at least 80% (100% for bags containing air or nitrogen, 80% for bags containing calibrated mixture). There was no observable effect of water vapour on the stability of the bag contents. No significant differences could be found between ‘dry’ (RH = ‘0’%) and ‘wet’ air (RH > 80%) (results not shown).
4. Conclusion

We have tested the suitability of black-layered Tedlar bags for breath sampling. The contents of the bags were found to be polluted, by two compounds giving rise to characteristic ions at masses 88 and 95 amu, identified by three independent groups as $N,N$-dimethylacetamide and phenol, respectively (Amann, private communication; Di Francesco, private communication). It should be kept in mind that these products are present in significant quantities when analysing the volatile content of the bags. The ion intensities and the variation therein essentially make them ‘blind spots’ in PTR-MS analysis of the bag contents, since the amount of pollution varies greatly from one bag to another.

Compounds from the standard gas mixture are found to display considerable and reproducible losses between 5 and 50% during filling. The losses possibly occur due to sticking effects on the septum in the fitting. After this drastic decrease during filling, all compounds except styrene show only a slow decrease in concentration, ranging from $\sim$10% within 2 days for isoprene to $\sim$25% for styrene, which shows by far the fastest decrease of all compounds tested (see figure 3). The compounds in the test mixture represent most chemical groups of volatile organic compounds usually found in breath (Amann 2005). An important other chemical group, sulphur compounds, has been investigated in more detail elsewhere (Nielsen 2002). The pure compounds tested here all display a half-life of $\sim$6 days or longer.

Immediate losses during filling can also be expected for breath VOCs, although these have not been tested here. Most of the losses during storage from compounds in breath were limited to less than 10% within 52 h. These observed decreases from VOCs in breath samples (compared are methanol, acetaldehyde, acetone and isoprene) are lower than the decreases measured for the compounds in the standard gas mixtures. This might be due to the relatively high concentrations (200 ppbv or higher) from the pure compounds in comparison to most values commonly found in breath. Diffusion, which we showed to be a loss mechanism, depends on the concentration gradient, decreasing the losses with decreasing differences in concentration between the inside of the bag and the environment around it.

Relative humidity showed not to have a significant effect on the lifetime of compounds in the bags. Groves and co-workers (Groves and Zellers 1996) did find a significant influence for methanol and acetone, but only at supersaturated humidity levels.

Even though two compounds are demonstrated to pollute the contents of the bags, these will not interfere with the analysis of other compounds contained in them. After immediate losses during filling, only slow and reproducible (comparable for all bags measured in two series of measurements) decreases in concentrations are observed. The variations in these losses are generally smaller than the inter-personal differences for most compounds present in breath (for these variations, see, for instance, Turner et al (2006a, 2006b, 2006c, 2006d)). To increase reproducibility, we advise a fixed point in time after sampling is chosen for the analysis of the contents. We conclude that, as long as the bags are characterized for the compounds measured in such a study and a fixed time point after collection is chosen, black-layered Tedlar bags are suitable to be used for sampling breath in clinical studies, where it is not possible to analyse the breath sample immediately.

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Endnotes

(1) Author: Please include references ‘Amann, private communication’ and ‘Di Francesco, private communication’ in the references list.
(2) Author: Please provide complete names of authors in references Deng (2004), Hyspler (2000), Myung (1997) and Sulyok (2001).
(3) Author: Please provide page number in reference Harren et al (1999).
(5) Author: Please check whether the sense of the sentence ‘The data at 2 h indicate average . . . .’ retains your intended sense in caption of Table 1.
(6) Author: Please check the value ‘1.8 × 10^2 ± 0.8 × 10^2 in the table as set.

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