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Vibrational spectroscopic analysis of body fluids: avoiding molecular contamination
using centrifugal filtration

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Abstract

The use of centrifugal filtration for protein purification and concentration represents an important improvement for the application of vibrational spectroscopy to analysis of bodily fluids based on protein fractions with specific molecular weight. Vibrational spectroscopic techniques are highly specific and can potentially detect small variations in the protein content indicating the presence of different diseases, therefore usable as markers for early diagnostic. However, due to the high sensitivity of the techniques, it is essential to verify that no molecular contamination occurs during the preparation of the samples. Concentration of the blood serum using commercially available centrifugal filters has been shown to result in an improvement in the spectral intensity and quality. However, the presence of contaminant features, identified as deriving from glycerine, can be found in the data collected after use of the centrifugal filters. It is demonstrated in the present work that careful washing of the filters is required before the analysis of body fluids based on concentrated samples. The demonstrated protocols for spectroscopic measurement of human serum are applicable to a range of bodily fluids and should accelerate potential clinical applications.

Keywords: Vibrational spectroscopy, ATR spectroscopy, bodily fluids, human serum, centrifugal filtration, glycerol, filter washing.

Graphical abstract: Problem of molecular contamination associated with centrifugal filters for analysis of human body fluids using vibrational spectroscopy

1. Introduction

The potential of vibrational spectroscopy, both Fourier transform Infrared (FTIR) and Raman, has been widely investigated for diagnostic purposes in cells and tissue analysis and the feasibility to use them for disease diagnostics has been extended to a wide range of bodily fluids¹ ranging from serum²⁻⁵ to tears^{6, 7} urine and saliva⁸⁻¹⁰. The past decade has seen significant development of both modalities for molecularly specific analysis on the micron and even nanoscale¹¹⁻¹⁷. Analysis of bodily fluids for diagnosis and health monitoring presents a less invasive approach than performing biopsies of organs for example during cancer screening¹⁸. In this context, blood serum presents a promising candidate for the application of vibrational spectroscopy, as the analysis of the proteins present in the blood stream can potentially deliver crucial information on patient health and indicate the presence of numerous pathologies¹⁹. Blood serum contains >20,000 different proteins, ranging from 50 g.L⁻¹ (serum albumin)^{20, 21}, to less than 1 ng.L⁻¹ (troponin)²², with an overall protein concentration of ~1 mM. Although other components such as lipids (lipoproteins) can also be found, protein sensing remains the main challenge for diagnostic purposes. The low molecular weight fraction of the serum, referred to by the term “peptidome”, is of particular interest for the potentially rich cancer-specific diagnostic information it contains^{23, 24}. Bound to high-abundance proteins such as albumin, it represents a fingerprint of the molecular events taking place within different organs or tissues related to the presence of cancers but also modification to their close micro-environment^{25, 26}. Therefore, the analysis of protein imbalances in the serum can be directly related to, and indicative of, disease states²⁷⁻³⁰.

In both Infrared and Raman spectroscopic studies, to date the analysis has predominantly been performed on air-dried drops of serum deposited on spectroscopically neutral substrates such as CaF_2 ^{31, 32}. The main reasons are the relatively low concentration of analytes in the serum, leading to poor signal to noise ratios (Raman spectroscopy) or the strong contribution of water in the spectra collected (Infrared spectroscopy). Such dried deposits are, however, extremely chemically and physically inhomogeneous and can lead to spectral saturation (in transmission or transmittance FTIR) or high variability in spectral recordings in point mapping modes of Raman or FTIR, requiring rigorous and lengthy sampling and averaging procedures. Moreover, using FTIR in transmission mode, in order to collect a spectrum, the serum is diluted 2/3 fold leading to longer sample preparation procedures and drying times with the possible introduction of external contaminants. ATR spectroscopy (Attenuated Total Reflection) is a particularly interesting alternative for the application of FTIR to body fluids analysis³³, as in this mode the sampling should average over the whole film in a single acquisition. It generally requires less than 10 μL to be deposited on the ATR crystal, which is then air dried to reduce the water contribution in the data collected. Due to the extremely small volume used, only a few minutes are necessary to obtain a thin dry biofilm that can be recorded^{24, 34}.

More recently, the use of commercially available centrifugal filters has emerged as an alternative strategy which allows concentration of the blood serum proteins resulting in an improvement in the spectral intensity and quality³⁵. Moreover, it presents the possibility to rapidly separate protein fractions with different molecular weight according to the cut-off points selected, which can considerably improve the

sensitivity of disease detection by removing the most abundant high molecular weight proteins, thus revealing spectral features usually hidden from the less abundant low molecular weight molecules³⁶⁻³⁸. It is likely that such centrifugal devices will become increasingly popular for applications of vibrational spectroscopy to analysis of bodily fluids and already preliminary work can be found in the literature^{24, 34}. However, many concerns have to be addressed before being able to evaluate their benefits for diagnostic purposes. The high sensitivity of the vibrational techniques is often used as an argument for their potential to visualize minor modifications in the molecular profiles from different samples. Therefore, the techniques have the capacity to detect and identify biomarkers specific of different pathologies. However, although vibrational spectroscopies are label free techniques, some requirements regarding the sample preparation have to be considered to collect relevant data. The high sensitivities also mean that any trace impurities introduced in the samples during the preparation process, for example centrifugation or filtration, could also contribute significantly to the spectral profiles collected. In the present work, it is demonstrated that, prior to blood serum analysis, it is essential that the centrifugal filters are thoroughly washed in order to avoid contamination of the samples by molecules present in the as-delivered filter membrane. Although the application of vibrational spectroscopy to human serum has been taken as an example, the same considerations apply to analysis of any of the human bodily fluids by such methods.

2. Materials and Methods

2.1 Sample preparation

Sterile filtered human serum from normal mixed pool (off the clot) was purchased from TCS Biosciences (Ireland). Commercially available centrifugal filtering devices, Amicon Ultra-0.5ml (Millipore - Merck, Germany), with cut-off points at 100K and 10K, were employed in this study as examples.

At first, 0.5 mL of the serum was placed in the 100K centrifugal filtration device, with no washing, and centrifuged at 14,000 x g for 30 mins. The filtrate obtained from the 100K device was then centrifuged using the 10K, under identical conditions. The filter devices were then placed upside down in a new Eppendorf and spun down at 1000g for 2 mins in order to collect the remainder of the serum (concentrate) retained in the filter devices. The concentrating factor is of the order of 10, with a resultant concentrate volume of ~50 µL. As a result, 2 fractions were obtained; the first representing proteins with a molecular weight higher than 100K; the second corresponding to the fraction between 100K and 10K.

As indicated by the manufacturer, the ultrafiltration membranes in Amicon® Ultra-0.5 devices “contain trace amounts of glycerine. If this material interferes with analysis, pre-rinse the device with buffer or Milli-Q® water. If interference continues, rinse with 0.1 M NaOH followed by a second spin of buffer or Milli-Q® water.” (<http://www.millipore.com/catalogue/module/c82301#1>)

However, no recommendation of washing procedures are provided. This study demonstrates that optimised washing of the centrifugal devices prior to serum analysis was achievable by spinning the Amicon Ultra-0.5ml once with a solution of NaOH (0.1M) followed by two rinses with Milli-Q water (Millipore Elix S). For both washing and rinsing, 0.5 mL of the respective liquid was added to the filters and the centrifugation was applied for 30 mins at 14 000g followed by a spinning with the

devices upside down at 1000g for 2 mins in order to remove any residual solution contained in the filter.

The pH of the NaOH 0.1M solution, Milli-Q water and 3 different buffer solutions (Fischer scientific) at pH 4, pH 7 and pH 10 were recorded using both a pH meter (Sension Quality Electrodes, Hach Company, USA) and pH-Fix test strips (Macherey Nagel, Germany), the latter due to the small volumes obtained from the centrifugal filters. Different ranges of pH indicator strips were used (pH-Fix 0-14, pH-Fix 7.5-9.5, pH-Fix 6.0-7.7) for better accuracy in the pH measurements. The buffer solutions were used to correlate the exact pH measured with the pH probes and different tints of color obtained on the pH indicators.

Additionally, glycerine (glycerol) (BDH, Ireland) was analysed as a reference for the identification of molecular contamination during the centrifugation.

2.2 FTIR Instrumentation : Data collection using the UATR

ATR spectra were recorded with the Perkin Elmer Spotlight 400N Universal Attenuated Total Reflectance (UATR) accessory of the spectrometer, which employs a 9-bounce diamond top-plate for this analysis. Sample penetration is both wavenumber and sample dependent, but is typically of order 1 μ m. In ATR mode, spectral data were the result of 4 scans, with a spectral resolution of 8 cm⁻¹. 5 μ l of the different liquid samples were deposited on the crystal and left to dry for 10 mins prior to recording. A background spectrum was also recorded and automatically subtracted by the software.

3. Results

The water contribution is a limiting factor for application of infrared spectroscopy to bodily fluid analysis due to the presence of a strong absorbance in the finger print region (figure 1C) ³⁵. Therefore, the protein features from the human serum are obscured and hardly identifiable, as can be seen in the ATR spectrum of a drop of serum, shown in Figure 1B.

Figure 1: Infrared spectra collected using the UATR accessory. A: Whole human serum after 10 mins drying; B: Whole human serum before drying and C: Milli-Q water. Spectra are offset for clarity

As the drop dries, however, the water features become less apparent, and the protein features more prominent, such that after 10 minutes drying, strong protein bands are identifiable in the fingerprint region, as shown in Figure 1A.

However, serum is a complex mixture and the highly abundant proteins dominate the spectral profile collected. Therefore, the use of centrifugal devices can greatly improve the sensitivity of the analysis by separating the different molecules according to their molecular weight ^{24, 39, 40}.

Figure 2 shows the spectrum of the dried complete serum sample (A), the >100K serum fraction (B), and the <100K serum fraction (D). Table 1 provides a list of band

assignments⁴¹⁻⁴⁷. Although the >100K serum fraction is very similar to that of the pure serum, consistent with the >100K fraction containing the most abundant proteins, the spectral profile of the 100K-10K fraction is significantly different, supporting the fractionation approach to provide more diagnostic information. However, the spectrum of the lower molecular weight fraction has strong contributions in the spectral region $\sim 1100\text{cm}^{-1} - 1000\text{cm}^{-1}$, which are not readily attributable to expected serum components. Notably, when diluted by a factor of 10, the spectrum of the >100K fraction also shows the same features (Figure 2C). The spectra of the filtrates were taken from samples which were processed using unwashed centrifugal filter devices.

Figure 2: Infrared spectra collected using the UATR accessory 10 mins air drying following. A: Whole serum; B: Concentrate from human serum using an unwashed 100K filter; C: Concentrate from 1:10 diluted human serum using an unwashed 100K filter and D: Filtrate from human serum using an unwashed 100K filter.

Human serum	
Wavenumbers (cm ⁻¹)	Assignments
3280	H–O–H stretching
2957	Asymmetric CH ₃ stretching
2920	Asymmetric CH ₂ stretching
2872	Symmetric CH ₃ stretching
1635	Amide I of proteins ν C=O (70–85%) / ν C–N / δ N–H
1537	Amide II of proteins δ N–H (40–60%) / ν C–N (18–40%)
1453	CH ₂ scissoring
1396	C=O stretch of COO-
1230-1330	Amide III (N-H bend in plane and C-N stretch)
1311	CH ₂ twist
1242	Asymmetric PO ²⁻ stretch
1170	Ester C–O asymmetric stretch
1110	ν (CN) / δ (CH)
1079	C–O stretch
1047	C–O stretch
926	C–C–N backbone, C–C stretch
Glycerine	
Wavenumbers (cm ⁻¹)	Assignments
3282	O-H stretching
2930	C-H stretching
2881	C-H stretching
1457	C-O-H bending
1409	C-O-H bending
1326	C-H deformation
1237	CH ₂ twisting
1108	C-O stretching
1030	C-O stretching
993	C-O stretching
922	O-H bending

Table 1: Band assignments for prominent FTIR peaks in Human serum and glycerine³⁴⁻⁴⁰.

A comparison of the infrared spectra collected from the serum filtered using unwashed Amicon centrifugal filters with the spectra obtained for Milli-Q water processed with similar unwashed devices clearly demonstrates that the features observed are not related to the protein content of the serum but are due to a contamination of the samples during the filtration process (Figure 3A and 3B).

*Figure 3: Infrared spectra collected using the UATR accessory after 10 mins air drying. **A:** human serum corresponding to the <100 KDa fraction (obtained after centrifugal filtration using the (unwashed) 100K cut-off point filter); **B:** Milli-Q water after centrifugal filtration using the (unwashed) 100K cut-off point filter and **C:** glycerol. Their respective high wavenumbers regions are plotted in D, E and F. Spectra are offset for clarity*

Notably, the peaks observed in the spectrum of the Milli-Q water correspond to those seen in spectra collected from a solution of pure glycerine (peak assignments also listed in Table 1), with weak contributions of residual water at $\sim 1675\text{cm}^{-1}$ (Figure 3C). Successive spinning of Milli-Q water through the same filter demonstrates that the amount of glycerine in the filtrates collected decreases according to the number of

centrifugation cycles, such that the contamination becomes negligible after the 4th cycle, indicating the possibility to wash off the traces of glycerine with water (figure 4).

*Figure 4: Infrared spectra collected using the UATR accessory collected from the filtrate from Milli-Q water after 10 mins air drying. The unwashed filter has been topped up with Milli-Q water between each centrifugation and the filtrate collected in separate Eppendorfs. **A:** first centrifugation; **B:** Second centrifugation; **C:** Third centrifugation and **D:** Fourth centrifugation. Spectra are offset for clarity*

Although the manufacturer's instructions specify that traces of glycerine can be found on the ultrafiltration membranes, the recommendations for their washing are rather imprecise: "If these materials interfere with analysis, pre-rinse the device with buffer or Milli-Q water. If interference continues, rinse with 0.1 N NaOH followed by a second spin of buffer or Milli-Q water" (<http://www.millipore.com/catalogue/module/c82301#1>) and no recommendations of spinning times, speed, volumes and repetition are provided. After testing different protocols, it has been found that the most efficient washing was achieved by spinning the devices with NaOH (once) and Milli-Q water (twice) for 30 mins at 14 000g. The process was followed by a spinning with the centrifugal devices upside down at 1000g for 2 mins in order to remove any residual solution contained in the filter.

Using a 0.1M NaOH solution, decontamination of the filters can be achieved more efficiently. While Milli-Q water alone requires 4 cycles, when washing the filters first with NaOH followed by rinsing with Milli-Q water, the traces of glycerine are completely removed from the filtrate after only 2 water rinse spinning cycles. However, the main concern when using NaOH is whether another source of molecular contamination is introduced in the data. Therefore, it is also crucial to verify that no traces of salt can be found in the data collected. The pH of the filtrates was monitored at the progressive points in the protocol. The pH of the NaOH solution and filtrate was found to be 13.5. After a first subsequent rinse with Milli-Q water, the pH of the filtrate was found to be 9. After a second rinse, the pH was found to be 7. Subsequently, a blank sample containing only Milli-Q water has been processed through the washed filters instead of the human serum. The data presented in figure 5C clearly highlight that, as seen in figure 4D, no traces of glycerine (Figure 3) or NaOH (Figure 5A) can be seen.

Figure 5: Infrared spectra collected using the UATR accessory collected after 10 mins air drying. A: washing solution of NaOH (0.1M); B: Milli-Q water processed through a centrifugal filter washed using NaOH. Spectra are offset for clarity

Using appropriate washing and rinsing procedures, the analysis of high and low molecular weight fractions can be achieved with a higher specificity and without

interferences from the glycerine and any other contaminant, as shown in figure 6, which compares the ATR spectra of the high fraction from a washed 100k (A) centrifugal filter with that of the true, uncontaminated low molecular weight fraction from a washed 10k filter (B).

*Figure 6 : Infrared spectra collected using the UATR accessory after 10 mins air drying. **A:** Human serum concentrate after centrifugal filtration using the washed 100k device concentrate; **B:** Human serum concentrate after centrifugal filtration using the washed 10k device concentrate. Their respective high wavenumber regions have been displayed in C and D. Spectra offset for clarity*

4. Discussion

A growing concern with vibrational spectroscopy is the absence of suitable controls when performing analysis. In most studies, the so-called normal samples are used as controls. However, as for colorimetric, dye based techniques, used for biomedical

application such as flow cytometry and *in vitro* cytotoxic assays, controls in the absence of active agents with only the monitoring reagents have to be also measured to make sure no false positives are recorded.

This is also the case for the spectroscopic analysis of filtered bodily fluids. As mentioned in the manufacturer's guidelines, traces of glycerine are present on the filter membrane, requiring washing prior to use, depending on the sensitivity of the analysis performed. Therefore, in the present case using centrifugal devices for analysis of bodily fluids using vibrational spectroscopy, additional controls containing only deionized water have to be prepared under the same conditions in order to verify that the washing is complete, and that no molecular contamination of the samples occurred. Although when applied to the concentration of biological samples containing antigens, antibodies, enzymes or DNA, rigorous elimination of the trace amounts of glycerine may not be necessary, vibrational spectroscopies are highly specific techniques, delivering a molecular finger print of the samples.. When applied to the analysis of human serum, the main interest remains the possibility to remove the highly abundant proteins with high molecular weights to reveal the smaller proteins or peptides potentially usable as biomarkers. As a result, the low molecular weight proteins are present in relatively small concentrations in the different fractions collected which, according to the data presented in figure 3, are even lower than the amount of trace glycerine collected from the filtering devices. This explains why glycerine features can appear prominently in spectra of centrifugally filtered fluids collected without performing an appropriate washing of the filters. Although the use of centrifugal devices is a promising approach for the field of vibrational spectroscopy applied to body fluids based diagnosis, the relevancy of preliminary studies that can be found in the literature is questionable due to the presence of molecular

contamination of the samples by glycerine following ultra-filtration, obviously greatly reducing the accuracy of the analysis^{24, 39, 40}. The true spectrum of the filtrate fraction is only achievable after appropriate decontamination. Although no attempt has been made to quantify the spectral features related to the constituent components in the serum, the study rather demonstrates that, using such centrifugal fractionation, potentially ultimately for medical diagnostics, it is essential to ensure that the washing protocols are optimised to avoid contamination of the sample. Glycerine itself has similar chemical moieties to peptides and proteins and thus the spectral profile overlaps that of the biomolecules of interest and thus rigorous elimination of the contamination is necessary.

5. Conclusion

The use of commercially available, Amicon ultra-0.5 centrifugal filter devices (Millipore - Merck, Germany) for application of vibrational spectroscopy to disease diagnostics purposes is perfectly suitable. However, to ensure relevancy of the results, it is crucial that the filtration devices are carefully washed beforehand to avoid any molecular contamination. Delivering a specific molecular fingerprint of the samples analysed, it is crucial to perform the measurements with all the suitable controls to validate the observations made and increase the relevancy of the data set collected. It has been shown that, due to the specificity of the techniques, the washing requirements are more rigorous than those prescribed by the manufacturers. However, as long as appropriate preparation protocols are established, such devices will contribute to expediting the realization of real clinical applications for spectroscopic analysis of bodily fluids.

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