Polarized Raman spectroscopic investigations on hemoglobin ordering in red blood cells

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Abstract. We have investigated the dependence of the Raman spectrum of an optically trapped red blood cell (RBC) on the orientation of the cell, relative to the polarization direction of the Raman excitation beam. The Raman scattered light polarized parallel to the polarization direction of the excitation beam was observed to depend upon the orientation of the cell. In particular, the heme bands at ~754 cm\(^{-1}\) and in the 1500 to 1700 cm\(^{-1}\) region were observed to become maximum when the cells’ equatorial plane was parallel to the excitation beam polarization direction and minimum when the cells’ plane was normal to the polarization direction. In contrast, no significant orientational dependence was seen in the Raman scattered light polarized orthogonal to the polarization direction of the excitation beam. Theoretical simulations carried out to investigate these observations suggest that inside the RBCs, the hemoglobin molecules must be present in an ordered arrangement, such that heme-porphyrin planes become preferentially orientated parallel to the RBCs’ equatorial plane.

Keywords: optical tweezers; polarized Raman spectroscopy; red blood cells; hemoglobin.

1 Introduction

The primary function of the red blood cells (RBCs) is the transport of oxygen from the lungs to tissues, which requires diffusion of oxygen into RBCs and their binding with intracellular hemoglobin (Hb). The intracellular Hb concentration determines the maximum amount of oxygen that can diffuse into the cell. While a high concentration of Hb as found in RBCs (~34%)\(^{1}\) ensures a large oxygen storage capacity, the resulting interactions between the neighboring Hb molecules strongly decrease Hb mobility. When compared with Hb in solution, the rotational and the translation diffusion coefficients for the intracellular Hb are lower by a factor of ~2 and ~5, respectively.\(^{2,4}\) This reduction in Hb mobility inside the cells can lead to a reduction in the intracellular “Hb-facilitated diffusion of oxygen”\(^{5,6}\) and consequently the oxygen transport through the cell. It has, therefore, been hypothesized that the intracellular Hb may be ordered to facilitate the diffusion of oxygen. Indeed, studies on low-angle scattering of x-rays from RBCs\(^{3}\) showed some evidence of ordering of Hb. Similarly, Perutz\(^{7}\) has shown that the high-intracellular concentration (~34%) of Hb molecules requires some ordering of these molecules inside the RBCs. However, there are no confirmatory studies showing the ordering of Hb within the RBCs.

Polarized Raman spectroscopy is a powerful technique to investigate the orientation of molecules in organized environments, such as crystal lattices, liquid crystals, and polymer samples,\(^{8–13}\) and thus can be useful for investigating the intracellular Hb ordering. The molecular orientation effects originating from the tensorial nature of the polarizability are reflected in the dependence of Raman bands’ intensities on the orientation of the electric field vectors of the incident and scattered light with respect to the molecular axes. Previously, some attempts\(^{14,15}\) have been made to understand the intracellular Hb ordering using this technique. In one of the studies,\(^{14}\) Raman spectra were recorded for two populations of RBCs; one having their equatorial planes normal to the beam propagation direction (population 1) and the other having their equatorial planes parallel to the beam propagation direction (population 2). These populations were observed to form spontaneously when a RBC suspension was poured onto a poly-L-lysine-coated substrate. In population 1, the RBCs had their equatorial planes parallel to the excitation beam polarization direction. However, in population 2, the equatorial planes of the RBCs could take any orientation relative to the beam polarization direction, while remaining parallel to the beam propagation direction. Hence, the mean spectrum taken over population 2 presented an averaged response taken over all possible orientations of the RBCs in the above-mentioned configuration. Although some orientation dependence of the RBC’s spectra was reported, the lack of precise control over the individual RBCs’ orientation limited the scope of this study. Raman optical tweezers\(^{16}\) are an attractive tool for addressing this issue. Because Raman measurements are made on a cell optically trapped in buffer media away from the substrate, the artifacts arising due to the immobilization of the cells on the substrate as well as due to the background originating from the substrate are avoided, leading to an improvement in the signal-to-noise ratio (SNR) of the Raman spectra. Dual trap tweezers have been used to simultaneously investigate two optically trapped cells to probe the dynamics and heterogeneity of interacting microbial cells\(^{17}\) and also to translate the trapped cell relative to the Raman excitation beam and thus acquire spatially resolved Raman spectrum\(^{18}\) etc. Rao et al.\(^{15}\) made use of a dual optical trap to orient the RBC with their equatorial plane...
parallel to the excitation beam polarization and to investigate the change in depolarization ratio of various Raman bands upon stretching the cells. The observed changes in the depolarization ratio were interpreted as an indication of an increase in Hb ordering inside the RBCs upon stretching.

A more thorough investigation of the anisotropic response of the Raman spectrum from the trapped RBC, which would help in gaining insight into the Hb ordering inside the cell, requires that the spectrum be acquired for different orientations of the equatorial plane of the RBC with respect to the polarization direction of the Raman excitation beam. This can be done using holographic optical tweezers, where the dynamic control on the traps can be used to rotate the trapped RBC about the Raman excitation beam in real time.

We report the results of a study conducted to record Raman spectra of an optically trapped RBC, where its equatorial plane was oriented at angles ranging from 0 deg to 180 deg with respect to the polarization direction of the excitation beam using a dual trap generated by holographic optical tweezers. We observed significant angular dependence for intensities of various Raman peaks. These experimental results and the theoretical simulations carried out to comprehend these results suggested that the Hb molecules within the cells may be present in an ordered arrangement, such that the heme-porphyrin planes acquire a preferential orientation parallel to the RBCs’ equatorial plane.

2 Materials and Methods

An integrated Raman-holographic optical tweezer was used to perform polarized Raman studies on single-trapped cells. Figure 1 shows a schematic of the experimental setup. A plane-polarized diode laser XTRA, TOPTICA Photonics AG, Graefelfing (Munich), Germany at 785 nm was used as the Raman excitation beam. For all experiments, the Raman spectra were recorded with an excitation power of 2 mW and an acquisition time of 30 s. The 785-nm laser beam was reflected off of a half-wave plate and polarizer placed before the spectrometer. For the polarized measurement, a combination of a polarizer and a analyzer. (b) A rendering showing the side view of a trapped RBC. The two holographic 1064-nm beams present near the edge of the cell trap and rotate the cell, whereas a 785-nm beam passing through the center of the cell excites the Raman spectrum. (c) An image of the trapped RBC captured by the CCD (Video 1, MPEG, 0.5 MB) [URL: http://dx.doi.org/10.1117/1.JBO.19.8.087002.1]. The red dots represent the 1064-nm beams, and the orange dot and arrow represent the Raman excitation beam and its electric field vector direction, respectively.

![Figure 1](attachment:image.png)
measurements, the half-wave plate was rotated by 45 deg. Sequential measurements were made on the two polarization components, because use of additional optics to separate the two polarization components for simultaneous measurements would require larger Raman excitation beam power, raising a concern for optically induced damage in the trapped cell. With the use of a high NA objective, induced depolarization effects are a concern. Therefore, we coupled the Raman excitation beam directly into the objective and did not utilize the full NA of the objective. Nevertheless to check the possible depolarization arising from the use of a high NA objective, the depolarization ratio of a chloroform line was measured using the 60×, 1.42, and 0.4 NA objectives. With the 1.42 NA objective, the depolarization ratio of the chloroform line at 667 cm⁻¹ was found to increase by ~3% when compared with the ratio obtained with the 0.4 NA objective. These results are consistent with the previous report, where an objective of NA 1.2 was used and depolarization effects were measured to be 1 to 2%.

Raman spectra were recorded from the cell in the spectral range of ~600 to ~2100 cm⁻¹ with a spectral resolution of about 6 cm⁻¹. The background was subtracted from each raw spectrum to obtain the actual cell spectrum. All Raman spectra were smoothed using Savitzky–Golay filters and the background was corrected using Lieber and Mahadevan–Jansen iterative polynomial fitting method.

Blood was collected by venipuncture from three healthy volunteers in glass tubes containing EDTA (5.4 mg/3 ml) as an anticoagulant. RBCs were separated from these blood samples by centrifugation at 3000 rpm for 3 min, and the separated RBCs were washed twice with phosphate buffer saline (PBS) by centrifugation at 3000 rpm for 3 min. A dilute suspension of RBCs in PBS containing 2% bovine serum albumin was used in the experiments.

### 3 Theory

The Raman spectrum of the RBCs is mostly contributed by the Hb molecules, the main cytoplasmic constituents of RBCs. Hb protein is an assembly of four globular protein subunits, each of which is composed of a protein chain tightly associated with a nonprotein heme group. A heme group is an iron-porphyrin complex in which an iron ion (Fe²⁺) lies in the center of a porphyrin macrocycle. Four of the six coordination sites around the iron are occupied by nitrogen atoms from the planar porphyrin ring. The fifth (axial) coordination site is occupied by a nitrogen atom from a histidine side chain on one of the amino acids in the protein. The last (axial) coordination site is available for binding with an O₂ molecule. Due to its planar and symmetric structure, the heme group has been approximated by a tetrahedral point group symmetry in literature and its vibrational modes have been accordingly assigned.

It may be noted that, due to the presence of heme charge transfer absorption band in the near-infrared region (NIR), some nontotally symmetric modes of B₁g symmetry are more enhanced in the NIR-excited Raman spectrum of the RBCs. Table 1 lists the band assignments, local coordinates, and symmetry terms for the prominent oxymyglobin Raman bands, observed with 785-nm excitation. In this table, the bands to which a symmetry term is assigned are due to the porphyrin ring vibrations and the bands with which no symmetry term is associated are due to the contribution from the porphyrin peripheral substituents such as vinyl group vibrations [ν(C˷a C˷b)] at ~1620 and ~1604 cm⁻¹ as well as the contribution from the surrounding globin proteins such as the phenylalanine peak at ~1003 cm⁻¹ and bending vibrations of CH₃/CH₂ groups at ~1450 cm⁻¹.

To gain insight into the RBCs’ orientation-dependent changes in the Raman spectra, we consider a hypothetical RBC which is packed with Hb molecules such that the planes of all the heme groups are oriented parallel to the RBCs’ equatorial plane. Although this hypothesis is a bit crude, the experimental observations indicate that this hypothesis may serve as a helpful starting point for analyzing the experimental results.

In Fig. 2, we show the chemical structure of the heme group and the set of axes used for the Raman tensors and intensity calculations, X, Y, and Z are the molecular axes, and x, y, and z are the laboratory axes. The Raman excitation beam is polarized along the x-direction. The z-axis is the direction of propagation of the Raman excitation beam. φ is the angle between the RBC equatorial plane and the Raman polarization direction.

The intensity of a Raman band depends on the orientation of the molecule and the polarization geometry of the optical configuration and can be expressed as

\[ I \propto |e_i \cdot \mathbf{M} e_s|^2, \]

where \( e_i \) and \( e_s \) are the unit polarization vectors of the electrical field of the incident and scattered laser beams, respectively, and \( \mathbf{M} \) is the Raman scattering tensor of the particular band under consideration. In our experiment, the polarization of the incident light was fixed in the x-direction and Raman scattered light polarized parallel and perpendicular to the excitation beam.

### Table 1: Band positions, assignments, local coordinates, and symmetry terms for the prominent oxymyglobin Raman bands

<table>
<thead>
<tr>
<th>Band position (cm⁻¹)</th>
<th>Assignment</th>
<th>Local coordinate</th>
<th>Symmetry</th>
</tr>
</thead>
<tbody>
<tr>
<td>1639</td>
<td>ν₁₀</td>
<td>ν(C˷a C˷b)</td>
<td>B₁g</td>
</tr>
<tr>
<td>1620</td>
<td>ν(C˷C˷)</td>
<td>ν(C˷aC˷b)</td>
<td>B₁g</td>
</tr>
<tr>
<td>1604</td>
<td>ν(C˷C˷)</td>
<td>ν(C˷aC˷b)</td>
<td>B₁g</td>
</tr>
<tr>
<td>1583</td>
<td>ν₃₇</td>
<td>ν(C˷aC˷b)</td>
<td>E₀</td>
</tr>
<tr>
<td>1563</td>
<td>ν₂</td>
<td>ν(C˷C˷)</td>
<td>A₁g</td>
</tr>
<tr>
<td>1547</td>
<td>ν₁₁</td>
<td>ν(C˷C˷)</td>
<td>B₁g</td>
</tr>
<tr>
<td>1448</td>
<td>δ(CH₂/CH₃)</td>
<td>δ(CH₂/CH₃)</td>
<td></td>
</tr>
<tr>
<td>1397</td>
<td>ν₂₀</td>
<td>ν(pyr half ring)</td>
<td>A₁g</td>
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<td>1371</td>
<td>ν₄</td>
<td>ν(pyr half ring)</td>
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</tr>
<tr>
<td>1337</td>
<td>ν₄1</td>
<td>ν(pyr half ring)</td>
<td>E₀</td>
</tr>
<tr>
<td>1266</td>
<td>ν₁₃ + ν₄₂</td>
<td>δ(CₚH)</td>
<td>B₁g or E₀</td>
</tr>
<tr>
<td>1212</td>
<td>ν₅ + ν₁₈</td>
<td>δ(CₚH)</td>
<td>A₁g + B₁g</td>
</tr>
<tr>
<td>1127</td>
<td>ν₁₅</td>
<td>ν(C˷Cm-methyl)</td>
<td>B₁g</td>
</tr>
<tr>
<td>1003</td>
<td>Phenylalanine</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Table 1: Band positions, assignments, local coordinates, and symmetry terms for the prominent oxymyglobin Raman bands.
polarization was detected. The unit polarization vectors can then be expressed as
\[ e_i = \begin{pmatrix} 1 \\ 0 \\ 0 \end{pmatrix}, \quad e_s^\parallel = \begin{pmatrix} 1 \\ 0 \\ 0 \end{pmatrix}, \quad e_s^\perp = \begin{pmatrix} 0 \\ 1 \\ 0 \end{pmatrix}, \]
where the superscript symbols \( \parallel \) and \( \perp \) refer to parallel and cross configurations, respectively. \( \mathbf{R} \) is the Raman scattering tensor of the particular band under consideration. In our experimental results, the significant orientation-dependent intensity changes were observed for the bands at \(~754\), \(1547\), \(1563\), and \(1621\) cm\(^{-1}\). Out of these, the bands at \(~754\) and \(1547\) cm\(^{-1}\) belong to the \(A_{1g}\) symmetry and the band at \(~1563\) cm\(^{-1}\) belongs to the \(A_{1g}\) symmetry. The band at \(1621\) cm\(^{-1}\) is due to \(\text{C} = \text{C}\) stretching vibrations of the porphyrin peripheral vinyl group substituents and does not belong to the porphyrin skeleton to which the \(D_{4h}\) point group symmetry has been assigned. Therefore, this band, not having any symmetry assignment, is excluded from our theoretical treatment. The second rank Raman tensor for \(A_{1g}\) and \(B_{1g}\) vibrational modes of a porphyrin ring having a \(D_{4h}\) point group symmetry can be expressed as\(^{29}\)
\[ \mathbf{R}_{A_{1g}} = \begin{pmatrix} a & 0 & 0 \\ 0 & a & 0 \\ 0 & 0 & b \end{pmatrix} \quad \text{and} \quad \mathbf{R}_{B_{1g}} = \begin{pmatrix} c & 0 & 0 \\ 0 & -c & 0 \\ 0 & 0 & 0 \end{pmatrix}. \]  

These tensors are referred to the principal molecular axes \(X\), \(Y\), and \(Z\), as has already been defined, and can be expressed in the laboratory axes system as follows:
\[ \mathbf{R}_{A_{1g}XYZ} = \Phi_{XYZ} \mathbf{R}_{A_{1g}} \Phi_{XYZ} \quad \text{and} \quad \mathbf{R}_{B_{1g}XYZ} = \Phi_{XYZ} \mathbf{R}_{B_{1g}} \Phi_{XYZ}, \]  
where \(\Phi_{XYZ}\) is an orthogonal transformation matrix, and \(\Phi_{XYZ}\) is the inverse, with direction cosines expressed in terms of Euler’s angles. These matrices transform the Cartesian system of coordinates associated with the molecular frame into that of the laboratory frame. For our experimental geometry,
\[ \Phi = \begin{pmatrix} \cos \phi & 0 & \sin \phi \\ -\sin \phi & 0 & \cos \phi \end{pmatrix}. \]  

From Eqs. (1)–(5), the Raman intensities pertaining to the \(A_{1g}\) and \(B_{1g}\) bands in parallel configuration can be expressed as a function of the angle \(\phi\) as follows:
\[ I_{A_{1g}} \parallel \propto |a \cos^2 \phi + b \sin^2 \phi|^2 \quad \text{and} \quad I_{B_{1g}} \parallel \propto |c \cos^2 \phi|^2. \]  

Similarly, for a perpendicular configuration,
\[ I_{A_{1g}} \perp \propto \left| \frac{(a - b)}{2} \sin 2\phi \right|^2 \quad \text{and} \quad I_{B_{1g}} \perp \propto \left| \frac{c}{2} \sin 2\phi \right|^2. \]  

The proportionality sign in Eqs. (6) and (7) arises from the dependence of the Raman intensity on the instrumental configuration used in the experimental setup.

4 Results
A total of 60 cells (20 cells each from the three blood samples) were investigated. Figures 3(a) and 3(b) present a mean spectrum of these cells for their different orientations. The orientation angle was measured as the angle between the RBCs’ equatorial plane and the excitation beam electric field vector. It can be seen from the figures that the intensity of various bands in the spectra shows the orientation-dependent changes. The most prominent orientation-dependent intensity changes occur in the band at \(~754\) cm\(^{-1}\), which is due to the breathing vibrations of the pyrrole rings, and in the spectral region \(1500\) to \(1700\) cm\(^{-1}\), which arises from the \(\text{C} = \text{C}\) bonds’ vibrations in the porphyrin ring and side chains. The intensities of these bands decreased as the RBC’s orientation angle was changed from 0 deg to 45 deg and was lowest at a 90 deg orientation angle. Upon further rotation, these changes reversed and the intensity of the bands again became highest at the 180 deg orientation. The spectra at 0 deg and 180 deg orientations were observed to be similar to each other as was expected. Further, the similarity of the spectra at the 0 deg and 180 deg orientations also ensures
that the cell was not photodamaged during the course of these measurements.

Another interesting observation was that while all the cells showed a decrease in the intensity of the above-mentioned Raman bands, a large number (39 cells, ~65%) of the cells showed a relatively greater decrease in the intensity of the band at \( \sim 1547 \text{ cm}^{-1} \) as compared with its adjacent band at \( \sim 1563 \text{ cm}^{-1} \) at a 90 deg orientation. This is shown in Fig. 4, where the Raman spectrum from a representative cell exhibiting this feature is presented. The reason for the differential response of the cells may be the inherent intercell variability in the cell age, shape, and Hb concentrations and oxygenation level. Figures 5(a) and 5(b) plot the average parallel-polarized Raman spectra of these 39 cells at the different orientations, and the differences between the spectra of these 39 cells and the total 60 cells are shown in Figs. 5(c) and 5(d). It can be noted that, while in the spectra shown in Fig. 4, almost all the bands show some orientation-dependent intensity changes, the average spectra shown in Figs. 5(a) and 5(b) show the prominent orientation-dependent changes only in the spectral region 1500 to 1700 cm\(^{-1}\) and in the band at \( \sim 754 \text{ cm}^{-1} \). The absence of the orientation-dependent changes in the relatively weaker bands may be due to their poor SNR not allowing inferring any reliable information from these bands.

To quantify the orientation-dependent changes observed in the Raman spectra, the peak intensities of the bands showing a prominent orientation-dependent change were calculated. Since the Raman band at \( \sim 754 \text{ cm}^{-1} \) is a well-isolated band [Fig. 6(a)], the peak intensities of this band were measured directly from the spectrum. However, the region 1500 to 1700 cm\(^{-1}\) is composed of eight bands: \( \nu_{38}, \nu_{11}, \nu_{10}, \nu_{7}, \nu_{27}, \nu_{38}, \nu_{11}, \nu_{10}, \nu_{7}, \nu_{27} \). Since these bands are closely spaced and strongly overlapped, their intensity cannot be directly determined from the spectra. To determine the peak positions and intensities of these bands, spectral deconvolution was performed. For this, eight Lorentzian peaks with intensity, width, and position as fitting parameters were used to obtain the best fit profile for the experimental spectral data. The value of \( R^2 \), which is an estimate of goodness of fit, was ~0.9. The fitted trace and the mean spectral data corresponding to 0 deg and 90 deg orientations are shown in Figs. 6(b) and 6(c). It can be seen that the fitted trace fit the observed data quite well. In Figs. 6(d)–6(g), the peak heights of the band at \( \sim 754 \text{ cm}^{-1} \) and some of the deconvoluted bands in the 1500 to 1700 cm\(^{-1}\) region, which showed statistically significant orientation-dependent intensity changes, are plotted as a function of the cell-orientation angle. To check for the statistical significance of the intensity changes observed for different orientation angles, a two-sample student \( t \)-test was performed. The bands at \( \sim 754 \) and \( \sim 1547 \text{ cm}^{-1} \) showed significant differences \( (p < 0.05) \) between the peak intensities observed for consecutive orientation angles [Figs. 6(d) and 6(e)]. However, the bands at \( \sim 1563 \) and \( \sim 1621 \text{ cm}^{-1} \) showed a significant difference \( (p < 0.005) \) only between the peak intensities observed for 0 deg (180 deg) and 90 deg orientations [Figs. 6(f) and 6(g)]. It may be noted here that no normalization was performed on the Raman spectra and the inherent intercell variation in Hb concentration as well as oxygenation level may be the reason for the large spread of the peak intensity values around their mean peak intensities.
present in Figs. 6(d)–6(g). Table 2 lists the mean peak intensities of the Raman bands whose angular dependence is plotted in Figs. 6(d)–6(g). It can be noted that of the four bands, the bands at ∼754 and 1547 cm$^{-1}$ undergo the largest change in intensity as the cells rotate from 0 deg to 90 deg orientation.

Next, we present the results of perpendicular-polarized Raman measurements performed on RBCs in their different orientations. It may be noted here that to avoid any photodamage effects that may occur due to prolonged laser irradiation, the parallel and perpendicular polarization measurements were made on different populations of the RBCs from the same blood samples. Figure 7 shows the perpendicularly-polarized Raman spectra at different cell orientations averaged over 60 cells (20 cells investigated from each blood sample). It can be seen from the figure that for this polarization, there are no significant orientation-dependent intensity changes in the spectra.

5 Discussions

From the results presented in Figs. 4–6, it can be noted that the intensities of the various bands in the parallel-polarized Raman spectra undergo significant changes as the cell is rotated to make different orientation angles relative to the excitation beam polarization direction. In particular, the intensity of the bands in the spectral region 1500 to 1700 cm$^{-1}$ and at ∼754 cm$^{-1}$ become highest when the RBCs' equatorial plane is parallel to the excitation beam polarization direction, i.e., 0 deg and 180 deg orientation, and lowest when the RBC plane is perpendicular to the excitation beam polarization direction, i.e., 90 deg orientation.

If the Hb molecules inside the RBCs are in totally disordered arrangement as in a solution, then the Raman spectrum of the RBC should not have any dependence on its orientation relative to the excitation beam polarization direction. Thus, the anisotropic behavior of Raman spectra provides strong evidence that the Hb molecules inside the cell must be in some ordered arrangement. It is pertinent to note here that the bands showing orientation-dependent intensity changes are the ones arising out of heme vibrations, whereas the bands contributed by the surrounding protein, such as the CH$_2$∕CH$_3$ bending vibration band at ∼1450 cm$^{-1}$ and the very prominent phenylalanine band at ∼1003 cm$^{-1}$, do not show any significant orientation-dependent changes. This suggests that it is the arrangement of heme groups within the RBCs that is responsible for the anisotropic spectral response of the RBCs. Because a heme molecule is planar, it is highly anisotropic and is expected to have maximum polarizability when the Raman excitation polarization direction is parallel to its plane. The experimental observation that the intensity of the prominent heme bands become highest when the RBCs' equatorial plane is parallel to the excitation beam polarization, i.e., 0 deg and 180 deg orientation, and lowest when it is
perpendicular to the excitation beam polarization direction, i.e., 90 deg orientation, suggests that the heme molecules may have some preferential orientation parallel to the RBCs’ equatorial plane.

Since the heme groups are embedded within the globin subunits of the Hb, any ordering of the heme group would also imply the ordering of Hb molecules. For such an ordered Hb arrangement, one would expect an anisotropic response from globin units as well. However, the molecular subgroups in the globin contributing to the Raman spectra, such as phenylalanine rings, are present in relatively high abundance, ~7 phenylalanine residues per globin subunit,31 and at different orientations with respect to the porphyrin plane32 (Protein Data Bank ID: 1HHO) within the quite complex structure of the globin folds, which might have resulted in randomization of the anisotropic response of individual phenylalanine rings and hence the isotropic response.

It is worthwhile to compare the results of our study with the polarized Raman spectroscopy (PRS) study carried out by Wood

![Fig. 6](https://www.spiedigitallibrary.org/journals/Journal-of-Biomedical-Optics)
et al. They compared the average unpolarized spectra recorded from the two populations: population 1 and population 2 (as described in Sec. 1) and analyzed the spectral changes using principal components (PC) analysis. They observed that the PC1, which explained 75% of the variance, mainly reflected the differences related to laser-induced effects and oxygen saturation levels of various RBCs and PC2 accounting for 13% of the variance exhibited for differences related to the orientation of the cell. From PC2, they suggested that the bands at ~1544, 1561, 1588, 1605, 1619, and 1634 cm\(^{-1}\) were most sensitive to the RBC orientation. The bands at ~1605, 1588, 1561, and 1544 cm\(^{-1}\) were found to be more intense in population 2, while the bands at 1634 and 1619 cm\(^{-1}\) were more intense in population 1. It may be noted here that these bands belong to the porphyrin skeletal vibration modes spectral region, i.e., 1500 to 1700 cm\(^{-1}\), and in our study most prominent changes were observed in this spectral region, too. A more direct comparison between the two studies is not possible mainly because of two reasons: (1) Wood et al. made a comparison between the unpolarized averaged spectra recorded from the two populations of RBCs, whereas we have investigated the polarized Raman spectra as a function of cell orientation. Although Wood et al. also performed polarized measurements, they did not make a comparison between the polarized spectra observed for the two populations. (2) In population 1, the RBCs had their equatorial planes parallel to the beam propagation direction while remaining parallel to the beam polarization direction. However, in population 2, the equatorial planes of the RBCs could take any orientation relative to the beam polarization direction while remaining parallel to the beam propagation direction. Therefore, the spectral response of a fraction of the cells from population 2 could overlap with the response of cells from population 1, whereas in our study there was no such overlapping between the spectra recorded at different orientations of the RBCs.

It may be noted here that the arguments presented in the preceding paragraphs to relate the experimentally observed anisotropic Raman response of the RBCs to the heme ordering within the RBC are very qualitative and cannot explain the subtle orientation levels of various RBCs and PC2 accounting for 13% of the variance exhibited for differences related to the orientation of the cell. From PC2, they suggested that the bands at ~1544, 1561, 1588, 1605, 1619, and 1634 cm\(^{-1}\) were most sensitive to the RBC orientation. The bands at ~1605, 1588, 1561, and 1544 cm\(^{-1}\) were found to be more intense in population 2, while the bands at 1634 and 1619 cm\(^{-1}\) were more intense in population 1. It may be noted here that these bands belong to the porphyrin skeletal vibration modes spectral region, i.e., 1500 to 1700 cm\(^{-1}\), and in our study most prominent changes were observed in this spectral region, too. A more direct comparison between the two studies is not possible mainly because of two reasons: (1) Wood et al. made a comparison between the unpolarized averaged spectra recorded from the two populations of RBCs, whereas we have investigated the polarized Raman spectra as a function of cell orientation. Although Wood et al. also performed polarized measurements, they did not make a comparison between the polarized spectra observed for the two populations. (2) In population 1, the RBCs had their equatorial planes parallel to the beam propagation direction while remaining parallel to the beam polarization direction. However, in population 2, the equatorial planes of the RBCs could take any orientation relative to the beam polarization direction while remaining parallel to the beam propagation direction. Therefore, the spectral response of a fraction of the cells from population 2 could overlap with the response of cells from population 1, whereas in our study there was no such overlapping between the spectra recorded at different orientations of the RBCs.

It may be noted here that the arguments presented in the preceding paragraphs to relate the experimentally observed anisotropic Raman response of the RBCs to the heme ordering within the RBC are very qualitative and cannot explain the subtle differences observed in the magnitude of the intensity modulation for the adjacent bands at ~1547 cm\(^{-1}\) (\(\nu_{11}\)) and at ~1563 cm\(^{-1}\) (\(\nu_{2}\)) and also the observation of no significant orientation-dependent intensity change in the perpendicularly polarized Raman spectra (Fig. 7). To explain these observations, a more quantitative approach may be required. Since the heme molecules can be approximated by \(D_{4h}\) point group symmetry, a theoretical model based on this symmetry was used to comprehend the experimental observations. In the model, a hypothetical RBC consisting of the planes of all heme molecules oriented parallel to the RBCs’ equatorial plane was considered. The cue for adopting such a configuration of heme molecules was taken from the observed experimental results, where the maximum intensity of heme bands was obtained when the RBC equatorial plane was parallel to the excitation beam polarization direction. In this modeling, only the bands showing significant orientation-dependent intensity modulation and having symmetry assignment, i.e., bands at ~754, 1547, and ~1563 cm\(^{-1}\), were considered. It can be noted from Table 1 that the bands appearing at ~754 and 1547 cm\(^{-1}\) belong to \(B_{1g}\) and the band at 1563 cm\(^{-1}\) belongs to the \(A_{1g}\) symmetry species. For the hypothetical RBC, the dependence of intensity of the bands belonging to these symmetry species on the RBCs’ orientation angle would be governed by Eq. (6). For \(B_{1g}\) modes, the intensity varies as \(|c \cos^2 \phi|^2\), where \(\phi\) is the orientation angle, and thus these modes exhibit a periodicity of 180 deg with a maximum intensity at 0 deg and 180 deg orientations and a minimum intensity at 90 deg. The experimental and theoretical results for orientation-dependent intensity variation for the 1547 cm\(^{-1}\) band are shown in Fig. 8. The theoretical results predict a minimum intensity value of zero at a 90 deg orientation, which can only be true in the case of perfectly oriented molecules. The agreement between the theoretical and experimental values appears to be very good.

For \(A_{1g}\) modes, the intensity is predicted to vary as \(|a \cos^2 \phi + b \sin^2 \phi|^2\), where \(a\) and \(b\) are the in-plane and out-of-plane components of the Raman tensor. For a perfectly planar molecule, ratio \(b/a\) is expected to be very small and can be ignored. However, for heme molecules to be perfectly planar molecules, the requirement is that both the axial ligands should be same and of perfect axial symmetry and the porphyrin macrocycle should be symmetrically substituted. None of these conditions are fulfilled exactly for the heme molecules and therefore the \(b\)-component cannot be neglected. Since the angle-dependent variations of the \(a\)- and \(b\)-components tend to cancel each other, the presence of a small \(b\)-component will act to reduce the magnitude of the intensity change that would have occurred if only the \(a\)-component was present. Therefore,
we may expect the $A_{1g}$ modes to show a similar angle-dependent behavior as that for $B_{1g}$ modes, but with a reduced amplitude modulation. This may be the reason for the experimental observation of a smaller change in the intensity at $\sim 1563$ cm$^{-1}$ ($A_{1g}$) as compared with the band at $\sim 1547$ cm$^{-1}$ ($B_{1g}$).

Further, it can be noted from the Eq. (7) that for both $A_{1g}$ and $B_{1g}$ modes, the intensity of the Raman scattered light polarized perpendicular to the excitation beam polarization direction varies as $\sin^2 \theta$, i.e., with a periodicity of 90 deg and become maximum at 45 deg and minimum at 0 deg and 90 deg orientations. However, the magnitude of variation is only ~one-fourth of that for the parallel-polarized case. This may be the reason for not observing any significant orientation-dependent intensity changes in the perpendicularly polarized Raman spectra.

Thus, the results of the measurement on the orthogonal-polarized component of Raman scattered light provide strong evidence that the heme molecules within the RBC have a preferential orientation parallel to the RBCs’ equatorial plane. However, a smaller amplitude modulation (~12%) observed for experimental data (Fig. 7) as compared with the theoretical prediction (100%) suggests that the degree of orderliness may be very small. The fact that the four heme groups within each oxyhemoglobin tetramer are approximately parallel to each other further strengthens the possibility that for some arrangement of Hb molecules, the heme groups of neighboring Hb molecules may become parallel to each other. Thus, the neighboring Hb molecules inside the RBCs may frequently acquire an arrangement which makes the heme molecules parallel to RBCs’ equatorial plane.

To conclude, we have observed that the parallel-polarized Raman spectrum of an RBC depends upon its orientation relative to the excitation beam polarization direction. Theoretical simulations carried out to comprehend these observations suggest that in RBC, the Hb molecules within the cells must be present in an ordered arrangement, such that heme-hemoporphyrin planes become preferentially oriented parallel to the RBCs’ equatorial plane, although the degree of orderliness appears to be very small.

References


Biographies of the authors are not available.