

RESEARCH OUTPUTS / RÉSULTATS DE RECHERCHE

Analysis of accuracy and ambiguities in spatial measurements of birefringence in uniaxial anisotropic media

Bouhy, Julie; Dekoninck, Augustin; Voué, Michel; Deparis, Olivier

Published in:
Applied Optics

DOI:
[10.1364/ao.463657](https://doi.org/10.1364/ao.463657)

Publication date:
2022

Document Version
Peer reviewed version

[Link to publication](#)

Citation for published version (HARVARD):

Bouhy, J, Dekoninck, A, Voué, M & Deparis, O 2022, 'Analysis of accuracy and ambiguities in spatial measurements of birefringence in uniaxial anisotropic media', *Applied Optics*, vol. 61, no. 27, pp. 8081-8090. <https://doi.org/10.1364/ao.463657>

General rights

Copyright and moral rights for the publications made accessible in the public portal are retained by the authors and/or other copyright owners and it is a condition of accessing publications that users recognise and abide by the legal requirements associated with these rights.

- Users may download and print one copy of any publication from the public portal for the purpose of private study or research.
- You may not further distribute the material or use it for any profit-making activity or commercial gain
- You may freely distribute the URL identifying the publication in the public portal ?

Take down policy

If you believe that this document breaches copyright please contact us providing details, and we will remove access to the work immediately and investigate your claim.

Analysis of accuracy and ambiguities in spatial measurements of birefringence in uniaxial anisotropic media

BOUHY JULIE,^{1,*} AUGUSTIN DEKONINCK,² VOUE MICHEL,³ AND DEPARIS OLIVIER¹

¹*Department of Physics, Namur Institute of Structured Matter, University of Namur, 61 Rue de Bruxelles, 5000 Namur, Belgium*

²*Department of Geology, Institute of Life-Earth-Environment, University of Namur, 61 Rue de Bruxelles, 5000 Namur, Belgium*

³*Physics of Materials and Optics Units, Research Institute for Materials Science and Engineering, University of Mons, 20 Place du Parc, 7000 Mons, Belgium*

**julie.bouhy@unamur.be*

Abstract: Accuracy and ambiguities in retardance and optical axis orientation spatial measurements are analyzed in details in the context of the birefringence imaging method introduced by Shribak and Oldenbourg [Appl. Opt. **42**,3009 (2003)]. An alternative formula was derived in order to determine the optical axis orientation more accurately, and without indetermination in the case of a quarter-wave plate sample. Following Shribak and Oldenbourg's experimental configuration using two variable retarders, a linear polarizer and five polarization probes, we examined the effect of the swing angle χ , which selected the ellipticity of each polarization state, on the accuracy of retardance (Δ) and axis orientation (ϕ) measurements. Using a quarter-wave plate, excellent agreement between measured and expected values was obtained for both the retardance and the axis orientation, as demonstrated by statistical analysis of Δ and ϕ spatial distributions. The intrinsic ambiguity in the determination of Δ and ϕ for superimposed layers of transparent anisotropic cello-tape is discussed in details and solutions are provided to remove this ambiguity. An example of application of the method on geological samples is also presented. We believe our analysis will guide researchers willing to exploit this long-standing method in their laboratories.

© 2022 Optica Publishing Group

1. Introduction

Birefringence is the property exhibited by uniaxial optical media, which present two refractive index values according to the polarization of light [1]. This phenomenon was first observed in 1669 in the Iceland spar (calcite) by Rasmus Bartholin. Birefringence origin can be either molecular or structural (form birefringence), or induced by mechanical strains, electric field (Pockels and Kerr effect) or magnetic field (Faraday effect) [1]. The measurement of birefringence or retardance is particularly significant not only in optical industry (quality control), but also in various domains such as medicine [2, 3], pharmacology [4] and geology [5]. In the field of photoelasticity [6], birefringence measurements give access to the distribution of mechanical stress in e.g. glass or plastics, which is of great interest for industrial or architectural applications. In the case of liquid crystals, retardance of which is controlled by the application of an electric field, birefringence must be accurately determined in order to provide the product with specifications, i.e calibration curves [7]. Also, birefringence measurements give direct information about solid-state phase transition [8]. Linear birefringence can be measured by means of different techniques such as interferometry [9], compensation [10, 11], polarimetry [12, 13] or modulation [8, 14]. In 2003, Shribak and Oldenbourg introduced an original technique for birefringence imaging [15], which

45 they further developed [16] and extended to diattenuation measurement [17]. Basically, the 2D
 46 spatial distributions of both the retardance and the optical axis orientation are determined over
 47 the beam spot-size. These techniques were commercialised under the trademark Polscope [18].
 48 Among the various versions of Polscope, we refer herein to the most precise version that uses two
 49 variable retarders, a linear polarizer and five polarization states to probe an anisotropic uniaxial
 50 transparent sample. As pointed by the authors, the method suffers from intrinsic ambiguities in
 51 determining retardance and axis orientation. Experimentally, it is worth recalling that the input
 52 polarization states is selected by the angle χ , which is called the swing angle [15]. The angle χ is
 53 a key parameter, since it tunes the ellipticity. In this article, we examined the influence of the χ
 54 parameter on measurement accuracy and found the latter could be improved by increasing χ value.
 55 We also investigated ambiguity issues and provided solutions to mitigate them. Hereafter, we
 56 first introduce the theoretical framework by recalling the basic principle of the method developed
 57 by Shribak and Oldenbourg and we derive an alternative formula for optical axis orientation
 58 measurements. In the next two sections, the experimental setup is presented as well as its
 59 calibration with commercial products. Then, on the basis of measurement results we obtained
 60 with various birefringent samples, we discuss ambiguity issues and propose solutions to mitigate
 61 them. Finally, we illustrate the usefulness of our method for the characterization of natural
 62 samples, in particular a composite quartz/tourmaline thin section.

63 2. Theoretical framework

64 Let us consider a general case, where a monochromatic polarized light beam propagates along
 65 the z axis into an anisotropic, uniaxial and inhomogeneous medium with varying orientations
 66 of its optical axis according to the lateral spatial position (Fig.1). We assume that the optical
 67 axis is everywhere parallel to both faces of the medium, i.e. it lies in the (x, y) plane. This
 68 hypothesis is fulfilled not only in polarizing optical components (retardation plates, etc.) but also
 69 in anisotropic films such as those resulting from the alignment of molecules or structures parallel
 70 to the surface. Therefore, in general, the medium exhibits a 2D spatial distribution of retardance
 $\Delta(x, y)$ and optical axis orientation $\phi(x, y)$.

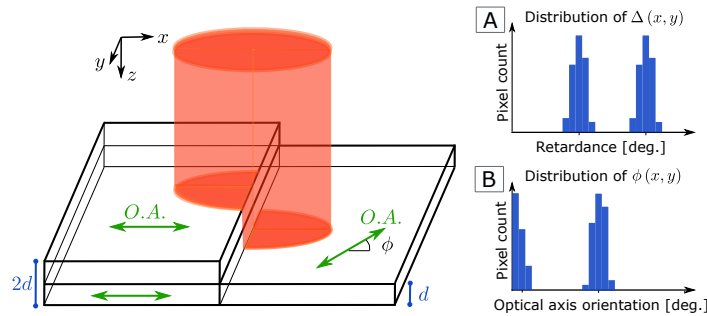


Fig. 1. Hypothetical samples, respectively composed of one layer (right) and two superimposed layers (left) of an anisotropic film of same retardance, have their optical axis oriented at different angles. (A) The retardance of the two superimposed layers is twice that of the single layer. (B) The optical axis orientations are 45° and 0° , respectively for one single layer and two superimposed layers. Blue bars around the mean values symbolize measurement inaccuracies or sample inhomogeneities.

71
 72 Five input polarization states are successively generated by a tunable elliptical polarizer in
 73 order to probe the medium of interest and calculate both its retardance and optical axis orientation.
 74 The calculation is based on the five-frame algorithm of Shribak and Oldenbourg, derived from

75 the Jones matrix formalism [19]. It is noteworthy to remind that, in the Jones formalism, light is
76 assumed to be entirely polarized.

77 The first input polarization state ψ_0 is circular whereas the four other ones (ψ_1, ψ_2, ψ_3 and ψ_4)
78 are elliptical, with the same ellipticity ϵ but with azimuthal angles ϑ equal to $0^\circ, 45^\circ, 90^\circ$ and
79 135° , respectively. The ellipticity ϵ of the input polarization states is controlled by the parameter
80 χ , as $\chi = 90^\circ - 2\epsilon$ [15]. Output polarization states, modified after propagation into the medium,
81 are analyzed by a circular analyzer, the handedness of which is opposite to the handedness of the
82 elliptical polarizer.

83 Any state of polarization can be decomposed in a basis of orthogonal polarization states
84 e.g. horizontal and vertical ones or left-handed and right-handed ones. Since circular and
85 elliptical polarization states were used, a circular polarization basis was chosen. Left-handed and
86 right-handed polarization states are given by the following Jones vectors $\vec{E}_L = \frac{1}{\sqrt{2}} [1, -i]^T$ and
87 $\vec{E}_R = \frac{1}{\sqrt{2}} [1, i]^T$, where T denotes transpose.

88 All polarization states can be expressed as a linear combination of these two polarization
89 states, with a different phase shift between the components. For instance, horizontal linear
90 polarization state is composed of in-phase left-handed and right-handed circular polarization
91 states: $\vec{E}_H = \vec{E}_R + \vec{E}_L$. Therefore, a right-handed circular analyzer will only select the right-handed
92 component of the linear polarization state and the transmitted normalized intensity will be equal
93 to 0.5. Left-handed polarization component will not be selected, so that the corresponding
94 normalized intensity will be equal to zero whereas right-handed one will be entirely selected,
95 giving a normalized intensity equal to one.

96 We can visualize states of polarization on the surface of the Poincaré sphere (Fig.2), where all
97 states lie in the absence of depolarization. Polarization states located on the poles are circular
98 and those lying on the equator are linear. All the other polarization states are elliptical. The
99 Southern hemisphere contains left-handed states and the Northern hemisphere right-handed ones.
100 On any latitude, the ellipticity is invariant and the azimuthal angle spans over a range from 0° to
101 360° . On any longitude, the azimuthal angle is fixed and the ellipticity spans from -45° to 45°
102 (the tangent of ellipticity is given by the ratio of the ellipse axes).

103 Let us take the handedness of the five input polarization states to be left and the handedness
104 of the circular analyzer to be right (or vice versa). By this choice, the polarization state ψ_0 is
105 located on the Southern pole, which is opposite to the right circular analyzer location, leading to
106 extinction configuration. The four other ones are located on the same latitude in the Southern
107 hemisphere. The latitude is determined by the angle χ , which will be taken as a free parameter
108 hereafter (Fig.2).

109 We now examine the influence of the χ parameter on the input polarization states and the
110 normalized intensity transmitted through the analyzer, in the absence of any medium. As we
111 work with a circular analyzer, only the ellipticity ϵ has an impact on the measured intensity,
112 independently of the azimuthal angle ϑ . Since the circular polarization state ψ_0 has the opposite
113 handedness to that of the circular analyzer, the intensity transmitted is equal to zero (i.e.
114 extinction), as mentioned earlier. The ellipticity of the four elliptical polarization states, given
115 by $\epsilon = 45^\circ - \frac{\chi}{2}$, determines the latitude on the Poincaré sphere. When χ is increased, the
116 polarization states ψ_1, ψ_2, ψ_3 and ψ_4 move away from the South pole. As a result, the transmitted
117 intensity measured by a detector is increased, hence the signal to noise ratio is enhanced.

118 The effect of retardation plates is to transform the polarization states at the output. For instance,
119 an half-wave plate (HWP) leads to a retardation of half a wavelength between ordinary and
120 extraordinary light waves. The initial left circular polarization state $\psi_0^{(i)}$ at input is transformed in
121 a final right circular polarization state $\psi_0^{(f)}$ at output, leading to full illumination of the detector,
122 i.e. normalized intensity equal to one. The four elliptical polarization states remain on the same
123 latitude, but are moved to the Northern hemisphere as the handedness is changed. As a result, the

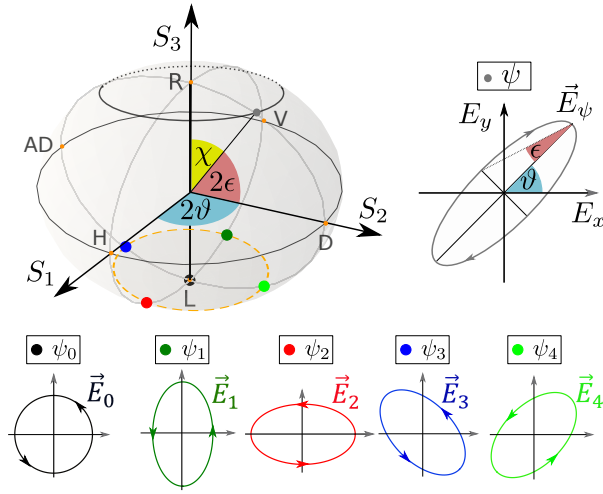


Fig. 2. All the polarization states on the surface of the Poincaré sphere are completely polarized. Those lying at the North and South poles are respectively right-handed circular (R) and left-handed circular (L), and those lying on the equator are linear. All the other polarization states are elliptical. Linear polarization states with azimuthal angle equal to 0° , 90° , 45° and 135° are horizontal (H), vertical (V), diagonal (D) and anti-diagonal (AD), respectively. The four elliptical polarization states ψ_1 , ψ_2 , ψ_3 and ψ_4 are located on the same latitude, which is determined by the χ parameter. The left-handed circular polarization state ψ_0 is located at the South pole. The ellipticity is defined by the angle ϵ and the ellipse inclination by the angle ϑ .

124 four intensities measured after the circular polarizer have to be the same. This particularity leads
 125 to an indetermination of the orientation of the optical axis of the half-wave plate (as it will be
 126 explained in section 3).

127 The intensity detected after the analyzer depends on the five input polarization states and the
 128 properties of the medium of interest, that is to say $\Delta(x, y)$ and $\phi(x, y)$. For the sake of simplicity,
 129 we consider a transparent medium, neglect depolarisation effects due to imperfect components
 130 (hardware dependent) and do not consider correction of background retardance obtained by
 131 removing the sample from the field of view [15]. These assumptions are equivalent, in eqs.7
 132 of [15], to setting the distribution of the depolarized background illumination $I_{\min}(x, y) = 0$,
 133 the distribution of the illumination intensity on the sample $I_{\max}(x, y) = 1$ and the isotropic
 134 transparency $\tau(x, y) = 1$. The intensities $I_j(x, y, \Delta, \phi)$, with j referring to the corresponding
 135 input polarization state $\psi_j^{(i)}$, measured at each location (x, y) by an imaging detector, are
 136 expressed by:

$$\begin{aligned}
I_0 &= I(\alpha = 90^\circ, \beta = 180^\circ) = \frac{1}{2} [1 - \cos \Delta], \\
I_1 &= I(\alpha = 90^\circ - \chi, \beta = 180^\circ) = \frac{1}{2} [1 - \cos \chi \cos \Delta + \sin \chi \sin (2\phi) \sin \Delta], \\
I_2 &= I(\alpha = 90^\circ + \chi, \beta = 180^\circ) = \frac{1}{2} [1 - \cos \chi \cos \Delta - \sin \chi \sin (2\phi) \sin \Delta], \\
I_3 &= I(\alpha = 90^\circ, \beta = 180^\circ - \chi) = \frac{1}{2} [1 - \cos \chi \cos \Delta - \sin \chi \cos (2\phi) \sin \Delta], \\
I_4 &= I(\alpha = 90^\circ, \beta = 180^\circ + \chi) = \frac{1}{2} [1 - \cos \chi \cos \Delta + \sin \chi \cos (2\phi) \sin \Delta],
\end{aligned} \tag{1}$$

137 where $\Delta \in [0, 180^\circ]$ and $\phi \in [0, 180^\circ]$. Here, α and β denote, respectively, the retardances of
138 two liquid crystals of the elliptical polarizer (see next section), which define the input polarization
139 state. Algebraic manipulation of these formula allows us to isolate Δ and ϕ , the two quantities
140 of interest. In order to simplify the expressions of Δ and ϕ , two intermediate quantities are
141 introduced (cf. eqs. 19 in [15]):

$$\begin{aligned}
A &\equiv \sin (2\phi) \tan \Delta = \frac{I_1 - I_2}{I_1 + I_2 - 2I_0} \tan \frac{\chi}{2}, \\
B &\equiv \cos (2\phi) \tan \Delta = \frac{I_4 - I_3}{I_3 + I_4 - 2I_0} \tan \frac{\chi}{2},
\end{aligned} \tag{2}$$

142 leading to the following expressions (cf. eqs. 20 in [15]):

$$\Delta = \begin{cases} \arctan \left(\sqrt{A^2 + B^2} \right), & \text{if } I_1 + I_2 - 2I_0 \geq 0, \\ 180^\circ - \arctan \left(\sqrt{A^2 + B^2} \right), & \text{if } I_1 + I_2 - 2I_0 < 0, \end{cases} \tag{3}$$

143

$$\phi = \frac{1}{2} \arctan \frac{A}{B}. \tag{4}$$

144 If we examine eqs. 1 and 2, we see that both denominators of A and B are equal: $I_1 + I_2 - 2I_0 =$
145 $I_3 + I_4 - 2I_0 = \cos \Delta (1 - \cos \chi)$. In order to determine the optical axis orientation, instead of
146 using eq. 4, we propose to use an alternative formula in which the common denominator of A
147 and B was simplified:

$$\phi = \frac{1}{2} \arctan \left(\frac{I_1 - I_2}{I_4 - I_3} \right). \tag{5}$$

148 The usefulness of this alternative expression will be highlighted later in this article. In the
149 particular case of a quarter-wave plate (QWP), the common denominator of A and B is equal
150 to zero. In this case, if we use eq. 4 to calculate the angle ϕ , the division by zero leads to an
151 indetermination.

152 The parameter χ controls the ellipticity of input polarization states and, therefore, the intensity
153 transmitted through the analyzer. We introduce here two additional parameters κ and η in order to
154 analyze the influence of χ on the intermediate quantities A and B used to calculate the retardance
155 and optical axis orientation. These parameters are defined by

$$\kappa \triangleq \sqrt{(I_1 - I_2)^2 + (I_4 - I_3)^2} = \sin \chi \sin \Delta, \tag{6}$$

156

$$\eta \triangleq I_1 + I_2 - 2I_0 = \cos \Delta (1 - \cos \chi). \tag{7}$$

157 The parameter κ allows us to quantify the separation between the intensities I_j according
158 to the retardance of the medium and the choice of the parameter χ . By dividing eq. 6 by eq.

159 7, we get $\tan \Delta = \frac{\kappa}{\eta} \frac{(1-\cos \chi)}{\sin \chi}$, which can be used as an alternative formula for determining the
 160 retardance. It is noteworthy that κ is equal to zero, both for a half-wave plate ($\Delta = 180^\circ$) and
 161 an isotropic medium ($\Delta = 0^\circ$) because the four output polarization states ($j = 1, 2, 3, 4$) lie on
 162 the same latitude on the Poincaré sphere. Therefore, all the four output intensities are equal, i.e.
 163 there is no separation between the intensities. In other words, when $\kappa = 0$, we face an ambiguity
 164 when distinguishing between an HWP or the absence of sample. The parameter κ monotonically
 165 rises with the increase of χ parameter when Δ spans from 0° to 90° whereas above $\Delta = 90^\circ$, κ
 166 decreases monotonically until it reaches zero at $\Delta = 180^\circ$ (Fig.3).

167 The parameter η defines the quadrant in which the retardance must lie according to eq. 3
 168 ($]0^\circ, 90^\circ[$ or $]90^\circ, 180^\circ[$) and is also used to discriminate between ambiguous retardance values,
 169 i.e. $\Delta = 0^\circ$ (isotropic medium) or $\Delta = 180^\circ$ (half-wave plate). In the case of isotropic medium,
 170 $\eta = (1 - \cos \chi)$ whereas for the half-wave plate, $\eta = (\cos \chi - 1)$. For a small χ value, e.g.
 171 $\chi = 10^\circ$, $\eta = \pm 0.015$ so that, in practice, it might be difficult to discriminate between an isotropic
 172 medium and an half-wave plate because of measurement noise. In such a situation, the choice of
 a higher χ value is recommended.

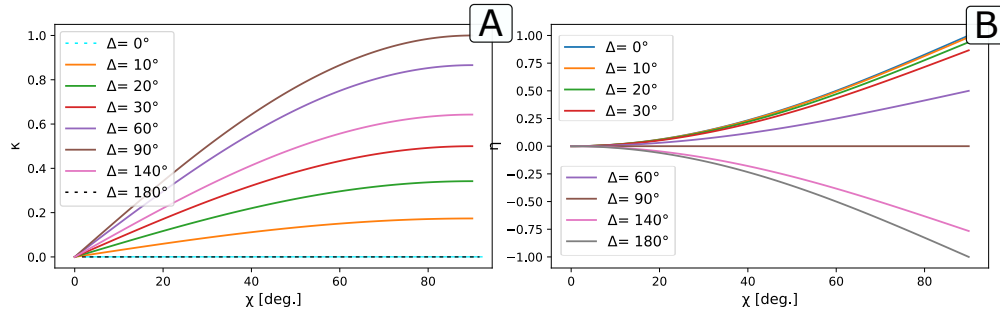


Fig. 3. A) The parameter κ , quantifying the separation between the measured intensities, is plotted as a function of parameter χ , for Δ between 0° and 180° . B) The parameter η , defining the quadrant in which Δ must lie, is plotted as a function of the parameter χ , for Δ between 0° and 180° .

173

174 3. Experimental setup

175 An experimental setup (Fig.4) was assembled from a HeNe laser (Thorlabs, Polarized HeNe
 176 Laser HNL150LB), an optical microscope (Olympus, BX53), light-polarising elements and a
 177 CCD camera (Olympus, SC50) in order to measure output light intensities corresponding to the
 178 five input polarization states and calculate the retardance and optical axis orientation distributions
 179 of the sample of interest.

180 Monochromatic light beam from the HeNe laser is collimated by two lenses and shaped by a
 181 pinhole. The collimated beam is elliptically polarized by a linear polarizer (Thorlabs, Glan-Taylor
 182 Polarizer) and two liquid crystal retardation plates LCA and LCB (Thorlabs, Uncompensated
 183 Half-Wave LC Retarder) with their optical axes respectively set at 0° and 45° . The retardance
 184 of both plates, noted respectively α and β , is controlled by applied voltage. The prepared
 185 polarization state is circular in the case of retardation values set to $\alpha = 90^\circ$ and $\beta = 180^\circ$.
 186 Otherwise, the ellipticity is induced by adding a positive or negative retardation χ to α and to β ,
 187 but not on both simultaneously. The four elliptical polarization states produced have all the same
 188 ellipticity $\epsilon = 45^\circ - \frac{\chi}{2}$, but different azimuthal angles equal to $0^\circ, 45^\circ, 90^\circ$ and 135° .

189 The beam, emerging from the elliptical polarizer, is reflected by a mirror tilted at 45° . It is
 190 noteworthy that the mirror reflection modifies the polarization and flips its handedness, so that

191 it is necessary to choose the voltage applied to liquid crystals in a way to compensate for the
 192 polarization modifications induced by the mirror. The handedness of input polarization states is
 193 chosen to be the same as the handedness of circular analyzer, since the mirror flips it.

194 The reflected beam is focused on the sample by the condenser. The objective lens enables
 195 a 10× magnification of the sample. The polarized light emerging from the sample impinges
 196 on the circular analyzer. The circular analyzer is composed of a quarter-wave plate (Thorlabs,
 197 Multi-Order Quarter-Wave Plate) and a linear polarizer (Olympus, U-AN360P), the optical axes
 198 of which are respectively oriented at 45° and 0°. Adjustment of the quarter-wave plate angle is
 199 carefully realized with a polarimeter (Thorlabs, Polarimeter PAX1000VIS/M) placed on the top
 200 of the microscope. For this purpose, the LCs are removed from the optical path so that the beam
 201 is linearly polarized. Once the optical axis of the quarter-wave plate is properly oriented at 45°,
 202 the transmitted polarization state becomes circular and the position of the QWP is settled. The
 203 linear polarizer is then added in the optical path, and its proper orientation is selected thanks to
 204 the polarimeter.

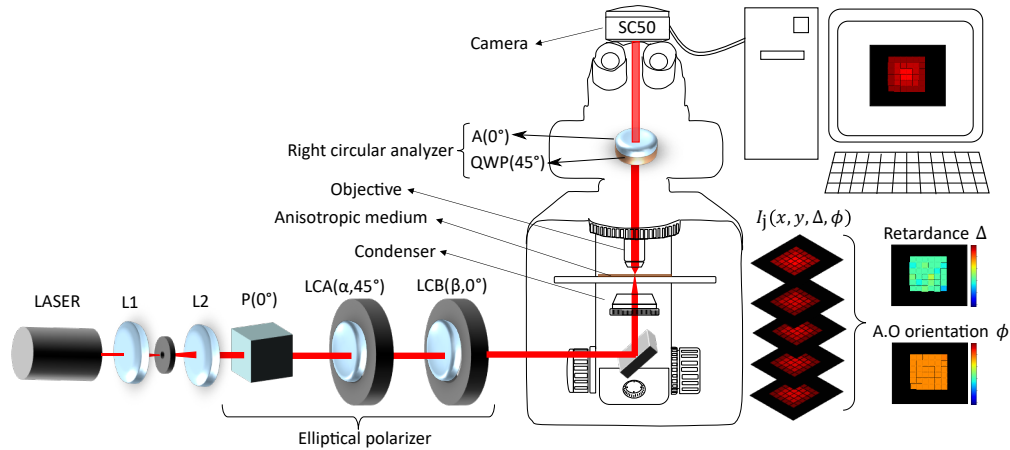


Fig. 4. Laser beam (632.8 nm) is collimated by two lenses (L_1, L_2) and elliptically polarized by a horizontal linear polarizer (P) and two liquid crystal retardation plates LCA and LCB, controlled by voltage. The optical axes of LCA and LCB are oriented at 45° and 0° respectively. The beam is reflected by a mirror oriented at 45°, focused by a condenser lens on the sample and transmitted throughout an objective lens (10×). The polarized beam is then directed towards the circular analyzer (A), composed of a quarter-wave plate and a linear polarizer (A) oriented so that the optical axis form an angle of 45°. The intensity of the beam at the output of the circular analyzer is measured by the CCD camera (SC50) of the microscope (BX53).

205 For each input polarization state $\psi_j^{(i)}$ where $j \in [0, 4]$, the intensity distribution $I_j(x, y; \Delta, \phi)$
 206 detected by 16-bit camera (2560×1920 pixels) is recorded for further processing. All
 207 measurements are made with a $\times 10$ objective lens and the image dimension is equal to
 208 $5632 \mu\text{m} \times 4224 \mu\text{m}$. The exposure time used to record the intensity distribution is extracted
 209 from image metadata for each configuration and used to calculate the intensity for an identical
 210 duration for all the configurations. This calculated intensity is directly used to determine the
 211 retardance and the orientation in each pixel of the image.

212 4. Calibration of measurement system

213 The calibration of the experimental setup consisted in determining the values of voltage to be
 214 applied to the liquid crystals in such a way to obtain the desired five input polarization states,
 215 with the ellipticity ϵ chosen beforehand. The calibration was carried out without sample and with
 216 two retardation plates as etalon: a quarter-wave plate (Thorlabs, WPQSM05-633) and half-wave
 217 plate (Thorlabs, WPHSM05-633). The quarter-wave plate allowed us to check the retardance
 218 and the orientation of the optical axis, whereas it was only possible to check the retardance of
 219 the half-wave plate, because of the indetermination on the orientation of the optical axis (see
 220 discussion below).

221 The retardation plate under test was placed at a particular angle and the control voltages applied
 222 to retardation liquid crystal plates were chosen to induce retardations equal to
 223 $[\alpha, \beta] \in \{(90^\circ, 180^\circ), (90^\circ - \chi, 180^\circ), (90^\circ + \chi, 180^\circ), (90^\circ, 180^\circ - \chi), (90^\circ, 180^\circ + \chi)\}$ (Fig.5).

224 Expected values for retardance of quarter-wave plate and half-wave plate are 90° and 180° ,
 225 respectively. Without sample, it must be measured $\Delta = 0^\circ$. In all cases, the recorded images
 must be ideally uniform.

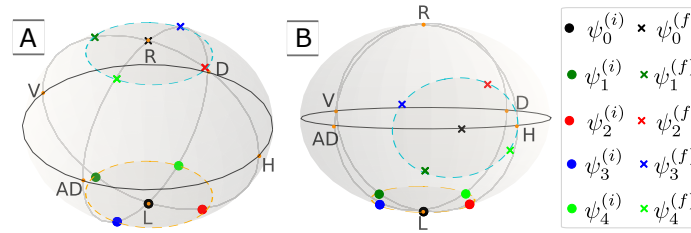


Fig. 5. The five input ($\psi_j^{(i)}$) and output ($\psi_j^{(f)}$) polarization states are depicted on the Poincaré sphere for A) HWP set at 0° B) QWP set at 120° (equivalent to -30°).

226 First order statistic analysis performed on Δ and ϕ values measured at $\chi = 60^\circ$ for the
 227 quarter-wave plate oriented at an angle equal to $\phi = -30^\circ$ gave mean values equal to $\langle \Delta \rangle = 89.77^\circ$
 228 and $\langle \phi \rangle = -30.17^\circ$, and standard deviations equal to $\sigma_\phi = 1.60^\circ$ and $\sigma_\Delta = 8.28^\circ$. The
 229 Gaussian-fitted probability density function (pdf) with standard deviation σ and mean value μ ,
 230 i.e. $f(x) = \frac{1}{\sigma\sqrt{2\pi}} \exp\left(-\frac{1}{2}\left(\frac{x-\mu}{\sigma}\right)^2\right)$, is plotted in Fig.6 for Δ and ϕ measured on the QWP oriented
 231 at -30° . The excellent agreement between measured and expected values for both the retardance
 232 and the optical axis angle validates our measurement procedure.

233 In the absence of sample or with the HWP, measurements were performed only for the
 234 retardance since the optical axis orientation suffers from intrinsic ambiguity. Mean values
 235 measured at $\chi = 60^\circ$ were equal to $\langle \Delta \rangle = 4.35^\circ$ and $\langle \Delta \rangle = 174.24^\circ$, respectively, and standard
 236 deviation values were equal to $\sigma_\Delta = 3.00^\circ$ and $\sigma_\Delta = 3.54^\circ$, respectively (Fig.6). In these
 237 cases, measured distributions departed from first-order statistics. As a result, peak values of the
 238 distributions did not match the mean values. The peak of the measured probability density was
 239 reached at $\Delta = 3^\circ$ in the absence of sample, and at $\Delta = 176^\circ$ with the HWP. The resolution limit
 240 of the experimental setup was therefore estimated to be around 4° , which corresponds to a noise
 241 level of $\Delta nd = \frac{\lambda}{2\pi} \frac{4\pi}{180} = 7.03 \text{ nm}$.

243 5. Results and discussion

244 Hereafter, we first examine the effect of the swing angle χ on measurement accuracy and then
 245 the ambiguities in determining optical axis orientation.

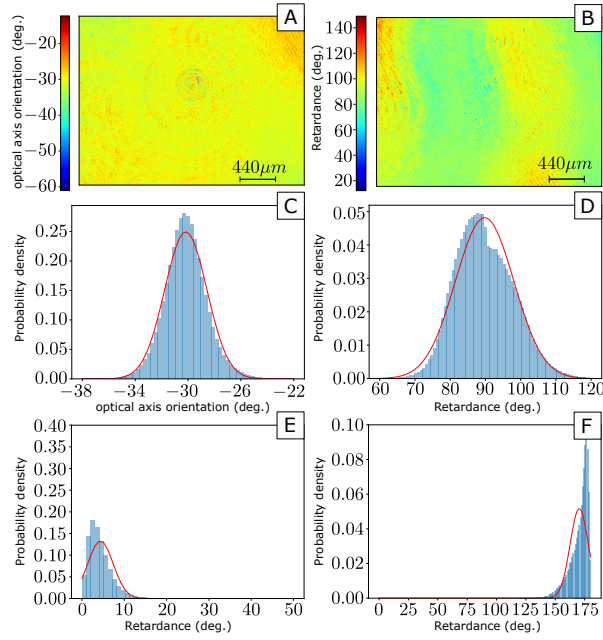


Fig. 6. Calibration with a QWP oriented at -30° (A-D), in the absence of sample (E) and with a HWP (F). The swing angle parameter of the measurement is $\chi = 60^\circ$. A) Optical axis spatial distribution for the QWP. B) Retardance spatial distribution for the QWP. C) Probability density of optical axis distribution for the QWP. D) Probability density of retardance distribution for the QWP. E) Probability density of retardance distribution in the absence of sample. F) Probability density of retardance distribution with an HWP oriented at 0° .

246 5.1. Effect of χ parameter on measurement accuracy

247 In order to highlight the effect of the parameter χ on the measurement accuracy, a retardation
 248 plate (Olympus, U-TP137) was chosen to measure both Δ and ϕ according to χ , with χ varying
 249 in the range $[0^\circ, 80^\circ]$. The mean and standard deviation obtained from first order statistics
 250 performed on Δ and χ measured distributions (Table A1 (A)) are compared for different χ values
 251 (Fig.7).

252 On overall, increasing the value of χ leads to higher accuracy in the determination of the
 253 retardance and the optical axis orientation. Increase of χ has much more pronounced effect on
 254 the accuracy of retardance measurement, especially in the range $\chi \in [10^\circ, 40^\circ]$ (Fig.7 (A)). The
 255 parameter χ intervenes directly in the retardance formula (eq. 2), whereas it is not present in
 256 the optical axis orientation formula (eq. 5). In theory, χ should therefore not have any impact
 257 on calculation of ϕ . However, a small effect on optical axis orientation measurement accuracy
 258 (Fig.7 (B)) is observed for $\chi \in [10^\circ, 40^\circ]$. This is due to the increase of measured intensities
 259 separation with increasing χ , as discussed in section 2.

260 As expected, in the absence of sample, the measured retardance is close to $\Delta = 0^\circ$ (Fig.8
 261 (A,B)), since the ellipticity of the output polarization states ψ_1, ψ_2, ψ_3 and ψ_4 are all the same
 262 and $\eta \geq 0$. The accuracy saturates with the increase of χ values (Table A1 (B)). In the case of
 263 the HWP (Table A1 (C)), the increase of χ removes the ambiguity between $\Delta = 0^\circ$ and $\Delta = 180^\circ$,
 264 as discussed in section 2 (Fig.8 (C,D)).

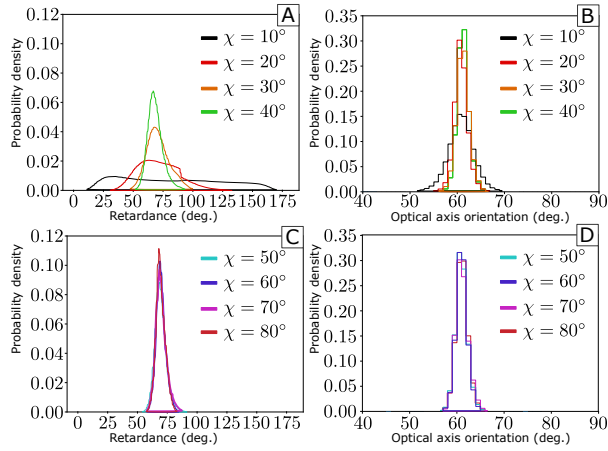


Fig. 7. Probability density drawn from pixel values of retardance (A,C) and optical axis orientation (B,D) for χ equal to $10^\circ, 20^\circ, 30^\circ$ and 40° (top charts) and χ equal to $50^\circ, 60^\circ, 70^\circ$ and 80° (bottom charts).

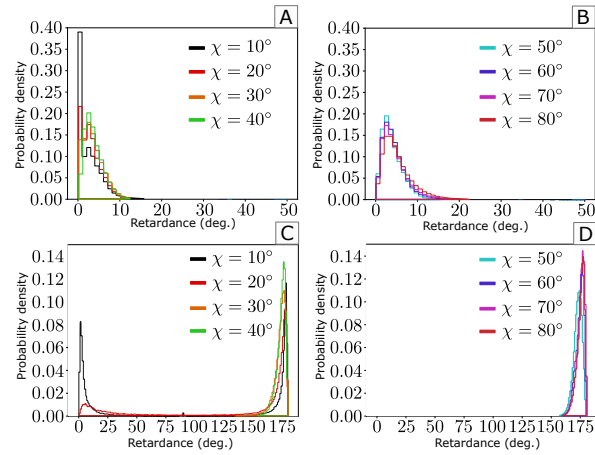


Fig. 8. Probability density drawn from pixel values of retardance in absence of sample ($\Delta = 0^\circ$) for χ equal to $10^\circ, 20^\circ, 30^\circ$ and 40° (A) and χ equal to $50^\circ, 60^\circ, 70^\circ$ and 80° (B). Probability density drawn from pixel values of retardance of a HWP for χ equal to $10^\circ, 20^\circ, 30^\circ$ and 40° (C) and χ equal to $50^\circ, 60^\circ, 70^\circ$ and 80° (D).

265 **5.2. Ambiguity on optical axis orientation**

266 If the angle ϕ is calculated from eq. 4, we already noted that the optical axis orientation suffers
 267 from indetermination in the case of a quarter-wave-plate (Thorlabs, WPMQ10M-633). This
 268 situation is illustrated in Fig.9 (A,C) for a quarter-wave plate oriented at -30° . However, thanks
 269 to our alternative expression of ϕ , i.e. (eq. 5), the indetermination on ϕ disappears (Fig.9 and
 270 Table A1 (D)). Both the map and the probability density obtained from eq. 4 show random ϕ
 271 values due to the indetermination. Those obtained from eq. 5 give almost uniform map with a
 narrow statistical distribution around the expected value.

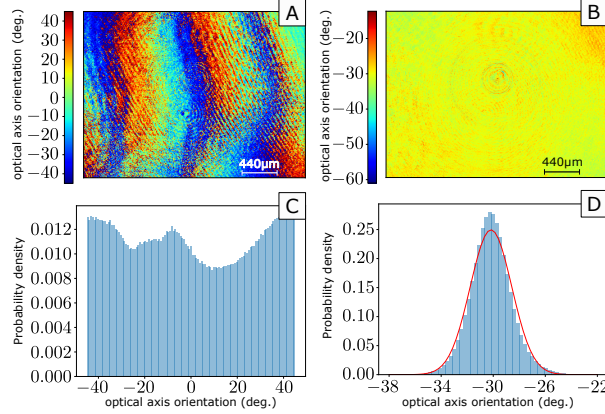


Fig. 9. Maps of the angle ϕ for a QWP oriented at $\phi = -30^\circ$ calculated using eq. 4 (A) or eq. 5 (B). Probability density (C, D) drawn from pixel values in images (A, B), respectively.

272
 273 Let us examine how is the indetermination removed thanks to our procedure. For an a priori
 274 unknown sample, the signs of the numerators of A and B allow us to determine the quadrant in
 275 which lies the angle that is solution of $\tan 2\phi = \left(\frac{I_1 - I_2}{I_4 - I_3} \right)$, with the angle 2ϕ being counted positive
 276 in anticlockwise direction from the horizontal axis. Two solutions exist, i.e. 2ϕ and $180^\circ + 2\phi$,
 277 the latter being located in the opposite quadrant with respect to the former. The numerators of A
 278 and B are given by, see eqs. 1:

$$\begin{aligned} I_1 - I_2 &= \sin(\chi) \sin(2\phi) \sin(\Delta), \\ I_4 - I_3 &= \sin(\chi) \cos(2\phi) \sin(\Delta). \end{aligned} \quad (8)$$

279 From now, we assume that Δ and χ both span from 0° to 180° , so that both $\sin \Delta$ and $\sin \chi$ are
 280 always positive. Therefore, the signs of the numerators of A and B are given by the signs of
 281 $\sin(2\phi)$ and $\cos(2\phi)$, respectively. A positive value of the product $(I_1 - I_2)(I_4 - I_3)$ indicates
 282 that the angle 2ϕ belongs either to the first or the third quadrant. Conversely, a negative value
 283 indicates that it belongs to either the second or the fourth quadrant.

284 We are interested to determine the angle ϕ , i.e. the optical axis orientation. Therefore, it is
 285 necessary to examine the signs of $I_1 - I_2$ and $I_4 - I_3$ independently in order to locate the angle
 286 in one of the four upper octants (Fig.10). Multiplication of the signs can lead to a flip between
 287 the fast and slow axis of the sample, e.g. the sign of $(I_1 - I_2)(I_4 - I_3)$ is positive for ϕ lying in
 288 the first and third octants, and negative in the second and fourth octants. For instance, without
 289 taking into account the signs of $I_1 - I_2$ and $I_4 - I_3$, an optical axis which forms an angle equal to
 290 $\phi = 100^\circ$ can be identified either as 10° (white octants in Fig.10) or 100° (light gray octants).
 291 Indeed, $2\phi = 200^\circ$ or $2\phi + 180^\circ = 20^\circ$ give the same value of the tangent. Now, taking into

292 account both the signs individually leads to angles equal either to $\phi = -80^\circ$ or 100° , which
 293 obviously define the same orientation (light gray octants).

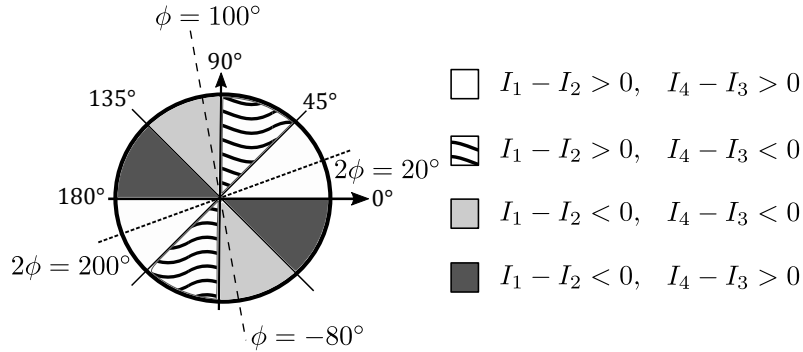


Fig. 10. Optical axis orientation is defined either by the angle ϕ or the angle $180^\circ + \phi$ (octants of the same color). The signs of both the numerators of A and B are examined in order to determine the correct optical axis orientation. An example is shown for $\phi = 100^\circ$.

293 In summary, the optical axis orientation can be determined without ambiguity when the signs
 294 of $I_1 - I_2$ and $I_4 - I_3$ are examined separately, as far as the retardance lies in the $[0^\circ, 180^\circ]$ range.
 295

296 5.3. Ambiguity on retardance

297 In principle, the signs of both $\sin \Delta$ and $\cos \Delta$ allow us to determine the quadrant in which Δ lies.
 298 For instance, Δ lies in the first quadrant when $\sin \Delta > 0$ and $\cos \Delta > 0$. In practice, however, we
 299 have only access to measurements of the intensities from which we deduce Δ and ϕ . A retardance
 300 higher than $\Delta = 180^\circ$ induce a flip of both the signs of $I_1 - I_2$ and $I_4 - I_3$ (eq. 8) which introduces
 301 an ambiguity in determination of ϕ if the range of Δ is not known a priori. This is why we had
 302 assumed $\Delta < 180^\circ$ in our previous discussion about optical axis orientation.

303 We illustrate hereafter the ambiguity in retardance determination by a simple experiment
 304 realized using commercial transparent cello-tape as sample. The tape consisted of an anisotropic
 305 polymer film with a smooth surface. We first observed cello-tape layers with a polarizing optical
 306 microscope and concluded that this material was birefringent [20], [21] (Fig.11 (A,B)). With
 307 its thickness of about $50 \mu\text{m}$, cello-tape is therefore a convenient sample for testing retardance
 308 measurement ambiguity. We then measured the retardance of one layer of cello-tape deposited
 309 on a glass plate and obtained $\Delta_{\text{cello},1} = 120^\circ$ (Fig.11 (C,D)). We can predict that superimposing n
 310 layers should induce a retardance equal to $\Delta_{\text{cello},n} = n\Delta_{\text{cello},1}$. Hence, two and three layers should
 311 give $\Delta = 240^\circ$ and $\Delta = 360^\circ$, respectively.

312 However, measurements of the retardance for two and three layers led to $\Delta = 120^\circ$ and $\Delta = 0^\circ$,
 313 respectively (Fig.11 B.). In the former case, the parameter η (determined from measured intensities,
 314 eq. 7) was found to be negative so that, according to eq. 3, $\Delta = 180^\circ - \arctan \sqrt{A^2 + B^2} =$
 315 $180^\circ - 60^\circ = 120^\circ$ and not 240° as expected.

316 Noting that $\tan \Delta = \sqrt{A^2 + B^2}$ is always positive, irrespective of the position of Δ in the
 317 quadrants, the formula of Δ must be adapted according to: $\Delta = 180^\circ + \arctan \sqrt{A^2 + B^2}$ when
 318 Δ lies in the third quadrant and $\Delta = 360^\circ - \arctan \sqrt{A^2 + B^2}$ when Δ lies in the fourth quadrant
 319 (Fig.12). In the first and second quadrants, the formula are the ones given by eq. 7. When
 320 $\Delta > 180^\circ$, the sign of $\cos \Delta$ allows us to discriminate whether the retardance lies in the third or
 321 the fourth quadrant. The sign can be extracted from the measured intensities if we figure out that,

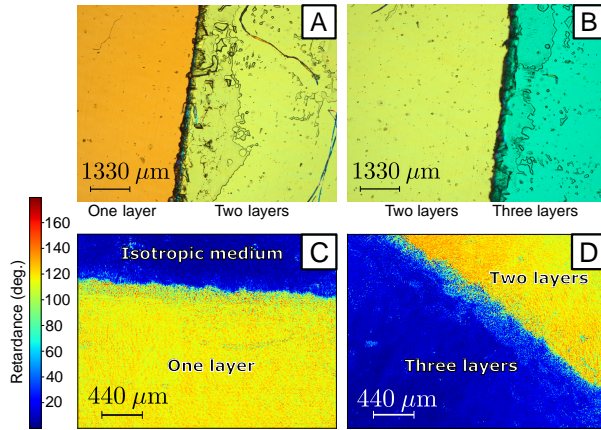


Fig. 11. (A,B) Polarizing optical microscope images of layers of cello-tape observed between crossed polarizers . A) One and two layers B) Two and three layers. Observation of colors indicates the birefringent nature of the sample. C) Map of the retardance of one cello-tape layer deposited on glass plate (isotropic medium). D) Map of the retardance imaging of two and three layers superimposed.

322 see eqs. 8

$$\cos \chi \cos \Delta = 1 - \frac{I_1 + I_2 + I_3 + I_4}{2}, \quad (9)$$

323 where χ is assumed to be in the range $]0^\circ, 90^\circ[$.

324 For two layers of cello-tape, we have $\cos \chi \cos \Delta < 0$. The retardance lies in the third quadrant
 325 and is given by $\Delta = 180^\circ + \arctan(\sqrt{A^2 + B^2})$, so $\Delta = 180^\circ + 60^\circ = 240^\circ$ as expected. In the
 case of the three layers, we have $\cos \chi \cos \Delta > 0$ and $\Delta = 360^\circ - 0^\circ = 360^\circ$, again as expected.

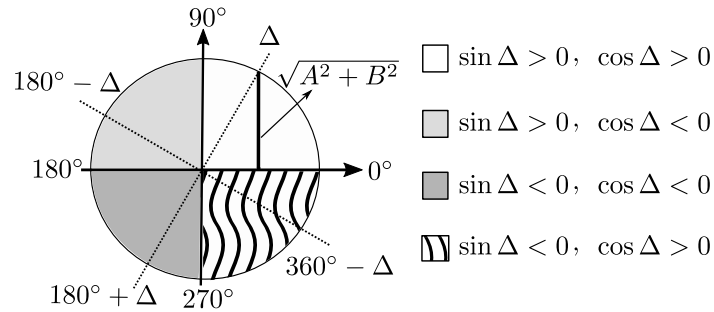


Fig. 12. Retardance, when calculated by $\Delta = \arctan \sqrt{A^2 + B^2}$, is always located in first quadrant. The formula used for calculates Δ must be adapted in the other cases: $180^\circ - \Delta$, $180^\circ + \Delta$ and $360^\circ - \Delta$ in the second, third and fourth quadrants.

326

327

328

329

330

331

In summary, the measurements of Δ and ϕ are unavoidably ambiguous for an a priori unknown sample because the optical axis orientation angle ϕ and retardance Δ appear together in the expressions of A and B in eqs. 2, i.e. the unknowns are coupled in the equations. The ambiguity can be removed, however, if we know a priori that Δ lies in the range $[0^\circ, 180^\circ]$ or in the range $[180^\circ, 360^\circ]$.

332 5.4. Application to geological samples

333 In order to illustrate the usefulness of our method in the context of natural samples, we analyzed
334 several geological thin sections with a thickness equal to $30\ \mu\text{m}$. The sample we selected for
335 illustration is composed of a mixture of quartz (SiO_2) and tourmaline
336 $[(\text{Ca}, \text{Na}, \text{K}, \text{Pb})(\text{Al}, \text{Fe}^{2+}, \text{Fe}^{3+}, \text{Li}, \text{Mg}, \text{Mn}^{2+}, \text{Ti})_3(\text{Al}, \text{Cr}, \text{Fe}^{3+}, \text{V})_6(\text{Si}_6\text{O}_{18})(\text{BO}_3)_3(\text{O}, \text{OH})_3(\text{F}, \text{O}, \text{OH})]$
337 crystals. Both silicate minerals belong to the trigonal crystal system and are uniaxial. However,
338 the signs of birefringence are different: quartz is positive and tourmaline is negative. Quartz
339 is translucent and uncoloured under polarized light with a weak birefringence of $\Delta n \approx 0.009$,
340 which enables recognition of quartz regions under optical microscope (Fig.13 A). Tourmaline
341 is translucent with colours varying within a single crystal depending on its chemistry under
342 polarized light (Fig.13 B). Its birefringence is stronger than quartz $\Delta n \in [0.015, 0.028]$. Under
343 optical microscope, it is easy to recognize tourmaline regions, as they appear colored (Fig. 13 A)

344 Due to the natural origin of the sample, it is not guaranteed that the optical axis lies in the plane
345 parallel to the sample surface, as assumed in the theoretical model (Fig.1). Moreover, quartz
346 regions are actually composed of grains whose axis orientations differ, as it can be observed
347 under polarized microscope (Fig.13 B). All these characteristics of composite natural samples
348 are retrieved by our measurements: namely, distinct retardance peaks corresponding to different
grains (Fig.13 (C,E)), with different optical axis orientations (Fig.13 (D,F)).

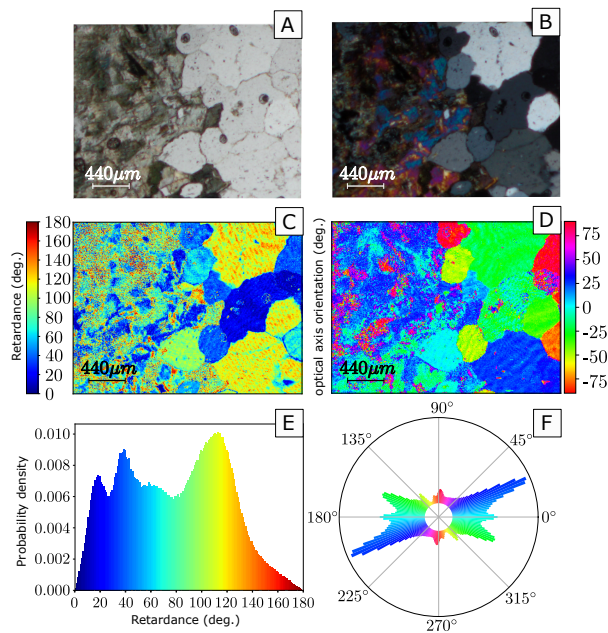


Fig. 13. Image of a $30\ \mu\text{m}$ thin section of a mixture of quartz and tourmaline observed (A) with an optical microscope. (B) with a polarizing optical microscope, between crossed polarizers. (C) Map of the retardance. (D) Map of the optical axis orientation. Probability density (E) drawn from pixel values in image (C). Optical axis orientation (F) drawn on circular histogram from pixel values in image (D).

349

350 6. Conclusion

351 The accuracy and ambiguities in birefringence measurements were examined in details and our
352 laboratory implementation was validated using reference samples. We proposed an alternative

353 formula for the calculation of the optical axis orientation, which improved the measurement
354 accuracy and avoided the indetermination in the case of a QWP. The effect of the swing value χ
355 on measurement accuracy was demonstrated and explained. We noted a clear improvement of
356 the accuracy, especially on retardance, with monotonically increase of χ parameter in the range
357 $[0^\circ, 90^\circ]$.

358 We examined the ambiguities in retardance determination thanks to a simple experiment based
359 on commercial cello-tape, which is a common example of anisotropic transparent film. Different
360 layers of cello-tape were superimposed, and the correct retardance value was identified according
361 to Δ was higher or lower than $\Delta = 180^\circ$.

362 We also applied our method for characterizing geological thin samples and showed it can help,
363 e.g. determining the orientation of grains in a composite sample.

364 In the future, we intend to apply this measurement procedure to the characterisation of
365 parchments, in the context of cultural heritage. Indeed, parchment, a non-opaque thin material
366 processed from animal skin, is known to exhibit birefringence and determination of spatial
367 distributions of local retardance and optical axis could bring valuable information on its state of
368 conservation [22].

369 **A. Appendix**

370 The mean and standard deviation values of the measured probability density of Δ and ϕ are
 371 displayed in the following tables, for three different retardation plates (UTP-137, QWP and HWP)
 and without sample.

Table A1. (A) Mean and standard deviation values of the measured probability density of the retardance and optical axis orientation for a retardation plate UTP-137 oriented at 60° . The angles calculated with the original and alternative formula are respectively noted ϕ and ϕ_{new} . (B) Mean and standard deviation values of the measured probability density of retardance ($\langle\Delta\rangle$, σ_Δ) and of the fitted (Gaussian) probability density function (μ , σ) in absence of sample. The angle ϕ is not calculated as it is undetermined in the absence of sample. (C) Mean and standard deviation values of the measured probability density of the retardance for an HWP. The angle is not calculated as it is undetermined in the case of a HWP. (D) Mean and standard deviation values of the measured probability density of the retardance and optical axis orientation for a QWP oriented at -30° .

| (A) UTP-137 oriented at 60° | | | | | | | | |
|--|--------|--------|--------|--------|--------|--------|--------|--------|
| $\chi [^\circ]$ | 10 | 20 | 30 | 40 | 50 | 60 | 70 | 80 |
| $\langle\Delta\rangle [^\circ]$ | 82.66 | 72.00 | 71.01 | 68.78 | 69.54 | 69.34 | 69.72 | 69.42 |
| $\sigma_\Delta [^\circ]$ | 42.21 | 18.69 | 10.39 | 6.96 | 5.32 | 4.46 | 4.90 | 4.06 |
| $\langle\phi\rangle_{\text{new}} [^\circ]$ | 60.77 | 60.83 | 61.17 | 61.24 | 61.00 | 61.01 | 61.12 | 61.12 |
| $\sigma_{\phi,\text{new}} [^\circ]$ | 3.43 | 1.55 | 1.50 | 1.30 | 1.30 | 1.27 | 1.38 | 1.31 |
| $\langle\phi\rangle [^\circ]$ | -17.78 | -17.80 | -25.87 | -28.35 | -28.42 | -28.39 | -28.58 | -28.56 |
| $\sigma_\phi [^\circ]$ | 25.59 | 22.68 | 13.33 | 6.74 | 4.47 | 2.58 | 1.40 | 2.07 |
| (B) Absence of sample | | | | | | | | |
| $\chi [^\circ]$ | 10 | 20 | 30 | 40 | 50 | 60 | 70 | 80 |
| $\langle\Delta\rangle [^\circ]$ | 15.75 | 5.98 | 4.47 | 3.90 | 4.03 | 4.35 | 4.68 | 5.42 |
| $\sigma_\Delta [^\circ]$ | 39.36 | 17.72 | 7.12 | 2.88 | 2.80 | 3.00 | 3.40 | 3.87 |
| $\mu [^\circ]$ | 9.85 | 4.97 | 4.07 | 3.90 | 4.03 | 4.35 | 4.68 | 5.42 |
| $\sigma [^\circ]$ | 32.61 | 16.31 | 6.91 | 2.88 | 2.80 | 3.00 | 3.40 | 3.87 |
| (C) HWP | | | | | | | | |
| $\chi [^\circ]$ | 10 | 20 | 30 | 40 | 50 | 60 | 70 | 80 |
| $\langle\Delta\rangle [^\circ]$ | 99.29 | 134.60 | 172.33 | 174.05 | 172.80 | 174.24 | 174.55 | 174.51 |
| $\sigma_\Delta [^\circ]$ | 82.47 | 64.27 | 8.90 | 4.22 | 3.82 | 3.54 | 3.41 | 3.45 |
| (D) QWP oriented at -30° | | | | | | | | |
| $\chi [^\circ]$ | 10 | 20 | 30 | 40 | 50 | 60 | | |
| $\langle\Delta\rangle [^\circ]$ | 90.23 | 87.70 | 87.94 | 89.48 | 89.23 | 89.77 | | |
| $\sigma_\Delta [^\circ]$ | 46.23 | 20.63 | 12.63 | 9.90 | 8.79 | 8.28 | | |
| $\langle\phi_{\text{new}}\rangle [^\circ]$ | -30.09 | -30.31 | -30.27 | -30.18 | -30.25 | -30.17 | | |
| $\sigma_{\phi,\text{new}} [^\circ]$ | 4.65 | 2.22 | 1.76 | 1.66 | 1.65 | 1.60 | | |

372

373 **Disclosures**

374 The authors declare no conflicts of interest.

375 **References**

- 376 1. M. C. T. Bahaa E. A. Saleh, *Polarization and Crystal Optics* (John Wiley and Sons, Ltd, 1991), chap. 6, pp. 193–237.
377 2. M. C. Pierce, J. Strasswimmer, H. Park, B. Cense, and J. F. de Boer, “Birefringence measurements in human skin
378 using polarization-sensitive optical coherence tomography,” *J. Biomed. Opt.* **9**, 287–291 (2004).
379 3. B. Cense, T. C. Chen, B. H. Park, M. C. Pierce, and J. F. de Boer, “In vivo depth-resolved birefringence measurements
380 of the human retinal nerve fiber layer by polarization-sensitive optical coherence tomography,” *Opt. Lett.* **27**,
381 1610–1612 (2002).
382 4. C. F. Chignell and D. K. Starkweather, “Optical studies of drug-protein complexes,” *Mol. Pharmacol.* **7**, 229–237
383 (1971).
384 5. A. M. Hofmeister, R. B. Schaal, K. R. Campbell, S. L. Berry, and T. J. Fagan, “Prevalence and origin of birefringence
385 in 48 garnets from the pyrope-almandine-grossularite-spessartine quaternary,” *Am. Mineral.* **83**, 1293–1301 (1998).
386 6. H. Mueller, “The theory of photoelasticity,” *J. Am. Ceram. Soc.* **21**, 27–33 (1938).
387 7. S.-T. Wu, U. Efron, and L. D. Hess, “Birefringence measurements of liquid crystals,” *Appl. Opt.* **23**, 3911–3915
388 (1984).
389 8. I. G. Wood and A. M. Glazer, “Ferroelastic phase transition in bivo4 i. birefringence measurements using the
390 rotating-analyser method,” *J. Appl. Crystallogr.* **13**, 217–223 (1980).
391 9. R. C. Miller and A. Savage, “Temperature dependence of the optical properties of ferroelectric LiNbO₃ and LiTaO₃,”
392 *Appl. Phys. Lett.* **9**, 169–171 (1966).
393 10. R. Evans, “Crystals and the polarising microscope by nh hartshorne and a. stuart,” *Acta Crystallogr.* **13**, 853–853
394 (1960).
395 11. J. C. Tolédano, “Phenomenological model for the structural transition in benzil,” *Phys. Rev. B* **20**, 1147–1156 (1979).
396 12. R. T. Harley and R. M. Macfarlane, “A determination of the critical exponent beta in TbVO₄ and DyVO₄ using linear
397 birefringence,” *J. Phys. C: Solid State Phys.* **8**, L451–L455 (1975).
398 13. W. T. Welford, “Principles of optics (5th Edition). M. Born, E. Wolf Pergamon Press, Oxford, 1975, pp xxviii + 808,
399 £9.50,” *Opt. Laser Technol.* **7**, 696 (1975).
400 14. D. J. Benard and W. C. Walker, “Modulated polarization measurement of structural phase transitions in kmnf₃,” *Rev.*
401 *Sci. Instruments* **47**, 122–127 (1976).
402 15. M. Shribak and R. Oldenbourg, “Techniques for fast and sensitive measurements of two-dimensional birefringence
403 distributions,” *Appl. Opt.* **42**, 3009–3017 (2003).
404 16. M. Shribak, “Complete polarization state generator with one variable retarder and its application for fast and sensitive
405 measuring of two-dimensional birefringence distribution,” *J. Opt. Soc. Am. A* **28**, 410–419 (2011).
406 17. S. B. Mehta, M. Shribak, and R. Oldenbourg, “Polarized light imaging of birefringence and diattenuation at high
407 resolution and high sensitivity,” *J. Opt.* **15**, 094007 (2013).
408 18. “OpenPolScope,” <https://openpolscope.org/>. Accessed: 2022-02-30.
409 19. R. C. Jones, “A new calculus for the treatment of optical systems. v. a more general formulation, and description of
410 another calculus,” *J. Opt. Soc. Am.* **37**, 107–110 (1947).
411 20. A. Beléndez, E. Fernández, J. Francés, and C. Neipp, “Birefringence of cellotape: Jones representation and
412 experimental analysis,” *Eur. J. Phys.* **31**, 551–561 (2010).
413 21. M. A. Blanco, M. Yuste, and C. Carreras, “Undergraduate experiment designed to show the proportionality between
414 the phase difference and the thickness of a uniaxial crystal,” *Am. J. Phys.* **65**, 784–787 (1997).
415 22. V. Vilde, M. Fourneau, C. Charles, D. V. Vlaender, J. Bouhy, Y. Poumay, and O. Deparis, “Use of polarised light
416 microscopy to improve conservation of parchment,” *Stud. Conserv.* **64**, 284–297 (2019).