Experimental quantum polarimetry using heralded single photons

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Abstract

We perform experimental quantum polarimetry using a heralded single photon to analyze the optical activity of linearly polarized light traversing a chiral medium. Three kinds of estimators are considered to estimate the concentrations of sucrose solutions from measuring the rotation angle of the linear polarization of the output photons. Through repetition of independent and identical measurements performed for each individual scheme and different concentration sucrose solutions, we compare the estimation uncertainty among the three schemes. The results are also compared to classical benchmarks for which a coherent state of light is taken into account. The quantum enhancement in the estimation uncertainty is evaluated and the impact of experimental and technical imperfections is discussed. In this work, we lay out a route for future applications relying on quantum polarimetry.

Keywords: optical activity, chirality, single photon, heralded single photon, polarimetry, quantum polarimetry, estimation

(Some figures may appear in colour only in the online journal)

1. Introduction

Optical activity causes rotation of the linear polarization of light when it traverses a chiral material, i.e. a material made from entities that lack mirror symmetry. The measurement of such optical activity has often been used in pharmacology, where it is important to study the chirality of drug molecules, which determines their toxicity and efficacy, originating from the relative handedness of the enantiomers [1, 2]. Chirality has also been explored in various other fields, including chemistry, life science, physics, and material science, where polarimetric schemes provide rich information about molecular or nanophotonic chiral structures [3–8].

A variety of detection schemes have been developed for measuring the optical activity, i.e. analyzing the rotation of the linear polarization of light upon propagation through a chiral medium [9–11]. This rotation of the polarization has its origin in the different propagation constants for the two counter-rotating circularly polarized plane waves, which are eigenmodes of the chiral medium and into which the linear polarization can be decomposed. The estimated direction of linear polarization will fluctuate due to the discrete nature of light even when the chirality is fixed and experimental noise is removed. These fluctuations determine the reliability of the measurement, often called the precision, which reflects the data quality. In general, the precision increases with the intensity $N$ (or the average photon number) of the light used. However, there exist situations in which the intensity of the incident light impinging on an analyte is required to be limited, when sample damage or any unwanted side effects occur in the high intensity regime, e.g. when the non-linear dependence is significant. Also, light itself can trigger unwanted chemical reactions in the molecular material to be detected [12–15]. The presence of such constraints has inspired the development of quantum metrological approaches over the last few decades. The aim is to improve the data quality while keeping the incident power in the low intensity regime [16, 17], i.e. improving the precision for a fixed intensity.
Probing analytes with quantum states of light promises detection with much reduced noise, below what is possible with classical states of light, thus allowing better performance in optical measurements [18-20]. In particular, the phase-sensitive N00N states have been adopted to measure the chirality. For example, the Faraday rotation in a rubidium vapor cell has been measured with polarization N00N states, showing that the standard quantum limit can be beaten [21]. The dispersion of optical rotation has also been analyzed by injecting polarization-entangled states with different and tunable wavelengths [22]. Chemical processes that change the chirality of molecules have been monitored in real-time with polarization N00N states with a protocol in the context of quantum phase estimation [23, 24]. In addition to schemes measuring the chirality, N00N states have been employed to achieve a quantum enhancement in other sensing scenarios, such as in the measurement of the refractive index of blood proteins [25], quantum lithography [26], and super-sensitive microscopy [27, 28]. The superior optical detection shown with the N00N states originates from the quantum entanglement of photons, leading to a higher signal-to-noise ratio ($\sqrt{N}$ times larger) when compared to measurements that rely on classical sources such as coherent states of light. The quantum advantage enabled by N00N states is known to be vulnerable to photon loss or decoherence [29].

In the context of polarimetry, the chirality of a molecular solution is measured by measuring the transmittance $T$ of a linearly polarized incident light passing through an analyte resolved in its co-or cross-polarized components. For definiteness we speak in the following of horizontal and vertical polarization. The presence of the chiral medium rotates the polarization weakly, which can be probed from the change of the intensity of the incidence polarization or from the difference between horizontally and vertically polarized light. From these measured intensities we can determine the concentration of molecules if the chirality itself is known. It is known that the photon number state $|N\rangle$ is the optimal state that minimizes the noise in the transmission measurement [34, 35]. This is because the photon number state has no uncertainty in its intensity, offering the most precise detection when monitoring a change of the intensity [36, 37]. Unlike N00N states, the photon number state always offers a quantum gain in noise reduction at any loss level $\gamma = 1 - \eta$, exhibiting a detection fluctuation $\Delta I = \sqrt{I(1 - \eta T)/N\eta}$ compared to $\Delta I = \sqrt{I/N\eta}$ that is obtainable when using coherent states with an average photon number of $N$. Therefore, the photon number state is not just optimal in reducing the noise or uncertainty, but also useful against the photon loss in realistic transmission measurements. As the generation of Fock states with an arbitrary photon number $N$ is not yet available with current technology, one can use instead $N$ single photons relying on the equivalence to the case using an $N$ photon number state in transmission measurement. Several studies have been carried out, such as absorption spectroscopy to analyze the organic dye molecule dibenzanthanthrene [38] or haemoglobin [39], and surface plasmon resonance sensing to measure the change of the refractive index of an analyte under investigation with such Fock states [40].

In this work, we study theoretically and experimentally how much quantum enhancement can be obtained in measuring the optical activity of chiral molecular materials with single photons as an input state. We use a heralding scheme to generate the single-photon state that is linearly polarized and that illuminates sucrose solutions with different concentrations. To estimate the concentration of the sucrose solutions, we consider three typical polarimetric schemes: measurement of (i) the intensity of the horizontally polarized outgoing photons, (ii) the intensity difference between horizontally and vertically polarized outgoing photons, and (iii) the intensity difference-to-sum ratio (DSR) between the two linearly polarized outgoing photons. We analyse and compare the performances of these three schemes that use single-photon states to each other, but also to the optimal precision achievable with a classical input state. We find that type (iii) leads to the minimal estimation uncertainty compared to the other two types. A quantum enhancement, i.e. an improvement of the estimation precision when compared to measurement with a classical state of light is experimentally observed for both type (i) and type (ii) measurements. In general, our study proves that the quantum enhancement is always obtained at any $\eta$ and $T$. The effect of experimental imperfections is also discussed, e.g. the extinction ratio of the optical components used in the experiment.

2. Experimental scheme

The experimental scheme used in this work is shown in figure 1(a). A continuous wave diode laser with a wavelength of 405 nm, filtered out at a later stage by a band pass filter and a long pass filter, pumps a non-linear crystal (a periodically poled potassium-titanyl-phosphate) to generate paired photons at the emission wavelengths of 809.6 nm and 810.4 nm with a full width at half maximum of 9.4 nm and 7.2 nm, respectively (see figure 1(b)). The two orthogonally polarized photons are spatially separated by a polarizing beam-splitter. One of the two photons, the vertically polarized idler photon, is directly sent to an avalanche photodiode single-photon detector (APD; SPCM-AQR-15, PerkinElmer). Quantum correlation between the photon pairs enables the click events from the idler detector APD$_I$ to herald the horizontally polarized signal photon that is fed into the polarimetry set-up under investigation. Rotation of the linear polarization occurs when the signal photon passes through a sucrose solution. The magnitude of this rotation depends on the concentration of the sucrose solution, eventually determining the extent to which the polarization of the outgoing photons rotates. To make the scheme operate with heralded single photons in detection, we adopt a coincident measurement between APD$_I$ and APD$_{H,V}$, i.e. the results of detection at APD$_{H,V}$ are recorded only when the single photons are detected at APD$_I$. Throughout this paper, let $N_{H,V}$ be the coincidence counts between APD$_I$ and APD$_{H,V}$.

The heralded single-photon scheme is valid in the low gain parameter regime, where the state generated via spontaneous parametric down-conversion (SPDC) can be written
The wavelength dependence in the specific rotation can be
where $[\alpha(\lambda)]$ is the specific rotation of the sample material.
The wavelength dependence in the specific rotation can be
modelled by Drude’s expression, written as
\[
[\alpha(\lambda)] = \sum_j \frac{A_j}{\lambda^2 - \lambda_j^2},
\]
where the summation is over multiple excitation transitions
with $\lambda_j$ and $\lambda$ being the resonance wavelength and the rotation
amplitude for the $j$th transition, respectively. For the case of
sucrose, the dispersion of the optical activity can be characterized
by a single transition (i.e., $j = 1$), for which $A_1 = 2.1648 \times 10^5$
$\text{deg nm}^2 \text{dm}^{-1} \text{g}^{-1} \text{ml}$ and $\lambda_1 = 146 \text{nm}$ [22, 41]. We thus
aim to estimate the concentration $C$ from the measurement of
the angle $\alpha(\lambda)$ for $\lambda = 0.1 \text{ dm}$ (the length of the cuvette used
in our experiment) and $[\alpha(\lambda)]$ given above.

In this work, we implement three kinds of estimators, yielding
the associated $f$ values, which are often used in classical polarimetry [9–11]: (i) the number $N_{\text{H}}$ of the horizontally
polarized photons transmitted through the sample, but
normalized by $\nu$; (ii) the difference $(N_{\text{H}} - N_{\text{V}})$ between
the horizontally and the vertically polarized photons transmitted
through the sample, normalized by $\nu$; and (iii) the difference
$(N_{\text{H}} - N_{\text{V}})$ normalized by the sum $(N_{\text{H}} + N_{\text{V}})$, i.e. the DSR,
which has recently been employed in imaging of non-uniform refractive profiles [42]. The expectation values of the above
tree estimators for $\nu$ single photons of incidence can be written,
respectively, as
\[
(i) \quad \langle f_{\text{s}} \rangle = \frac{\langle N_{\text{H}} \rangle}{\nu} = \eta_{\text{H}} T_{\theta_{\text{out}}},
\]
\[
(ii) \quad \langle f_{\text{diff}} \rangle = \frac{\langle N_{\text{H}} - N_{\text{V}} \rangle}{\nu} = \eta_{\text{H}} T_{\theta_{\text{out}}} - \eta_{\text{V}} R_{\theta_{\text{out}}},
\]
\[
(iii) \quad \langle f_{\text{DSR}} \rangle = \frac{\langle N_{\text{H}} - N_{\text{V}} \rangle}{N_{\text{H}} + N_{\text{V}}} \approx \frac{\eta_{\text{H}} T_{\theta_{\text{out}}} - \eta_{\text{V}} R_{\theta_{\text{out}}}}{\eta_{\text{H}} T_{\theta_{\text{out}}} + \eta_{\text{V}} R_{\theta_{\text{out}}}},
\]
where $\langle . \rangle$ denotes an average with respect to the output state
being measured and $\eta_{\text{H,V}}$ denote the efficiencies of transmission
from the PPKTP to the detectors $\text{APD}_{\text{H,V}}$ including the
detection efficiencies. Here, $T_{\theta_{\text{out}}} = \cos^2 \theta_{\text{out}}$ and $R_{\theta_{\text{out}}} = \sin^2 \theta_{\text{out}}$
denote the transmittance and the reflectance for the
outgoing single photon with a polarization of $\theta_{\text{out}} = \theta_{\text{in}} + \alpha$
to be transmitted through and reflected from the PBS$_2$ in figure
1, respectively. In particular, equation (5) asymptotically

![Figure 1. (a) Experimental scheme. A continuous-wave (CW) diode laser pumps a non-linear crystal (periodically poled potassium-titanyl-phosphate (PPKTP)) to produce photon pairs. The paired photons are orthogonally polarized and separated by the first polarizing beam-splitter (PBS$_1$). The pump beam is removed from the optical system using a band pass filter (BPF) and a long pass filter (LPF). The detection of a vertically (V) polarized photon at an avalanche photodiode in the idler channel (APD$_i$) heralds the generation of horizontally (H) polarized photons in the signal channel. The incident H-photons are rotated at a defined angle by a half-wave plate (HWP), and sent to a cuvette containing the sucrose solution under investigation. The polarization of the outgoing photons is analyzed via detection at APD$_s$ placed in the output ports of the PBS$_2$. (b) The spectra of the photons generated by the SPDC process in the PPKTP. The central wavelengths of the signal and the idler photons are located at 809.6 nm and 810.4 nm, respectively.](image-url)
holds for a large sample size \( \nu \), according to Jensen’s inequality [43], which is the case in our experiment. In an ideal case where \( \eta_H = \eta_V, N_H + N_V = \nu \), so that equation (5) becomes equation (4). The values \( \eta_{4HV} \approx 0.25 \) are explicitly measured by the coincidence count \( N_{4HV} \) with scanning over \( \theta_{\text{in}} \) when pure water is in the cuvette, i.e. \( \alpha = 0 \) (see the black curves and circles in figure 2), which exploits the correlated features of the SPDC photons as used in the Klyshko method [44–46].

In what follows, equations (3)–(5) are used to estimate the angle of rotation \( \alpha \) from the measurement of the output polarization \( \theta_{\text{out}} = \theta_{\text{in}} + \alpha \) for a given input polarization \( \theta_{\text{in}} \). The estimation of \( \alpha \) subsequently leads to the estimation of the concentration \( C \) of sucrose solutions through equation (1). The impact of a finite concentration of the sucrose solution can be seen as a shift of the curve (for example, see the red curves and triangles in figure 2 for the measurement of the sucrose solution with a concentration of 0.5 g ml\(^{-1}\)). Assuming the values of [\( \alpha \)], \( \eta_{4HV} \), and \( \theta_{\text{in}} \) to be accurately known beforehand, one can determine the estimation uncertainty \( \Delta C \) directly by the estimation uncertainty of the output polarization \( \Delta \theta_{\text{out}} \) extracted from the measurement of the outcome \( f \).

3. Results and discussion

\( \Delta \theta_{\text{out}} \) is experimentally obtained by the standard deviation of the estimated values \( \theta_{\text{out}} \) from \( \mu \) measurements with a sample size of \( \nu \), i.e. \( \Delta \theta_{\text{out}} = \sqrt{\frac{\sum f_{\text{out}}^2}{\mu} - (\sum f_{\text{out}}/\mu)^2} \). The experimentally measured value \( \Delta \theta_{\text{out}}^{\exp} \) is compared to the theoretical prediction \( \Delta \theta_{\text{out}} \), which can be calculated by considering the linear error propagation from the variance of the outcomes \( \Delta f = \sqrt{\langle (\Delta f)^2 \rangle} \) [47], written as

\[
\Delta \theta_{\text{out}} = \frac{\Delta f}{\theta_{\text{out}}}.
\]

The variance \( \langle (\Delta f)^2 \rangle \) can be calculated as

\[
\langle (\Delta f)^2 \rangle = \sum_{j,k \in \{H,V\}} \frac{\partial f_j}{\partial N_j} \frac{\partial f_k}{\partial N_k} \text{Cov}(N_j,N_k),
\]

where \( \text{Cov}(N_j,N_k) \) is the covariance between the photon counts \( N_j \) and \( N_k \). Equation (7) is an exact function for \( f_{\text{single}} \) and \( f_{\text{diff}} \), but is an approximate function for \( f_{\text{DSR}} \) since equation (5) is a non-linear differential function of \( N_{4HV} \) [48]. In our experiment, \( \nu \) heralded single photons are used, and the theoretical expression of \( \Delta \theta_{\text{out}} \) of equation (6) can be written, respectively, as

\[
\Delta \theta_{\text{out}}^{\text{single(q)}} = \frac{1}{\sqrt{\nu}} \sqrt{\frac{1 - \eta_H T_{\text{out}}}{4 \eta_H (1 - T_{\text{out}})}},
\]

\[
\Delta \theta_{\text{out}}^{\text{diff(q)}} = \frac{1}{\sqrt{\nu}} \sqrt{\frac{(\eta_H T_{\text{out}} + \eta_V R_{\text{out}}) - (\eta_H T_{\text{out}} - \eta_V R_{\text{out}})^2}{4(\eta_H + \eta_V)^2 T_{\text{out}} R_{\text{out}}}},
\]

\[
\Delta \theta_{\text{out}}^{\text{single(q)}} > \Delta \theta_{\text{out}}^{\text{diff(q)}},
\]

\[
\Delta \theta_{\text{out}}^{\text{DSR(q)}} = \frac{1}{\sqrt{\nu}} \sqrt{\frac{\eta_H T_{\text{out}} + \eta_V R_{\text{out}}}{4 \eta_H \eta_V}}.
\]

The uncertainties \( \Delta \theta_{\text{out}}^{\text{single(q)}} \) and \( \Delta \theta_{\text{out}}^{\text{diff(q)}} \) diverge as \( T_{\text{out}} \to 1 \) (unless \( \eta_H = 1 \)) and \( T_{\text{out}} \to 0 \) or 1 (unless \( \eta_H = \eta_V = 1 \)), but decrease as \( T_{\text{out}} \to 0 \) and \( T_{\text{out}} \to 1 \). The uncertainties \( \Delta \theta_{\text{out}}^{\text{DSR(q)}} \) is maximum at \( T_{\text{out}} = 1 \) (unless \( \eta_H = \eta_V = 1 \)). The minima and maxima are reversed when \( \eta_H > \eta_V \). The uncertainty \( \Delta \theta_{\text{out}}^{\text{DSR(q)}} \) takes a constant value \( 1/(\sqrt{\nu} \sqrt{4\eta_H}) \) regardless of \( \theta_{\text{in}} \). Also note that

\[
\text{min} \Delta \theta_{\text{out}}^{\text{DSR(q)}} = \text{min} \Delta \theta_{\text{out}}^{\text{single(q)}} < \text{min} \Delta \theta_{\text{out}}^{\text{diff(q)}}
\]

when \( \eta_H > \eta_V \), whereas

\[
\text{min} \Delta \theta_{\text{out}}^{\text{DSR(q)}} < \text{min} \Delta \theta_{\text{out}}^{\text{diff(q)}} < \text{min} \Delta \theta_{\text{out}}^{\text{single(q)}}
\]

Figure 2. Open symbols show the measured \( f_{\text{DSR}} \) of equation (5) while varying the input polarization angle \( \theta_{\text{in}} \) for deionized water (black circles) and for sucrose solution with \( C = 0.5 \) g ml\(^{-1}\) (red triangles). The inset shows the measured values in a full range of \( \theta_{\text{in}} \) from –100° to 100° in steps of 10°. The horizontal displacement between the two sets of data exhibits the optical activity induced by the chiral material, enabling estimation of the rotation angle \( \alpha \). Solid curves are the fitted values using equation (10). The error bars are the standard deviation in the histogram over \( \mu \) times of repetition at each value of \( \theta_{\text{in}} \).
\[ \Delta \theta_{\text{out}}^{\text{diff}}(c) = \frac{1}{\sqrt{V}} \sqrt{\frac{\eta_1 T_{\text{out}} + \eta_2 R_{\text{out}}}{4(\eta_1 + \eta_2)^2 T_{\text{out}} R_{\text{out}}}}, \]  
\[ \Delta \theta_{\text{out}}^{\text{DSR}}(c) = \frac{1}{\sqrt{V}} \sqrt{\frac{\eta_1 T_{\text{out}} + \eta_2 R_{\text{out}}}{4\eta_1 \eta_2}}, \]  
respectively. The relative quantum enhancement \( R \) can be quantified by the ratio of \( \Delta \theta_{\text{out}}^{(c)} \) to \( \Delta \theta_{\text{out}}^{(q)} \) written as

\[ R = \frac{\Delta \theta_{\text{out}}^{(c)}}{\Delta \theta_{\text{out}}^{(q)}}. \]

Therefore, a value of \( R \) greater than unity exhibits a quantum enhancement. For the three schemes considered above, one can show that \( R^{\text{single}} = (1 - \eta_1 T_{\text{out}})^{-1/2} \), \( R^{\text{diff}} = [1 - (\eta_1 T_{\text{out}} - \eta_2 R_{\text{out}})/(\eta_1 T_{\text{out}} + \eta_2 R_{\text{out}})]^{-1/2} \), and \( R^{\text{DSR}} = 1 \). Interestingly, \( R \) is always equal to or greater than unity for all schemes considered in our experiment. These behaviors of the ratio \( R \) are presented in figure 3(b), given by the experimentally measured noise \( \Delta \theta_{\text{out}} \) and the theoretical value \( \Delta \theta_{\text{out}} \) that would be obtainable via the optimal classical polarimetry.

The estimators of equations (3)–(5) are unbiased under ideal conditions. One may wonder if the above estimation uncertainties evaluated using equation (6) reach the lower bound given by the Cramér–Rao (CR) inequality being satisfied for any unbiased estimator [49, 50]. The CR inequality can be written as \( \Delta \theta_{\text{out}} \geq 1/\sqrt{V} F(\theta_{\text{out}}) \) with \( F(\theta_{\text{out}}) \) being the Fisher information (FI) for a parameter \( \theta_{\text{out}} \). It depends on the measurement setting that is performed [51]. From the expressions of the FI (see appendix A) for both the classical and quantum schemes considered in this work, one can clearly see that the uncertainties evaluated using equation (6) are the same as the inverse of the squared FI regardless of \( T_{\text{out}}, R_{\text{out}}, \) and \( \eta_1, \eta_2 \) when estimating the parameter with the single-mode measurement scheme (i.e. \( f_{\text{single}} \)). However, the uncertainties of equation (6) with the two-mode measurement schemes (i.e. \( f_{\text{diff}} \) and \( f_{\text{DSR}} \)) are equal to or larger than the inverse of the squared FI. This implies that a better estimator than using \( f_{\text{diff}} \) and \( f_{\text{DSR}} \) exists when the photon-number-counting measurement is performed at the two modes.

When the measurement setting is optimized, the CR inequality can be further developed to be that called the quantum CR inequality, where the maximized FI is called the quantum FI (QFI) [47, 52]. For the output state decomposed into \( \hat{\rho}_{\text{out}} = \sum_k |\psi_k\rangle \langle \psi_k| \) with \( \langle \psi_k | \psi_k \rangle = \delta_{jk} \), it is known that the FI is the same as the QFI if \( \Delta \theta_{\text{out}} | \psi_k \rangle = 0 \) \( \forall \psi \). Interestingly, the latter is the case for both the classical and quantum schemes considered in this work. This indicates that the photon-number-counting measurement is the optimal measurement scheme for both the classical and quantum approaches. This may not be true for other kinds of probe states, i.e. an individual optimal measurement scheme needs to be found and performed for maximization of quantum enhancement when other types of quantum states are illuminated into the sample.

Actual estimation of the concentration of sucrose solutions can be made via the relation \( \theta_{\text{out}} = (\theta_{\text{in}}) + \alpha \) and equation (1).
Figure 4. Experimentally estimated concentrations of sucrose solutions with six different concentrations are obtained by using the estimator of equation (5). We exclude the shaded regions as in figure 3, where the estimated concentrations are misleading due to a non-ideal extinction ratio (see appendix B for verification via a Monte-Carlo simulation). The error bar at individual points represents the standard deviation of the histogram obtained over μ times measurements.

Figure 4 shows the results of the scheme using the DSR estimator of equation (5) for sucrose solutions with different concentrations. The average estimated values C of the concentration are shown together with the estimation uncertainties ΔC. The latter reflects the type A uncertainties caused by probabilistic photon number counting. Further evaluation regarding uncertainties of other factors can be made, but we leave rigorous investigation of those minor contributions as a future work while providing the type B uncertainty budget in appendix B (see table B1). The shaded areas in figure 4 represent fallible regions which would lead to erroneous estimation, so we exclude those regions from the estimation.

It is worth discussing two experimental imperfections that led to the exclusion of the fallible regions in figures 3 and 4. First, the polarization control by experimental components may not be ideal, e.g. an incident H-photon to the polarizing beam splitter can exit through the V-photon output port. Its rate can be quantified by the polarization extinction (PE) ratio rPE, which is measured to be about 1000/1 in our experiment. This means that even when \( \theta_{out} = 0 \), a few photons are found in APD\( \nu \) although no photon is supposed to be detected there in the ideal case. Such an imperfection becomes more significant as \( T_{\theta_{in}} \rightarrow 0 \) or 1, around which the estimation of the concentration C is likely to be more inaccurate. The effect of the PE is investigated via Monte-Carlo simulation (see the details in appendix C). We also identify that the imperfect functioning of experimental components can contribute to type A uncertainty. Second, particularly when the estimation is made with equation (3) at each incidence angle \( \theta_{in} \), the experimentally measured efficiency \( \eta_{H} \) is inserted into equation (3). This works reasonably well except for the cases when more than \( \eta_{H} \nu \) photons are detected in the measurement counting the photon number \( N_{H} \). For example, consider the extreme yet possible case that all \( \nu \) photons are detected in \( N_{H} \) even when \( \eta_{H} < 1 \). Note that photon loss occurs probabilistically with a probability \( \eta_{H} \). In this case, \( N_{H}/\nu \) can be greater than unity and thus \( T_{\theta_{in}} > 1 \), leading to unphysical estimation. This arises because the value \( N_{H}/\nu \) is divided by \( \eta_{H} \), implying that the probabilistic nature associated with the efficiency \( \eta_{H} \) is regarded as a deterministic process that supposes only a definite number \( \eta_{H} \nu \) of photons passes through an analyte, which is of course wrong and misses the probabilistic nature. Such an extreme case often takes place when \( T_{\theta_{in}} \rightarrow 1 \) and similarly when \( T_{\theta_{in}} \rightarrow 0 \). The same concern also applies to the estimation with equation (4). Therefore, we exclude the range of \( \theta_{out} \) such that \( \eta_{H} \nu \langle \theta_{out} \rangle - \langle N_{H} \nu \theta_{out} \rangle < 3 \Delta N_{H} \nu \langle \theta_{out} \rangle \) for all the estimation schemes considered in this work in order to avoid inaccurate or unphysical estimation. The corresponding regimes are represented by shaded areas in figures 3 and 4.

The first issue could be alleviated by using optical components with as high a PE ratio rPE as possible or at least a much better ratio on a logarithmic scale. The second issue, on the other hand, requires a more sophisticated modification to the estimation schemes. For example, one could perform simultaneous estimation of the concentration C and the efficiencies \( \eta_{H} \nu \) from the measurement with scanning through the incidence angles. This can typically be done by fitting equations (3)–(5) to the measured data, which we leave for future study.

4. Conclusion

We have experimentally investigated quantum polarimetric schemes using single-photon inputs to analyze the concentration of sucrose solutions. The horizontally and vertically polarized outgoing photons have been counted for each polarization of incidence and each concentration of sucrose solution in the experiment, leading to the determination of the optical activity. It has been shown that the minimal estimation uncertainty is generally achieved when the concentration is analyzed with a normalized difference between the horizontal and vertical photons, i.e. equation (5). We have discussed the quantum gain in the detection of the optical activity with respect to the three typical polarimetric schemes and the effects of the experimental and technical imperfections are identified.

When probing the optical activity with more intense light than a single-photon state and when it is, in addition, allowed and preferred for practical purposes, one may exploit bright squeezed states that can carry high optical powers [53]. In this case, the unit quantum enhancement per photon is still smaller when compared to single photons, but the overall quantum enhancement would be much higher due to the use of increased optical power (i.e. much larger N). Using bright squeezed states would provide practical quantum polarimetry in the high intensity regime, which we leave for future study. Quantum theory for the characterization of an arbitrary polarization in terms of Stokes parameters has recently been discussed and would be useful for further development of various quantum polarimetric schemes [54–56].
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Appendix

Appendix A. Fisher information

Here we provide the calculation of the FI when the photon-number-counting measurement is performed at single-mode H or V, or at both modes, respectively. The FI can be written in terms of the probabilities of the measurement outcomes as

$$F_{N_{H},N_{V}}(\theta_{\text{out}}) = \sum_{N_{H},N_{V}} \frac{1}{p(N_{H},N_{V}|\theta_{\text{out}})} \left( \frac{\partial p(N_{H},N_{V}|\theta_{\text{out}})}{\partial \theta_{\text{out}}} \right)^2,$$

(A1)

when $\theta_{\text{out}}$ is estimated from measurement at the two modes. Similarly, when measurement is performed at either mode H or mode V, the FI can be written as

$$F_{N_{s}}(\theta_{\text{out}}) = \sum_{N_{s}} \frac{1}{p(N_{s}|\theta_{\text{out}})} \left( \frac{\partial p(N_{s}|\theta_{\text{out}})}{\partial \theta_{\text{out}}} \right)^2,$$

(A2)

where $s \in \{H, V\}$. It can be easily shown that the associated probabilities are written as

$$p^{(s)}(N_{H},N_{V}) = \binom{N}{N_{H}} (1-\eta_{H} T_{\theta_{\text{out}}})^{N_{H}} (\eta_{V} R_{\theta_{\text{out}}})^{N_{V}} \times (1-\eta_{H} T_{\theta_{\text{out}}}-\eta_{V} R_{\theta_{\text{out}}})^{N-N_{H}-N_{V}},$$

(A3)

$$p^{(s)}(N_{H}) = \binom{N}{N_{H}} (\eta_{H} T_{\theta_{\text{out}}})^{N_{H}} (1-\eta_{H} T_{\theta_{\text{out}}})^{N-N_{H}},$$

(A4)

$$p^{(s)}(N_{V}) = \binom{N}{N_{V}} (\eta_{V} R_{\theta_{\text{out}}})^{N_{V}} (1-\eta_{V} R_{\theta_{\text{out}}})^{N-N_{V}},$$

(A5)

for an $N$-photon number state of light, and

$$p^{(s)}(N_{H},N_{V}) = e^{-\eta_{H} T_{\theta_{\text{out}}} N_{H}} (\frac{\eta_{H} T_{\theta_{\text{out}}} N_{H}}{N_{H}!}) e^{-\eta_{V} R_{\theta_{\text{out}}} N_{V}} (\frac{\eta_{V} R_{\theta_{\text{out}}} N_{V}}{N_{V}!}),$$

(A6)

$$p^{(s)}(N_{H}) = e^{-\eta_{H} T_{\theta_{\text{out}}} N_{H}} (\frac{\eta_{H} T_{\theta_{\text{out}}} N_{H}}{N_{H}!}),$$

(A7)

$$p^{(s)}(N_{V}) = e^{-\eta_{V} R_{\theta_{\text{out}}} N_{V}} (\frac{\eta_{V} R_{\theta_{\text{out}}} N_{V}}{N_{V}!}),$$

(A8)

for a coherent state of light with an average photon number $N$. Substituting these probabilities into equations (A1) and (A2), one can write the FI for the quantum scheme as

$$F_{N_{H},N_{V}}^{(q)}(\theta_{\text{out}}) = N \left( \frac{\eta_{H} T_{\theta_{\text{out}}} + \eta_{V} R_{\theta_{\text{out}}}}{\eta_{H} T_{\theta_{\text{out}}} - \eta_{V} R_{\theta_{\text{out}}}} \right)^2$$

$$+ \frac{N(\eta_{H} - \eta_{V})^2}{1 - \eta_{H} T_{\theta_{\text{out}}}} \left( \frac{\partial T_{\theta_{\text{out}}}}{\partial \theta_{\text{out}}} \right)^2,$$

(A9)

$$F_{N_{H}}^{(q)}(\theta_{\text{out}}) = \frac{N \eta_{H}}{T_{\theta_{\text{out}}} - \eta_{V} R_{\theta_{\text{out}}}} \left( \frac{\partial T_{\theta_{\text{out}}}}{\partial \theta_{\text{out}}} \right)^2,$$

(A10)

$$F_{N_{V}}^{(q)}(\theta_{\text{out}}) = \frac{N \eta_{V}}{R_{\theta_{\text{out}}} - \eta_{H} T_{\theta_{\text{out}}}} \left( \frac{\partial T_{\theta_{\text{out}}}}{\partial \theta_{\text{out}}} \right)^2,$$

(A11)

and for the classical scheme as

$$F_{N_{H},N_{V}}^{(c)}(\theta_{\text{out}}) = N \left( \frac{\eta_{H} T_{\theta_{\text{out}}} + \eta_{V} R_{\theta_{\text{out}}}}{\eta_{H} T_{\theta_{\text{out}}} - \eta_{V} R_{\theta_{\text{out}}}} \right)^2,$$

(A12)

$$F_{N_{H}}^{(c)}(\theta_{\text{out}}) = \frac{N \eta_{H}}{T_{\theta_{\text{out}}} - \eta_{V} R_{\theta_{\text{out}}}} \left( \frac{\partial T_{\theta_{\text{out}}}}{\partial \theta_{\text{out}}} \right)^2,$$

(A13)

$$F_{N_{V}}^{(c)}(\theta_{\text{out}}) = \frac{N \eta_{V}}{R_{\theta_{\text{out}}} - \eta_{H} T_{\theta_{\text{out}}}} \left( \frac{\partial T_{\theta_{\text{out}}}}{\partial \theta_{\text{out}}} \right)^2,$$

(A14)

where $(\partial T_{\theta_{\text{out}}}/\partial \theta_{\text{out}})^2 = 4 T_{\theta_{\text{out}}} R_{\theta_{\text{out}}}$ and $F(\theta_{\text{out}}) = F(T_{\theta_{\text{out}}}) \times (\partial T_{\theta_{\text{out}}} T_{\theta_{\text{out}}})^2$ is used with $F(T_{\theta_{\text{out}}})$ being the FI for the parameter $T_{\theta_{\text{out}}}$ for mathematical convenience. It is clear that the FI for the quantum scheme with a single-mode measurement is always greater than that for the classical scheme regardless of the values of $T_{\theta_{\text{out}}}$, $R_{\theta_{\text{out}}}$, and $\eta_{H,V}$. The quantum scheme with two-mode measurement, on the other hand, is beneficial compared to the classical scheme only when $\eta_{H} \neq \eta_{V}$. Note again that the FI is the same as the QFI for both the classical and quantum schemes considered in this work, so the lower bounds to the uncertainty associated with equations (A9)–(A14) are ultimately optimal within the framework of the CR inequality.

Appendix B. Uncertainty budget

Here we provide the uncertainty budget for type A and B uncertainties [57], which have the potential to influence the estimation of concentration $C$. The relevant quantities in this work are the wavelength-dependent specific rotation, the length of the cuvette, and the angles such as $\theta_{\text{in}}$ and $\theta_{\text{out}}$. For the specific rotation, we have used a fixed value of the wavelength in equation (2), but in reality no light is entirely monochromatic (i.e., see figure 1(b)), so that modulation of the wavelength in the reference calculation of the specific rotation needs to be considered in the estimation. Also, any realistic measurement of the length of the cuvette is far from ideal, so
the length might be slightly different from the assumed value of 0.1 dm. A similar behavior would be observed in the calibration of $\theta_{in}$. These effects are treated as type B uncertainty, whereas the measurement of $\theta_{out}$ is actually repeated, leading to type A uncertainty evaluation. Such relevant uncertainties are listed in table B1 for further information.

**Appendix C. Monte-Carlo simulation**

To understand the effects of the experimental imperfections involved in the results shown in figures 3 and 4, we perform a Monte-Carlo simulation by varying the relevant parameters, in an arbitrary range, which are typically constrained in a real experiment. The concentration is estimated from the data obtained through the numerical simulation by varying the input polarization $\theta_{in}$ from $0^\circ$ to $10^\circ$ in steps of $1^\circ$, providing a detailed study in the shaded area in figures 3 and 4. In order to see the effect of the PE in the estimation, we compare the case of an ideal extinction ratio ($r_{PE} \rightarrow \infty$) with the case of a realistic extinction ratio ($r_{PE} = 1000/1$) for $C = 0.1, 0.3,$ and $0.5 \text{ g ml}^{-1}$, shown in figure C1. It is shown that the estimated concentration is equal to the true value up to the numerical precision of sampling in the ideal case, whereas it drops down at $\theta_{out} = \theta_{in} + \alpha \approx 0^\circ$—the effect of PE is maximal, indicating an erroneous estimation in the realistic case. The effect of PE becomes less significant as $T_{\theta_{out}}$ approaches $1/2$ since the PE of H- and V-photons takes place almost symmetrically and thus compensates each other. Furthermore, note that a little deviation from the true value occurs at $\theta_{out} \approx 0^\circ$ even in the ideal case. Such an error can be shown to be independent of $\nu$ and $\mu$, and rather is a technical problem related to using the inverse function of equation (5) to estimate $\theta_{out}$. It has been overwhelmed by the effect of PE ($r_{PE} \sim 1000/1$) in the measured data shown in figures 3 and 4.

The observed behaviors shown in figure C1 can be explained via an appropriate modification to the theoretical development made in the main text. The effect of PE, i.e. the erroneous exchange between the H-photon and the V-photon with a rate $r_{PE}$, can be included in the theory by replacing $T_{\theta_{out}}$ by $T_{\theta_{out}} + r_{PE} \times \delta T_{\theta_{out}}$, where $\delta T_{\theta_{out}} = 1 - 2T_{\theta_{out}}$. Equations (3)–(5) can thus be written up to the first order of $r_{PE} \ll 1$ as

\begin{align}
(i) \quad & \frac{\langle N_H \rangle}{\nu} \approx \langle f_{\text{single}} \rangle + r_{PE} \times \delta f_{\text{single}}, \\
(ii) \quad & \frac{\langle N_H - N_V \rangle}{\nu} \approx \langle f_{\text{diff}} \rangle + r_{PE} \times \delta f_{\text{diff}}, \\
(iii) \quad & \frac{\langle N_H - N_V \rangle}{N_H + N_V} \approx \langle f_{\text{DSR}} \rangle + r_{PE} \times \delta f_{\text{DSR}},
\end{align}

where

\begin{align}
\delta f_{\text{single}} &= (1 - 2T_{\theta_{out}})\eta_H, \\
\delta f_{\text{diff}} &= (1 - 2T_{\theta_{out}})(\eta_H + \eta_V), \\
\delta f_{\text{DSR}} &= (1 - 2T_{\theta_{out}})\left[\frac{2\eta_H\eta_V}{[T_{\theta_{out}}\eta_H + (1 - T_{\theta_{out}})\eta_V]^2}\right].
\end{align}

**Table B1.** The uncertainty budgets of type A and B in our study. N and R represent the normal and rectangular distributions, respectively. This table is considered at $\theta_{in} = 40^\circ$ and $C = 0.5 \text{ g ml}^{-1}$.

| Source of uncertainty | Value | Unit | Divisor (distribution) | Sensitivity | Standard uncertainty (g ml$^{-1}$) |
|-----------------------|-------|------|------------------------|-------------|-----------------------------------|
| $\theta_{out}$ obtained by $f_{\text{single}}$ | 0.2708 | degree ($^\circ$) | 1 (N) | 0.293 | 0.0793 |
| $\theta_{out}$ obtained by $f_{\text{diff}}$ | 0.1841 | degree ($^\circ$) | 1 (N) | 0.293 | 0.0539 |
| $\theta_{out}$ obtained by $f_{\text{DSR}}$ | 0.1763 | degree ($^\circ$) | 1 (N) | 0.293 | 0.0517 |

Specific rotation [$\alpha(\lambda)$] 0.41 degree ($^\circ$) -1 g$^{-1}$ ml |

Length of cuvette $l$ 0.0005 dm |

Input polarization $\theta_{in}$ 0.03 degree ($^\circ$) |
It is clear to see that the effect of the PE for the three estimators is maximal when \( T_{\theta_{\text{out}}} = 0 \) or 1, whereas it vanishes when \( T_{\theta_{\text{out}}} = 1/2 \).

All the errors \( \delta \) in equations (C4)–(C6) lead to an equal amount of error in the estimation of \( \theta_{\text{out}} \), i.e. \( \theta_{\text{out}} \rightarrow \theta_{\text{out}} + r_{\text{PE}} \delta_{\theta_{\text{out}}} \), where the error

\[
\delta_{\theta_{\text{out}}} = \frac{\cos 2\theta_{\text{out}}}{|\sin 2\theta_{\text{out}}|} = \frac{2T_{\theta_{\text{out}}} - 1}{2 \sqrt{T_{\theta_{\text{out}}}(1 - T_{\theta_{\text{out}}})}}.
\]  

(C7)

The error \( \delta_{\theta_{\text{out}}} \) caused by the PE diverges as \( T_{\theta_{\text{out}}} \rightarrow 0 \) or 1, while it decreases as \( T_{\theta_{\text{out}}} \) approaches 1/2.

The PE consequently causes an error in the uncertainty \( \Delta \theta_{\text{out}} \) of equations (8)–(10), i.e. \( \Delta \theta_{\text{out}} \rightarrow \Delta \theta_{\text{out}} + r_{\text{PE}} \times \delta \Delta \theta_{\text{out}} \), where \( \Delta \theta_{\text{out}} \) can be found as

\[
\delta \Delta \theta_{\text{out}}^{\text{single}(q)} = \frac{1}{\sqrt{2}} \frac{1 - 2T_{\theta_{\text{out}}}}{8} \frac{1}{(\eta H - 1)} \frac{1}{(1 - T_{\theta_{\text{out}}})^2} \frac{1}{\Delta \theta_{\text{out}}^{\text{single}(q)}}.
\]  

(C8)

\[
\delta \Delta \theta_{\text{out}}^{\text{diff}(q)} = \frac{1}{\sqrt{2}} \frac{1 - 2T_{\theta_{\text{out}}}}{8} \frac{1}{(\eta H + \eta V)} \left( \frac{\eta H (1 - \eta H) - \eta V (1 - \eta V)}{(1 - T_{\theta_{\text{out}}})^2} \right) \frac{1}{\Delta \theta_{\text{out}}^{\text{diff}(q)}}.
\]  

(C9)

\[
\delta \Delta \theta_{\text{out}}^{\text{DSR}(q)} = \frac{1}{\sqrt{2}} \frac{1 - 2T_{\theta_{\text{out}}}}{8} \frac{1}{\eta V} \frac{1}{\eta H} \frac{1}{\Delta \theta_{\text{out}}^{\text{DSR}(q)}}.
\]  

(C10)

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