Optical Amplification of Spin Noise Spectroscopy via Homodyne Detection

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Spin noise (SN) spectroscopy measurements on delicate semiconductor spin systems, like single (In,Ga) As quantum dots, are currently not limited by optical shot noise but rather by the electronic noise of the detection system. We report on a realization of homodyne SN spectroscopy enabling shot-noise-limited SN measurements. The proof-of-principle measurements on impurities in an isotopically enriched rubidium atom vapor show that homodyne SN spectroscopy can be utilized even in the low-frequency spectrum, which facilitates advanced semiconductor spin research like higher order SN measurements on spin qubits.

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I. INTRODUCTION

Traditional optical experiments study the spin dynamics in semiconductors by the optical excitation of spinpolarized carriers, such as in polarization-resolved photoluminescence or pump-probe Faraday rotation experiments [1,2]. However, these techniques are problematic in spin systems where the perturbation by optical excitation changes the intrinsic carrier spin dynamics. This adverse effect applies to weakly interacting and few-particle spins [3-5], as well as spin systems where an additional interaction is induced by optically injected carriers [6,7]. For such fragile semiconductor spin systems, spin noise (SN) spectroscopy has been established as the method of choice to avoid unnecessary optical excitation [8]. The experimental technique was transferred from quantum optics to semiconductor physics in 2005 [9] and is now used to study, e.g., the spin dynamics of single carriers and trions in single (In,Ga)As quantum dots (QD) [5,10] or the complex interaction of localized electrons with nuclear spins [11]. In this context, SN spectroscopy is used either as a weakly disturbing measurement method at thermal equilibrium [12] or to study quantum spin systems which are strongly driven by a resonant light field [10,13]. In the first case, very low light intensities are required [4,5] and, in the second case, SN at high frequencies [14,15] and high sensitivity [10] is of special interest. In both scenarios, the dominant experimental noise source is not SN or optical shot noise but rather extrinsic electronic noise from the electro-optical detectors entailing long measuring times. This constraint, inter alia, impedes fundamental measurements like high-frequency single spin dynamics or the efficient measurement of higher correlation SN.

The impact of electronic noise can be diminished by optical amplification. The technique is known in quantum optics as homodyne detection, and we show in this paper that homodyne detection [16–18] can be transferred smoothly to a standard SN spectroscopy setup even in the regime of low noise frequencies. For high frequencies, Cronenberger and Scalbert recently demonstrated quantum-limited heterodyne detection of SN in GaAs [18] in a nonstandard SN setup. Here, we demonstrate the optical amplification of the SN signal on 85Rb impurities in an isotopically enriched ⁸⁷Rb vapor, which is a wellcharacterized reference system [13,19,20]. The same technique will also be applicable to delicate, low-frequency semiconductor spin systems like a single, charged (In,Ga) As quantum dot at low temperatures, where the electronic noise is a dominating obstacle in fast and precise measurements of the spin dynamics. The setup and data presented in Fig. 1 demonstrate the typical dominance of electronic noise in current state-of-the-art QD SN measurements. Such low-dimensional semiconductor spin systems with long spin lifetimes are of special interest due to their prospective potential as spin qubits in quantum information processing [21–23].

II. METHODS

A. Spin noise spectroscopy

Spin noise spectroscopy in semiconductors maps the stochastic spin fluctuations of carriers at thermal equilibrium onto the polarization of laser light that is quasiresonant to an optical transition [8]. Figure 1(a) depicts a typical schematic experimental setup for SN measurements on a single semiconductor QD in reflection. Here, linearly polarized laser light is focused on a single (In,Ga)As QD which is embedded in an asymmetric AlAs/GaAs Bragg resonator and charged by a single hole. The stochastic spin polarization of the hole yields a Faraday rotation of the linear polarization of the laser light [8] which is measured by a polarization bridge and a balanced

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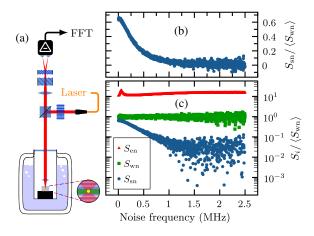


FIG. 1. (a) Schematic of a typical SN spectroscopy setup for single-quantum-dot SN measurements in reflection (see Ref. [10] for details). (b) Typical SN power spectrum of a single (In,Ga)As quantum dot charged by a hole in units of the probe laser shot noise power. (c) Comparison on a logarithmic scale of the broadband electronic noise of the balanced receiver $S_{\rm en}$, the white photon shot noise $S_{\rm wn}$, and the SN $S_{\rm sn}$ for a typical quantum-dot SN measurement.

receiver. The time-continuous electrical signal is amplified, digitized, and Fourier transformed. The resulting Fourier power spectra are averaged (S_{fg}) , and an averaged background spectrum (S_{bg}) is subtracted. The background spectrum is measured by applying a transverse magnetic field which shifts the SN to higher frequencies outside of the detection bandwidth for the measurement depicted in Fig. 1. Figure 1(b) shows exemplarily such a single-quantum-dot SN spectrum for a laser intensity of 1.1 μ W/ μ m², while Fig. 1(c) compares the corresponding noise contributions of the electronic noise $(S_{en}, \text{ the red dots})$, the white optical shot noise $(S_{wn}$, the green dots), and the weak SN $(S_{sn}$, the blue dots). The dominant electronic noise results mainly from the commercial low-noise balanced photoreceiver, which, in this typical SN OD experiment, is about 1 order of magnitude larger than the optical shot noise; i.e., the electronic noise increases the necessary measurement time by 2 orders of magnitude. Intensitydependent measurements (not shown) reveal that, nevertheless, the depicted SN spectrum is strongly influenced by the light field despite the rather low probe laser intensity [5]. In fact, orders of magnitude lower laser intensities are desirable for such SN experiments whereat optical amplification by homodyne detection becomes essential.

B. Homodyne detection

In this section, we briefly introduce the principle of homodyne amplification in the context of standard SN spectroscopy. Optical homodyne detection utilizes two phase-locked, monochromatic laser beams which can be described as classical electromagnetic plane waves. For SN spectroscopy, only the electric fields and the polarization states of the two beams are relevant, both of which can be represented by two Jones vectors in a linearly polarized basis [24]. The vertically polarized signal beam travels through the sample and experiences a small stochastic Faraday rotation by an angle θ_F . The resulting electric-field vector is

$$\mathbf{E}_{\text{sig}} = E_{\text{sig},0} R(\theta_F) \cdot \mathbf{e}_V = E_{\text{sig},0} \begin{pmatrix} \sin(\theta_F) \\ \cos(\theta_F) \end{pmatrix}, \quad (1)$$

where $E_{\rm sig,0}$ is the complex amplitude, $R(\theta_F)$ is the regular Euclidean rotation matrix, and \mathbf{e}_V is the polarization vector. The second beam yields the local oscillator field and acts as an unchanged polarization reference,

$$\mathbf{E}_{lo} = \sqrt{2I_{lo}/\epsilon_0 c} \cdot \mathbf{e}_V = \sqrt{2I_{lo}/\epsilon_0 c} \cdot \begin{pmatrix} 0\\1 \end{pmatrix}, \quad (2)$$

where $I_{\rm lo}=p\epsilon_0c|E_{\rm sig,0}|^2/2=pI_{\rm sig}$ is the intensity of the local oscillator in terms of the intensity of the signal beam and p is the corresponding proportionality factor. For SN spectroscopy, the interference of both beams ${\bf E}_{\rm if}$ is decomposed into a linearly polarized basis rotated by $\pi/4$ (diagonal ${\bf e}_D$ and antidiagonal ${\bf e}_A$), giving the components $E_{\rm if}$, and $E_{\rm if}$,. The balanced receiver measures the difference of the intensities of these two components:

$$\Delta I = |E_{if,D}|^2 - |E_{if,A}|^2$$

$$= 2I_{sig} \sin(\theta_F) [\cos(\theta_F) + \sqrt{p} \cos(\varphi_{sig,lo})]$$

$$= 2I_{sig} \theta_F [1 + \sqrt{p} \cos(\varphi_{sig,lo})] + O(\theta_F^2), \qquad (3)$$

where $\varphi_{\rm sig,lo}$ is the phase between ${\bf E}_{\rm sig}$ and ${\bf E}_{\rm lo}$ which is typically set to a multiple of 2π for optimal amplification. Typical SN measurements involve very small Faraday rotation angles and, for this reason, the use of small angle approximation is justified. In this case, the intensity difference is linear in θ_F and $(1+\sqrt{p})$. Finally, the SN signal is proportional to ΔI^2 and consequently scales by the factor $\eta_p=(1+\sqrt{p})^2$. This dependence allows the signal-to-noise ratio of the SN signal to be increased without disturbing the measured system by increasing the local oscillator intensity up to the point where optical shot noise dominates over electronic noise. In practice, this kind of amplification is limited only by the maximally allowed optical power incident on the photodiodes of the balanced receiver.

C. Noise power and signal-to-noise ratio

Spin noise is typically evaluated by the SN power density spectrum $S_{\rm sn}(f)$, which follows, for

monoexponential spin-relaxation dynamics, a normalized Lorentzian profile of area 1 multiplied by the SN power amplitude A. The Lorentzian is shifted in a transverse magnetic field by the Larmor spin-precession frequency f_L and has a width γ_s given by the spin dephasing rate:

$$L_{f_L,\gamma_s}(f) = \frac{2\gamma_s}{\gamma_s^2 + 4\pi^2 (f - f_L)^2}.$$
 (4)

The *measured* SN power, i.e., the integral over $S_{\rm sn}$, scales with the square of the probe laser power [8], and thus the SN power density scales with the same factor if γ_s is constant. Homodyne detection amplifies $S_{\rm sn}$ by approximately η_p . Laser photon shot noise yields an additional broadband white-noise contribution that scales linearly with the laser power on the photodiodes. The total laser power at the homodyne operation point of constructive interference ($\varphi_{\rm sig,lo}=0$) reads

$$P_L = (\sqrt{P_{\rm sig}} + \sqrt{pP_{\rm sig}})^2 = \eta_p P_{\rm sig}.$$

The electronic noise is, in general, independent of the incident laser power, resulting in the total noise power density spectrum

$$S(f) = AL_{f_t, \gamma_s} \alpha^2 P_{\text{sig}}^2 \eta_p \quad \text{(spin noise } S_{\text{sn}})$$
 (5a)

$$+W\hbar\omega_L\alpha P_{\rm sig}\eta_p$$
 (shot noise $S_{\rm wn}$) (5b)

$$+S_{\rm en}$$
 (electronic noise), (5c)

where W is a constant, $\hbar\omega_L$ is the laser photon energy, and $\alpha(f)$ is the (unitless) detector response function.

The signal-to-noise ratio is given by the spin noise signal $S_{\rm sn}$ [Eq. (5a)] divided by the nonsignal contributions $S_{\rm wn}$ [Eq. (5b)] and $S_{\rm en}$ [Eq. (5c)]. Here, however, in order to consistently analyze the performance of homodyne detection, we define the signal-to-noise ratio $r_{\rm SNR}^{\star}$ as the ratio of the SN amplitude and the spectral mean of the remaining broadband noise contributions:

$$r_{\rm SNR}^{\star} = \frac{\max(S_{\rm sn})/P_{\rm sig}}{\langle S_{\rm en}\rangle + \langle S_{\rm wn}\rangle} = \frac{AL_{f_L,\gamma_s}(f_L)\alpha^2(f_L)P_L}{\langle S_{\rm en}\rangle + W\hbar\omega_L\langle\alpha\rangle P_L}, \quad (6)$$

where the SN amplitude is divided by the probe laser power $P_{\rm sig}$ such that $r_{\rm SNR}^{\star}$ depends directly on the total laser power P_L detected on the photodiodes. This definition makes $r_{\rm SNR}^{\star}$ robust against fluctuations in $P_{\rm sig}$, and the auxiliary parameters $P_{\rm sig}$ and P_L can be easily recorded during the measurement. The denominator of $r_{\rm SNR}^{\star}$ is the mean of the background spectrum $S_{\rm bg}$ that is measured by shifting the spin noise outside of the spectral window considered, while the numerator is recovered from the

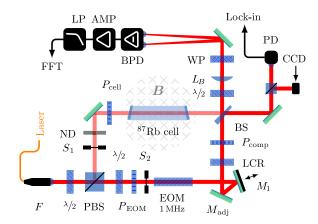


FIG. 2. Schematic drawing of the homodyne setup with fiber output from the laser source (F), a polarizing beam splitter (PBS), a high-extinction linear polarizer (P_i) , a mechanical shutter (S_i) , an electro-optic phase modulator (EOM), a mirror with a piezoelectric actuator (M_1) , a liquid-crystal retarder (LCR), a neutral density filter (ND), a low-wave-front-distortion nonpolarizing beam splitter (BS), a Wollaston prism (WP), a lens (L_B) , a high-speed balanced photoreceiver for SN (BPD) whose output is amplified (AMP) and low-pass filtered (LP), a photodetector for phase stabilization (PD), and an imaging CCD array for interferometer adjustment (CCD). A solenoid underneath the 87 Rb vapor cell yields an adjustable, transverse magnetic field B in order to shift the Larmor frequency (the center frequency of $S_{\rm sn}$).

measured difference spectrum $S_{\rm sn} \approx S_{\rm diff} = S_{\rm fg} - S_{\rm bg}$ divided by the measured laser probe power $P_{\rm sig}$. With this procedure $r_{\rm SNR}^{\star}$ can be easily estimated in the experiment as

$$r_{\rm SNR}^{\star} = \max{[(S_{\rm fg} - S_{\rm bg})/P_{\rm sig}]/\langle S_{\rm bg}\rangle}. \tag{7}$$

III. SETUP

A. Interferometer

Figure 2 depicts the homodyne SN spectroscopy setup, which is derived from a Mach-Zehnder interferometer. A half-wave retarder and a polarizing beam splitter (PBS) split the incoming laser beam with a power ratio between the transmitted and reflected beams of approximately 5:1. In this configuration, the transmitted beam forms the local oscillator (LO) path, and the reflected beam the signal path. A neutral density filter reduces the intensity in the signal path by 3 orders of magnitude so that the probe intensity at the ⁸⁷Rb sample cell is, for all of the presented experiments, about 250 μ W/cm² (approximately 8 μ W for a beam of 2 mm diameter). A linear polarizer in front of the ⁸⁷Rb cell ensures a well-defined polarization. The laser light in the LO path is passed through a linear polarizer and an electrooptic phase modulator in order to (a) precisely redefine the linear polarization and (b) modulate the phase of the electric field with 1.111 MHz producing the error signal for the phase stabilization feedback. A mirror mounted on a piezoelectric actuator allows us to adjust the phase by up to 8 rad and is used to stabilize the interferometer at the desired operating point. A liquid-crystal retarder and an additional polarizer P_{comp} allow for the adjustment of the laser power in the LO arm. The signal and the LO path overlap on a wedged nonpolarizing beam splitter (BS) and proceed jointly to the SN measurement port (the top port in Fig. 2) and the control port (the right port in Fig. 2). The power in the control port is sampled by a photodetector (PD) and demodulated by a lock-in amplifier which effectively detects the phase between \mathbf{E}_{sig} and \mathbf{E}_{lo} . Using M_1 the phase is steered by a proportional-integral-derivative loop towards the operating point, i.e., constructive interference in the measurement port with an accuracy of $\delta \varphi \approx 0.13$ rad. Power measurements on both paths are realized by blocking the other path by one of two mechanical shutters, S_1 or S_2 , and sampling the remaining power on the photodetector. The polarization alterations in the measurement port are detected by a polarization bridge consisting of a halfwave retarder, a Wollaston prism, and a high-speed balanced photoreceiver. The output of the balanced photoreceiver [25] is postamplified and low-pass filtered to a bandwidth of 5 MHz (3 dB). The resulting voltage signal is sent to an analyzing computer that samples the voltage with a fast, 12-bit digital-to-analog converter operating at 20 MHz and performs a discrete fast Fourier transform. Please note that only electronic noise from the transimpedance amplifier in the first stage of the balanced photoreceiver has a significant influence on the experiment, whereas the electronic noise contribution of the intermediate voltage amplifier, 2.5 nV/ $\sqrt{\text{Hz}}$ ("postamplification"), and the digitizer, $80 \text{ nV}/\sqrt{\text{Hz}}$, can be neglected. We selected at the balanced photoreceiver two different transimpedance gains of $1 \times 10^4 \text{ V A}^{-1}$ and $1 \times 10^5 \text{ V A}^{-1}$ corresponding to noise-equivalent-power densities of 10 and 2.5 pW/ $\sqrt{\text{Hz}}$, respectively, in order to change the ratio of electronic versus optical shot noise present in the detection. Both amplifications have a bandwidth larger than the auxiliary low-pass filter of 5 MHz. Thus, we alter via a diametrically set amplification by the subsequent voltage amplifier the relative contribution of electronic noise with respect to spin and shot noise.

B. Sample and experimental details

The sample is a ${}^{87}\text{Rb}$ vapor cell with helium buffer gas and a purity of the ${}^{87}\text{Rb}$ isotope of > 98%. The goal of the presented setup is the optical amplification of extremely small SN signals. Therefore, the weak SN signature of the remaining ${}^{85}\text{Rb}$ isotope is analyzed and the measurement is carried out at room temperature, where the Rb vapor partial pressure is very low. The vapor density of $n \approx 1.2 \times 10^{10}\text{cm}^{-2}$ and optical probe power of $8\,\mu\text{W}$ used in the

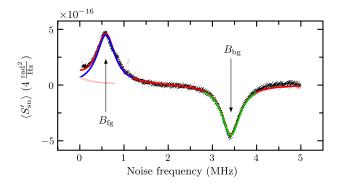


FIG. 3. Composite average of the measured SN difference spectrum of 85 Rb (the dots) [cf. Eq. (8)] in units of Faraday rotation power spectral density. The solid red line shows a fit with two Lorentzians which are shifted by the Larmor frequencies corresponding to $B_{\rm fg}$ and $B_{\rm bg}$. The field values of $B_{\rm fg} = 120~\mu{\rm T}$ and $B_{\rm bg} = 800~\mu{\rm T}$ are chosen so that both peaks fit well inside the available bandwidth. The folding of the foreground peak bears no drawback for the evaluation and nicely demonstrates the conservation of spin noise power.

presented measurements are significantly lower than the optimal values reported on in Ref. [26]. These parameters have been chosen purposely, in order to simulate the unfavorable conditions present in the single-quantum-dot setups that are the main scope of our technique.

The laser source used in this setup is an external cavity diode laser stabilized by a Fizeau interferometer and the laser photon energy is blueshifted from the 85 Rb D_2 resonance by about 0.3 GHz. A solenoid below the vapor cell produces a transverse magnetic field which modulates the stochastic spin orientation by the Larmor frequency f_L . In order to separate the weak SN signature from other noise contributions, a foreground and a background SN spectrum are measured at a magnetic field of $B_{\rm fg}=120~\mu{\rm T}$ and $B_{\rm bg}=800~\mu{\rm T}$, respectively, and the two spectra are subtracted from each other. All other noise contributions are independent of the magnetic field and average out in the difference of the foreground and background spectra, whereat only the SN contribution remains.

IV. RESULTS

Figure 3 depicts a high-quality SN spectrum which is obtained by averaging over SN spectra [27] measured at different amplifications factors η_p and normalized by P_{sig}^2 :

$$S'_{\rm sn} = \frac{S(B = B_{\rm fg}) - S(B = B_{\rm bg})}{P_{\rm sig}^2 \eta_p}.$$
 (8)

The solid (red) line is a fit to the spectrum consisting of a positive (foreground; the blue line) and a negative (background; the green line) Lorentzian, whereat part of the foreground Lorentzian is folded back at zero frequency (the

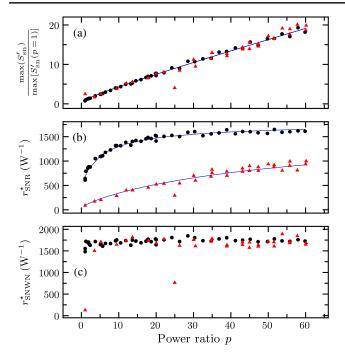


FIG. 4. (a) Measured SN amplitude in dependence on p normalized to the SN amplitude at p=1. The solid line is a plot of η_p/η_1 . (b) Signal-to-noise ratio for weak (black circles) and large electronic noise (red triangles). The solid line is a fit according to Eq. (6). (c) Measured ratio of SN and photon shot noise.

pink line). The extremely accurate fit allows the extraction of the noise powers, the center frequencies f_L , and the full widths at half maximum γ_s with high precision. Both parameters, f_L and γ_s , are constant in the experiments such that the spin noise amplitude is a direct measure of optical amplification in the following evaluation of the measurements.

First, we study the optical amplification of the SN that depends on the light intensity of the local oscillator. Here, we change the power ratio p only, not $P_{\rm sig}$. Figure 4(a) shows the amplitude of the Lorentzian SN peak normalized to the amplitude at p=1 as a function of p. As predicted by Eq. (5), the amplitude scales with $\eta_p=(1+\sqrt{p})^2$ (the solid line). This is a direct experimental proof of optical amplification but does not yet prove to be an improvement on the signal-to-noise ratio.

All of the experiments shown in Fig. 4 are carried out for two different scenarios, (i) high electrical amplification of the balanced receiver of 6×10^4 V W⁻¹ and low postamplification of 30 dB (the black circles), and (ii) lower electrical amplification of the balanced receiver of 6×10^3 V W⁻¹ and higher postamplification of 45 dB (the red triangles). The typical amplifier characteristic of the balanced receiver yields for the case (i) lower electronic noise than for the case (ii); i.e., the black circles and the red triangles show measurements with different effective electronic noise contributions. The data for the optical

amplification of SN is depicted for both cases in Fig. 4(a) and shows that the optical amplification of SN is independent of the background electronic noise.

Next, we demonstrate that homodyne amplification also increases the signal-to-noise ratio $r_{\rm SNR}^{\star}$ if electronic noise is the dominating noise source. Figure 4(b) clearly shows the increase of $r_{\rm SNR}^{\star}$ with an increasing power ratio p. In the case of moderate electronic noise (the black circles), $r_{\rm SNR}^{\star}$ increases with an increasing p value and approaches saturation for a high p value since white photon shot noise dominates over electronic noise; i.e., homodyne detection with nonsqueezed light cannot beat the photon shot noise limit here [28]. In the case of higher electronic noise (the red triangles), $r_{\rm SNR}^{\star}$ also increases with an increasing p value but is significantly smaller at low p and does not reach saturation for the maximal p value in this experiment.

Finally, Fig. 4(c) shows the ratio $r_{\rm SNWN}^{\star}$, which is obtained by scaling the measured SN power normalized by $P_{\rm sig}$ with the measured optical shot noise power. This ratio is consistently constant since the optical shot noise and the SN power increase for homodyne detection by the same proportionality factor η_p , with $P_{\rm sig}$ staying constant. Here, $r_{\rm SNWN}^{\star}$ being independent of p affirms the assumptions and the accuracy of our model.

V. CONCLUSIONS

In this paper, we show in a proof-of-principle experiment that the resulting constraints by electronic noise can be successfully circumvented by optical homodyne amplification of spin noise in a standard spin noise spectroscopy setup—even at low frequencies. This low-frequency limit is especially important for semiconductor systems with long spin coherence times, which will be relevant in the framework of future semiconductor spin information processing. Here, spin noise spectroscopy on fragile spin systems—particularly on single (In,Ga)As quantum dots which is usually limited by the finite electronic noise of the available balanced receivers, can be made feasible. The experiments were successfully carried out with a noncommercial diode laser down to SN frequencies of tenths of kilohertz, which is sufficient for most spin noise experiments. Nevertheless, even lower spin noise frequencies should be easily accessible utilizing stabilized ultranarrow linewidth lasers.

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APPENDIX: AN ALTERNATIVE TO PHASE STABILIZATION

An active phase stabilization loop significantly increases the setup complexity. Here, we propose an alternative scheme that explicitly averages over a changing phase $\varphi_{\rm sig,lo}$.

The homodyne amplification has a cosine-type phase dependence which can be averaged out if the phase modulation (a) acts on a different time scale than the spin dynamics and (b) is faster than the characteristic drift of the setup. For a modulation much slower than the rate $f_{\rm rec}$ at which spectra are recorded, the phase has to be effectively constant. For a modulation much faster than $f_{\rm rec}$, the phase dependence must average out for each individual spectrum. For the measured noise power density [roughly the square of Eq. (3)], it is easy to show that

$$\overline{\Delta I^2} = \int_0^{n2\pi} \Delta I^2 \frac{d\varphi}{n2\pi} \propto I_{\text{sig}}^2 \theta_F^2 \left(\frac{2+p}{2}\right) + O(\theta_F^3). \quad (A1)$$

This kind of *phase dithering* reduces the effective amplification factor of the spin noise signal to $\eta_{\text{dith},p} = [(2+p)/2]$, while keeping all other properties of the homodyne detection. For the technical realization of this scheme, the piezoelectrically actuated mirror M_1 has to be replaced ideally by a phase modulator with a very high modulation depth to ensure an optimal phase averaging.

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- [27] The SN difference spectra often exhibit a small drift term S_{drift} which arises from the imperfect cancellation of the different noise sources. This drift term can be easily subtracted from the individual difference spectra.
- In order to guarantee the most consistent results, all presented data are corrected accordingly in the same manner.
- [28] V. G. Lucivero, R. Jimenez-Martinez, J. Kong, and M. W. Mitchell, Squeezed-light spin noise spectroscopy, Phys. Rev. A **93**, 053802 (2016).