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Observing and Verifying the Quantum Trajectory of a Mechanical Resonator

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Continuous weak measurement allows localizing open quantum systems in state space and tracing out their quantum trajectory as they evolve in time. Efficient quantum measurement schemes have previously enabled recording quantum trajectories of microwave photon and qubit states. We apply these concepts to a macroscopic mechanical resonator, and we follow the quantum trajectory of its motional state conditioned on a continuous optical measurement record. Starting with a thermal mixture, we eventually obtain coherent states of 78% purity—comparable to a displaced thermal state of occupation 0.14. We introduce a retrodictive measurement protocol to directly verify state purity along the trajectory, and we furthermore observe state collapse and decoherence. This opens the door to measurement-based creation of advanced quantum states, as well as potential tests of gravitational decoherence models.

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Within the Copenhagen interpretation of quantum mechanics, the quantum state of an isolated physical system is represented by its wave function. This mathematical object encodes the probability of possible measurement outcomes and contains the maximum possible knowledge about the system. Under the usually inevitable coupling of the system to an unknown environment, the state evolves into a statistical mixture of quantum states in a process known as decoherence. The mixture is described via a density matrix ρ, which again encodes measurement probabilities, while accounting for the ignorance about system-environment interactions. Decoherence entails the disappearance of some of the most salient, and useful, features of quantum mechanics, such as superposition and entanglement. However, if information becomes available on how the system has interacted with the environment, it is possible to restore and retain the purity of the quantum state (i.e., the extent to which the mixture is dominated by a single random but known wave function). Measurements can yield such information; over a finite time interval, however, the obtained information is often incomplete [1,2]. The density matrix can anyways be updated, however, the obtained information is often incomplete [1,2]. The density matrix can anyways be updated, however, the obtained information is often incomplete [1,2]. The density matrix can anyways be updated, however, the obtained information is often incomplete [1,2]. The density matrix can anyways be updated, however, the obtained information is often incomplete [1,2]. The density matrix can anyways be updated, however, the obtained information is often incomplete [1,2]. The density matrix can anyways be updated, however, the obtained information is often incomplete [1,2]. The density matrix can anyways be updated, however, the obtained information is often incomplete [1,2]. The density matrix can anyways be updated, however, the obtained information is often incomplete [1,2]. The density matrix can anyways be updated, however, the obtained information is often incomplete [1,2]. The density matrix can anyways be updated, however, the obtained information is often incomplete [1,2]. The density matrix can anyways be updated, however, the obtained information is often incomplete [1,2].

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In the continuous limit of many weak subsequent measurements carried out over short times, the state conditioned on a measurement record traces out the system’s quantum trajectory in time [3]. Observing pure quantum trajectories is a challenging task, so far it has been achieved only in very clean settings such as cavity [4] and circuit [5,6] QED. Here, we extend these ideas to measurements of the motion of a macroscopic mechanical resonator [7–12]. In this setting, pure conditional states are obtained through measurements of high efficiency ηmeas = Γmeas/(γ + Γqba), where Γmeas is the measurement rate; and γ and Γqba are decoherence rates induced by a thermal bath and the measurement quantum backaction, respectively [13,14]. Prior experiments on motional state estimation have remained confined to the classical regime, due the fast decoherence by the thermal bath [15–18]. In contrast, by probing the system with a measurement for which the efficiency reaches ηmeas ≈ 67%, we are able to observe individual quantum trajectories of highly pure conditional states. Moreover, building on recent theoretical and experimental work on retrodiction [10,11,19,20] and past quantum states [21,22], we introduce a retrodiction-based trajectory-verification technique, and we use it to confirm the purity of the conditional state by statistical analysis of ensembles of trajectories. This allows us to directly observe the collapse of the conditional state, as
well as the decoherence that occurs in the absence of information from the measurement.

We explore these ideas in an optomechanical system based on an ultracoherent soft-clamped membrane resonator [23]. The mechanical mode of interest (at the frequency $\Omega_m/2\pi = 1.14$ MHz) corresponds to a localized defect mode created within a phononic crystal. This design simultaneously reduces radiative energy loss and avoids loss-inducing mode curvature, resulting in an extremely low mechanical energy dissipation rate, $\Gamma_m$. For the device used here, we find $Q = \Omega_m/\Gamma_m = 1.03 \times 10^9$ at temperature $T = 11$ K. This motion is dispersively coupled to the frequency of a Fabry-Perot cavity mode [linewidth of $\kappa/(2\pi) = 18.5$ MHz] at a characteristic vacuum optomechanical coupling rate of $g_0/(2\pi) = 129$ Hz. Populating the cavity with a large coherent field (with $\bar{n}_{cav}$ average photons) leads to a linearized field-enhanced coupling at a rate of $\bar{g} = \sqrt{\bar{n}_{cav}}g_0$. A probe laser that drives this cavity resonantly will acquire phase modulation sidebands proportional to the mechanical displacement, which we detect via a balanced homodyne receiver. The total detection efficiency (including cavity outcoupling) is $\eta_{det} = 74\%$, ensuring that minimal mechanical information is lost.

In addition to a resonant probe beam, we also utilize an auxiliary beam to provide some precoupling of other modes of the membrane, via both sideband and feedback cooling [24]. The effect of this beam on the main mode of interest is simply to change its effective thermal environment. Small, residual detuning of the probe similarly provides some damping. In the following, we account for both of these and refer to the effective energy damping rate and bath occupancy as, respectively, $\Gamma_m/2\pi \approx 130$ Hz and $\bar{n}_{th} = 2$ such that $\gamma = \Gamma_m\bar{n}_{th} \approx 2\pi \times 260$ Hz.

The quantum measurement backaction of the probe is manifest as radiation pressure force fluctuations, leading to additional mechanical decoherence at a rate of $\Gamma_{qba} = 4\bar{g}^2/\kappa \approx 2\pi \times 2.54$ kHz. Similarly, the mechanical displacement measurement can be characterized by a measurement rate of $\Gamma_{meas} = 4\eta_{det}\bar{g}^2/\kappa \approx 2\pi \times 1.88$ kHz. Thus, the experimental system studied here can achieve measurements in which quantum backaction dominates thermal motion, and the measurement rate approaches the total decoherence rate (i.e., the measurement efficiency $\eta_{meas} \rightarrow 1$).

The quantum trajectory of the mechanical resonator is derived from quadratures $\hat{f} = (\hat{x}, \hat{y})$ of the homodyne photocurrent, $I(t)$, demodulated at frequency $\Omega_m$. This demodulation, which occurs in postprocessing, uses a high-order low-pass filter for which the $\sim 120$ kHz bandwidth is significantly larger than the total decoherence rate ($\gamma + \Gamma_{qba} \approx 2\pi \times 2.80$ kHz) such that it has a negligible effect on the mechanical signal [7,25]. The continuous data stream is subdivided into 3.2 ms segments: each of which is treated as an individual experimental realization. Examples of the raw photocurrent and one demodulated quadrature are shown in Fig. 1(b).

These photocurrent quadratures form the measurement channels from which we extract the quantum trajectory. This is done according to a stochastic master equation (SME), which simultaneously describes unitary evolution of the system, environmental coupling, and Bayesian updates according to the measurement record [12,26,27]. Note that, although our system is composed of both an optical mode and a mechanical mode, the cavity field can be adiabatically eliminated because $\kappa \gg \Omega_m$ [7,28]. For a high-$Q$ resonator, it is also convenient to move to the interaction picture at frequency $\Omega_m$ and make a rotating wave approximation. Thus, we describe the system in terms of the slowly varying quadratures $\hat{f} = (\hat{X}, \hat{Y})$, where the mechanical position is $\hat{q} = \hat{X}\cos(\Omega_m t) + \hat{Y}\sin(\Omega_m t)$. The corresponding SME is [7]

$$d\rho = (\mathcal{L}_{th} + \mathcal{L}_{qba})\rho dt + \sqrt{\Gamma_{meas}(\mathcal{H}[\hat{X}]\rho dW_X + \mathcal{H}[\hat{Y}]\rho dW_Y),$$

(1)

where $\mathcal{L}_{th}$ and $\mathcal{L}_{qba}$ describe interactions with the thermal and quantum optical baths, respectively [25]. The final term, written in terms of the measurement superoperator $\mathcal{H}$ and two independent Wiener processes, $W = (W_X, W_Y)$, describes the conditioning of the state upon the measurement record $i$. We define $\vec{f} = \text{tr}(\hat{f}\rho)$ as the vector of the quadrature expectation values. Assuming Gaussian statistics, the conditional state is fully characterized by these expectation values along with the covariance matrix $V_{ij} = \text{tr}((\hat{r}_i - \bar{r}_i)(\hat{r}_j - \bar{r}_j)\rho)/2$, where $\{\cdot\}$ is the anticommutator. For our purposes, this covariance can be written in terms of a single number: $V_{ij} = V_{ij}$. The dynamics of these first and second moments are given by

$$d\vec{f} = -\frac{\Gamma_m}{2}\vec{f} dt + \sqrt{4\Gamma_{meas}}V(t) dW, \tag{2a}$$

$$\dot{V}(t) = -\Gamma_m V(t) + \Gamma_m \left(\bar{n}_{th} + \frac{1}{2}\right) -\Gamma_{qba} - 4\Gamma_{meas}V(t)^2, \tag{2b}$$

where the Wiener process is given by the measurement record $dW = i dt - \sqrt{4\Gamma_{meas}}\vec{f} dt$ such that the quantum trajectory as defined in Eq. (2b) represents a filtering of the measurement record $i$. Note that, although the expectation value evolution is stochastic (driven by the stochastic term $dW$), the conditional variance evolves deterministically, decaying to a steady-state value

$$V = \sqrt{1 + \frac{16V_{bath}\Gamma_{meas}/\Gamma_m - 1}{8\Gamma_{meas}/\Gamma_m}},$$

(3)
where $V_{\text{bath}} = \left( \eta_{\text{th}} + 1/2 + \Gamma_{\text{qba}}/\Gamma_{m} \right)$ is the total bath variance [13]. The reduction of the conditional variance [Fig. 2(a)] to its steady state happens in a characteristic collapse time, which depends inversely on the measurement rate [8]. In the regime of fast, efficient measurement $\Gamma_{\text{meas}} \gg \gamma \gg \Gamma_{m}$, which is relevant to this work, this variance approaches $V \approx 1/(2\sqrt{\eta_{\text{meas}}})$. Thus, in the limit of highly efficient measurements ($\eta_{\text{meas}} \to 1$), the measurement process is able to project the initial thermal state into a pure coherent state $(V = 1/2)$. We note that Eqs. (2a) and (2b) are formally equivalent to a Kalman filtering problem [8], with constraints on the measurement and process noises imposed by quantum mechanics. Like the state estimate of a Kalman filter, the quantum state $\rho(t)$ enshrines the most accurate possible prediction of subsequent measurement outcomes.

In Fig. 1(d), we show an example of a quantum trajectory $\bar{\mathbf{r}}(t)$, as calculated according to Eq. (2a). The region $0 < t < 100 \mu s$ indicates the collapse time of the measurement; after which, the conditional variance has decayed to its steady-state $V$. For the large probe strength used here (corresponding to $\eta_{\text{meas}} = 67\%$), the predicted conditional variance is $V = 0.61$, which is only 20% larger than the zero-point fluctuations. This corresponds to a coherent state with purity $\mathcal{P} = tr(\rho^2) = 1/(2V) = 0.82$, which is an improvement of almost two orders of magnitude as compared to the initial thermal state purity of $\mathcal{P} = 0.02$.

The extracted quantum trajectory depends on the model and the parameters chosen to describe the experiment. Successful modeling of data acquired in earlier experiments testifies to a good understanding of our system, as well as reliable methods to extract its key parameters [24]. We can gain further confidence in the validity of the state estimation by analyzing the experimental innovation function $v dt = i dt - \sqrt{4\Gamma_{\text{meas}} \bar{r}} dt$. For correct modeling, this function represents a white noise with a flat power spectral density [16]. This matches the assumption, made above, that the measurement record is driven by a Wiener process. With all system parameters estimated independently, we have calculated the innovation function and checked that, indeed, it has a flat spectrum [25].

Unlike $\bar{r}$, the conditional variance $V$ cannot be immediately obtained from the experimental data: averaging individual trajectories approximates only the unconditional variance, $V_{\text{bath}}$ in this regime [25] (see Fig. 1). To actually verify the prepared conditional state at time $t_0$, an experimenter could make a strong projective measurement at that time. Here, we approximate this by a positive-operator valued measure (POVM) measurement based on subsequent data collected for $t > t_0$ [20,21]. To do so, we backpropagate an effect operator $E$ from future times to the past time $t_0$. The role of the effect operator is to refine, in a Bayesian sense, the probabilities for measurement outcomes, as determined by a density matrix $\rho(t_0)$. Together, $\rho$ and $E$ define the past quantum state [20,21] from which the
expectation value of an operator $\hat{A}$ is calculated as
$\text{Tr}(\hat{A}\rho E)/\text{Tr}(\rho E)$. The (unconditional) thermal state could be
chosen for $\rho(t_0)$, disregarding the data collected before $t_0$. If any prior (to $t_0$)
information about the system is ignored, then $\rho(t_0) \propto 1$ and $E$ contains all
information about the quantum state at time $t_0$—as determined exclusively
from measurements at later times.

For states for which the evolution $\rho(t)$ is restricted to Gaussian states, the
effect operator $E$ can be characterized by the expectation value $\bar{r} = \text{tr}(rE)$
and the covariance matrix $(V_E)_{ij} = \text{tr}((\hat{r}_i - \bar{r}_i)(\hat{r}_j - \bar{r}_j)E)/2$.
For the mechanical measurement performed here, $V_E$ can be written as
$(V_E)_{ij} = V_E \delta_{ij}$; and the first and second moments evolve according to [10,11,20]

$$-d\bar{r}_i = \frac{\Gamma_m}{2} \bar{r}_i dt + \sqrt{4\Gamma_{\text{meas}}V_E(t)}dW_i, \quad (4a)$$

$$-V_E(t) = \frac{V_E(t-dt) - V_E(t)}{dt}$$

$$= \Gamma_m V_E(t) + \Gamma_m \left(\bar{n}_\text{th} + \frac{1}{2}\right) + \Gamma_\text{qba} - 4\Gamma_{\text{meas}}V_E(t)^2, \quad (4b)$$

where $dW_i = idt - \sqrt{4\Gamma_{\text{meas}}}d\bar{r}_i$ is a stochastic variable,
and the steady-state conditional variance is $V_E = V + \Gamma_m/(4\Gamma_{\text{meas}}) \approx V$. It is this retrodicted trajectory
(determined only by the measurement record after $t_0$) that we will use to verify individual quantum trajectories such as the one shown in Fig. 1.

In Fig. 2(a), we show predicted and retrodicted trajectories, $\bar{r}(t)$ and $\tilde{r}(t)$, in a time interval in which both conditional variances have reached the steady state. These trajectories are compared at an arbitrary common end point, $t_0$. Various pairs of $\bar{r}(t_0)$ and $\tilde{r}(t_0)$, from different experimental trials, are shown in Fig. 2. We calculate, over
this ensemble of experimental realizations, the covariance matrix of the relative trajectories $\sigma^2 = \text{Cov}[(\bar{r}(t_0) - \tilde{r}(t_0))]$.

As expected, we always find that $\sigma^2_{\tilde{X}} \approx 0$ and $\sigma^2_{\tilde{X}X} \approx \sigma^2_{\tilde{Y}Y}$
(within 2%), and henceforth report only the average diagonal term, $\sigma^2$. This experimental variance provides a direct measurement of the desired quantum conditional variance $V$. As we derive in the Supplemental Material
[25], the ensemble variance is given by a simple sum of the variances of the operators characterizing the pre- and retrodicted quantum states, respectively,

$$\sigma^2 = V + V_E \approx 2V, \quad (5)$$

corresponds to a purity of $p = 0.78$. (This includes a correction due to a 6% systematic error introduced by demodulation filter correlations [25]). This corresponds to a displaced thermal state of occupation of $\bar{n}_\text{cond} = 0.14$. In this sense, this process is sometimes referred to as “cooling by measurement” [15]. Force feedback based on the predicted quantum state can, in principle, entirely undo the displacement to yield a zero-mean low-entropy state [7,29].

We can extend this retrodiction-verification protocol to study the measurement process in more detail, including its dynamics. In particular, we can observe both the measurement-induced collapse of the conditional state and decoherence in the absence of measurement conditioning. To do so, we compare retrodicted quadrature values with the forward-calculated ones at a time $t_0$, which now varies within the interval $0 < t_0 < 3$ ms. The resulting relative trajectories $[\bar{r}(t_0) - \tilde{r}(t_0)]$ and their ensemble variance are shown in Figs. 3(a) and 3(b), respectively. Note that the retrodiction always begins at 3.2 ms such that its conditional variance is in the steady state throughout the displayed time interval. In contrast, the predictions all
For comparison, the unconditional variance \[ \sigma^2 \] calculated from Fig. 3(b). Next, to visualize the dynamics of the system in behavior revealed by the retrodictive state verification in unconditional thermal state. Indeed, this is exactly the variance \( \langle \vec{r}(t) \rangle = \vec{r}(0) \) is expected, per Eq. (2b), to decay from an unconditional thermal state. A few traces are highlighted in black for illustration purposes. (b) Relative variance \( \sigma^2(t) \) of the ensemble of relative trajectories from Fig. 3(a). For comparison, the unconditional variance [calculated from \( \vec{r}(t) \) only] is shown in blue. The theoretical expectation \( (V + V_E) \) is shown in gray, and the variance of a pure coherent state is shown in black.

begin at \( t = 0 \mu s \). Thus, up to \( \approx 50 \mu s \), the conditional variance \( V(t) \) is expected, per Eq. (2b), to decay from an unconditional thermal state. Indeed, this is exactly the behavior revealed by the retrodictive state verification in Fig. 3(b). Next, to visualize the dynamics of the system in the absence of measurement, we set the measurement efficiency to zero in Eq. (2a) from a time of 0.7 ms onward. That is, we stop conditioning the predicted quantum trajectory based on the measurement record. Thus, the conditional variance rethermalizes, and our nearly pure state decoheres into a statistical mixture.

The ability to effect strong projective displacement measurements via continuous weak measurement opens the door to various measurement-based protocols. Applying the same underlying machinery to modified measurement schemes allows, for example, the production of conditional mechanical squeezed states, as well as conditional entanglement and quantum steering [10,11,30]. Moreover, although the analysis presented here was implemented in the postprocessing of data, there is no fundamental obstacle to implementing this protocol as a real-time filter—e.g., using a field-programmable gate array (FPGA). Real-time feedback that removes the (known) mean displacement can then yield unconditional states with the desired quantum correlations. Finally, the application of an optimal quantum-limited state estimation to such ultracoherent mechanical resonators may be of interest to test and constrain models of spontaneous wave function collapse. Indeed, the resonator in this work already suggests experimental bounds for continuous spontaneous localization that are competitive with the state of the art [25,31,32], and the prospects should improve dramatically at millikelvin temperatures.

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