Large Quantum Delocalization of a Levitated Nanoparticle Using Optimal Control: Applications for Force Sensing and Entangling via Weak Forces

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We propose to optimally control the harmonic potential of a levitated nanoparticle to quantum delocalize its center-of-mass motional state to a length scale orders of magnitude larger than the quantum zero-point motion. Using a bang-bang control of the harmonic potential, including the possibility of inverting it, the initial ground-state-cooled levitated nanoparticle coherently expands to large scales and then contracts to the initial state in a time-optimal way. We show that this fast loop protocol can be used to enhance force sensing as well as to dramatically boost the entangling rate of two weakly interacting nanoparticles. We parameterize the performance of the protocol, and therefore the macroscopic quantum regime that could be explored, as a function of displacement and frequency noise in the nanoparticle's center-of-mass motion. This noise analysis accounts for the sources of decoherence relevant to current experiments.

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A levitated nanoparticle in high vacuum is a promising system to explore quantum mechanics at large scales for three reasons. (i) The center-of-mass motion of a nanoparticle of mass m in the regime of 10^8 to 10^{11} atomic mass units can be prepared in a pure quantum state via groundstate cooling in a tight harmonic potential of frequency ω_0 $({\sim}2\pi \times 100 \text{ kHz})$ [1–[7\].](#page-4-1) The corresponding zero-point motion $x_0 \equiv \sqrt{\hbar/(2m\omega_0)}$ is minute (∼10⁻¹² m). (ii) The harmonic potential can be tuned to induce dynamics in which the position probability distribution is expanded to scales larger than x_0 , ideally approaching the length scale given by the size of the nanoparticle $(\sim 10^{-7} \text{ m})$ [8–[11\]](#page-4-2). (iii) Levitation in ultra-high vacuum provides a high degree of isolation of its center-of-mass motion, enabling the induced dynamics to be coherent. In this Letter, we combine these three ingredients in an optimal way to propose feasible nanomechanical experiments [\[12\]](#page-4-3) exploring and exploiting quantum mechanics at scales orders of magnitude larger than the zero-point motion (see Fig. [1](#page-0-0)). This proposal is particularly timely given the recent experiments on cooling an optically levitated nanoparticle into the quantum regime $[5-7,13,14]$. In contrast to other significantly more challenging proposals [\[8,10,15](#page-4-2)–17], here we do not require the preparation of quantum superposition states.

More specifically, we propose to dynamically manipulate the harmonic potential of a nanoparticle, including the possibility of inverting it, to first coherently expand its state to large scales and then compress it back to its initial state in a time-optimal way. By measuring the purity of the final state (e.g., via its center-of-mass temperature) as well as the position probability distribution at the point of maximum

FIG. 1. Applications of the proposed loop protocol. (a) coherent expansion to large scales, (b) static force sensing, and (c) entangling two particles via weak forces. (d) Dimensionless variance $v_x \equiv \langle \hat{X}^2 \rangle / x_0^2$ (solid black line, left axis in logarithmic scale) and ω^2/ω_0^2 of the harmonic potential (solid blue line, right axis in linear scale) as a function of time in the optimal bang-bang loop.

expansion (on different experimental runs), one could conclude that the center-of-mass expanded coherently to scales larger than the zero-point motion [Fig. [1\(a\)](#page-0-0)]. In addition, returning to the initial contracted state is beneficial both to minimize decoherence, as the expanded state is very fragile, and to facilitate the required repetition of experimental runs. The loop protocol can be used to explore quantum physics of massive objects at large scales, thereby falsifying collapse models [\[9,18\].](#page-4-5) The loop protocol can also be used for force sensing [\[11\],](#page-4-6) as the presence of a static force critically alters the final state of the protocol [see Fig. [1\(b\)\]](#page-0-0). Furthermore, if two nanoparticles interact via a weak force (e.g., Coulomb [\[19\],](#page-4-7) Casimir [\[20\],](#page-4-8) or gravity [\[21,22\]\)](#page-4-9), the loop protocol dramatically enhances the entangling rate due to the expansion of the wave function [see Fig. $1(c)$]. In the following, we describe and analyze these ideas in detail by taking into account the effect of displacement and frequency noise in the nanoparticle, which encompass the relevant sources of decoherence in current experiments with levitated nanoparticles [\[23,24\]](#page-4-10).

Let us consider the center-of-mass motion of a nanoparticle along the x axis. The motion along the other axes as well as rotational degrees of freedom are assumed to be either decoupled or to effectively induce displacement and frequency noise. We assume the experimental possibility of dynamically controlling a harmonic potential in the form $V(X, t) = m\omega^2(t)X^2/2$ with $\omega^2(t) \in [-\omega_0^2, \omega_0^2]$. That is, the harmonic frequency that determines the strength of the spring constant is upper bounded to ω_0 , but the potential can be inverted. We are interested in the optimal control problem of transitioning between states in a time-optimal way [\[25\]](#page-5-0) in order to be faster than decoherence. With potentials of the form $V(X, t)$, the so-called bang-bang solutions are known to be time optimal [\[26,27\]](#page-5-1). Bang-bang solutions make $\omega^2(t)$ alternate, with sudden changes, between the two extreme available values $\pm \omega_0^2$. Hereafter, we thus consider the two Hamiltonians given by

$$
\hat{H}_{\pm} = \frac{\hat{P}^2}{2m} \pm \frac{1}{2} m \omega_0^2 \hat{X}^2 = \frac{\hbar \omega_0}{4} (\hat{p}^2 \pm \hat{x}^2). \tag{1}
$$

Here, $\hat{X} = \hat{x}x_0$ and $\hat{P} = \hat{p}p_0$, with $[\hat{X}, \hat{P}] = i\hbar$, are the position and momentum operators. We have defined dimensionless operators using the zero-point motion position x_0 and momentum $p_0 = \hbar/(2x_0)$. We define the thermal state of \hat{H}_+ as $\hat{\rho}_{\text{th}} = \sum_{n=0}^{\infty} P(n) |n\rangle\langle n|$, where $P(n) = [\bar{n}/(1+\bar{n})]^n (1+\bar{n})^{-1}$, |n) with $n = 0, 1, ...$ being a Fock state $[\hat{H}_+|n\rangle = \hbar\omega_0(n+1/2)|n\rangle$ and \bar{n} the phonon mean number.

The loop protocol proposed in this Letter [see Fig. [1\(d\)](#page-0-0)], is based on the following key equality

$$
e^{-i\hat{H}_{-}t_{I}/\hbar}e^{-i\hat{H}_{+}t_{H}/\hbar}e^{-i\hat{H}_{-}t_{I}/\hbar}=e^{-i\hat{H}_{+}t_{H}/\hbar},\qquad(2)
$$

which holds for $t_H\omega_0 = \pi(2l + 1)/2$ with $l = 0, 1, \dots$ and arbitrary t_I (see [\[28\]\)](#page-5-2). The left-hand side of Eq. [\(2\)](#page-1-0) is a unitary operator, labeled by \hat{U}_{loop} , that describes the time evolution of the loop protocol from $t = 0$ to $t = T \equiv 2t_I + t_H$ in three steps: (i) inverted harmonic potential \hat{H} for $0 < t < t_I$, (ii) harmonic potential \hat{H} for $t_1 < t < t_1 + t_H$, and (iii) inverted harmonic potential for $t_I + t_H < t < 2t_I + t_H$. The loop protocol starts and ends with the particle in the harmonic potential, that is, one has \hat{H}_{+} for $t < 0$ and $t > T$. Equation [\(2\)](#page-1-0) shows that an initial state $\hat{\rho}(0)$ evolves during the loop protocol into $\hat{\rho}(T) = \hat{U}_{\text{loop}}^{\dagger} \hat{\rho}(0) \hat{U}_{\text{loop}}^{\dagger} = e^{-i\hat{H}_{+}t_H/\hbar} \hat{\rho}(0) e^{-i\hat{H}_{+}t_H/\hbar}$. The state at $t = T$ is thus very similar to the one at $t = 0$. In fact, $\hat{\rho}(T) = \hat{\rho}(0)$ whenever $\hat{\rho}(0)$ is diagonal in the Fock basis (e.g., thermal state). However, the state at $t = T/2$ is dramatically different as it has expanded and squeezed due to the action of \hat{H} . To quantify the expansion, we define the coefficient $\eta = \Delta x(T/2)/\Delta x(0)$, where $\Delta x(t) \equiv [\langle \hat{x}^2(t) \rangle - \langle \hat{x}(t) \rangle^2]^{1/2}$. If the initial state is thermal $\hat{\rho}(0) = \hat{\rho}_{th}$, the expansion coefficient is given by $\eta = \exp(t_I \omega_0)$, that is, it exponentially grows with the time invested in the inverted potential [\[32\]](#page-5-3). The motional quantum state at $t = T/2$ is a highly squeezed state with large spatial extension, which is very different from a large coherent state with small spatial extent that has been realized with an ion using a bang-bang protocol where the center of the trap is displaced [\[33\]](#page-5-4).

Let us discuss how we model noise and decoherence in the loop protocol. We remark that during the protocol $0 < t < T$, there is neither active nor passive cooling acting on the nanoparticle. Hence, we consider displacement $(\nu = 1)$ and frequency $(\nu = 2)$ noise, as described by the master equation

$$
\partial_t \hat{\rho} = \frac{1}{i\hbar} [\hat{H}_{\pm}, \hat{\rho}] - \Gamma_1[\hat{x}, [\hat{x}, \hat{\rho}]] - \Gamma_2[\hat{x}^2, [\hat{x}^2, \hat{\rho}]]. \tag{3}
$$

This master equation is the result of averaging the noise [\[34\]](#page-5-5) described by the stochastic Hamiltonian \hat{H}_{\pm} + $\hat{H}_{\text{noise}}(t)$ with

$$
\hat{H}_{\text{noise}}(t) = \frac{\hbar \omega_0}{4} [2f_1(t)\hat{x} + f_2(t)\hat{x}^2].
$$
 (4)

Here, $f_{\nu}(t)$ is a stochastic variable with zero mean and power spectral density given by $S_{\nu}(\omega) \equiv$ $(2\pi)^{-1} \int_{-\infty}^{\infty} d\tau \langle f_{\nu}(t) f_{\nu}(t+\tau) \rangle e^{i\omega \tau}$ (here, $\langle \cdot \rangle$ is an ensemble average over the stochastic variable). In Eq. [\(3\)](#page-1-1), the decoherence rates are then given by $\Gamma_{\nu} \equiv \pi \omega_0^2 S_{\nu} (\nu \omega_0) / 4^{\nu}$. Equation [\(3\)](#page-1-1) can also be derived by tracing out an Ohmic bath modeled as a set of quantum harmonic oscillators in the high-temperature limit [\[29\]](#page-5-6) (see [\[28\]](#page-5-2)). Standard sources of decoherence in experiments with levitated nanoparticles (e.g., laser recoil heating, blackbody radiation, vibrations, intensity field fluctuations) can be modeled using Eq. [\(3\)](#page-1-1) with the corresponding contribution to the decoherence rate Γ_{ν} [\[23,24\]](#page-4-10). In addition, note that fluctuations in the switching times are already taken into account as they are equivalent to frequency noise. This is explicitly shown in [\[28\].](#page-5-2) Decoherence due to scattering of background gas particles [\[9,35\]](#page-4-5) can be neglected whenever $\gamma_{gas}T \ll 1$, where $\gamma_{\text{gas}} = 16\pi\sqrt{2\pi}PR^2/[\sqrt{3}m_{\text{gas}}\bar{v}_{\text{gas}}]$ is the gas scattering rate with R the nanoparticle's radius, P the gas pressure, and m_{gas} and \bar{v}_{gas} the averaged mass and thermal velocity of a gas particle, respectively. It is one of the key aspects of our proposal that the loop protocol is fast enough to easily guarantee $\gamma_{\text{gas}} T \ll 1$ at ultra-high vacuum $P \sim 10^{-9}$ mbar [\[36\]](#page-5-7), that is, that the probability for a single scattering event in each experimental run is negligible. The dynamics induced by Eq. [\(3\)](#page-1-1) lead to closed equations of motion for the first and second moments [\[28\]](#page-5-2) that can thus be easily solved. Note, however, that the dissipator modeling frequency noise generates mixed non-Gaussian states. Hereafter, we use expressions that are valid only for Gaussian states. In the presence of frequency noise, we have numerically checked that they are an excellent approximation in the relevant parameter regimes. Nonharmonic terms in the potential might become relevant for large expansions. Experiments with optically levitated nanoparticles [\[37\]](#page-5-8) show that these nonlinear contributions are not significant for expansions of up to $\eta \sim 10^4$, where the spatial extent of the nanoparticle's center-ofmass position is of around 10 nm. Furthermore, one could consider shallower traps to prevent nonlinearities provided the protocol is short enough such that $\gamma_{\text{gas}}T \ll 1$ and hence decoherence due to gas scattering is kept irrelevant.

In Fig. [2](#page-2-0), we show the purity $P = \text{tr}[\hat{\rho}^2(T)]$ of the state after the loop protocol, where $\hat{\rho}(T)$ is calculated using Eq. [\(3\)](#page-1-1) with $\hat{\rho}(0) = |0\rangle\langle0|$. In Fig. [2\(a\)](#page-2-0) [Fig. [2\(b\)\]](#page-2-0), we consider the impact of displacement (frequency) noise only. A coherent expansion of $\eta = 10^2$, namely a quantum delocalization over a length scale $100 \times x_0$ requires $\Gamma_1/\omega_0 \approx 10^{-7}$ and $\Gamma_2/\omega_0 \approx 10^{-11}$, that is, a displacement noise $x_0\sqrt{S_1} \sim 10^{-18}$ m/ \sqrt{Hz} and a frequency noise of $\sqrt{S_2} \sim 10^{-8} / \sqrt{Hz}$ $\sqrt{S_2} \sim 10^{-8} / \sqrt{Hz}$ $\sqrt{S_2} \sim 10^{-8} / \sqrt{Hz}$. Figure 2 quantifies the experimental challenge (decoherence rate levels) required to explore macroscopic quantum physics. It might be convenient in order to reduce decoherence to consider protocols where the harmonic frequency in \hat{H} = is smaller than in \hat{H}_+ . While this will make the protocol longer in time (larger T), it might be convenient if decoherence due to gas scattering is still negligible $\gamma_{\text{gas}}T \ll 1$. These modifications can be analyzed and optimized for a given experimental scenario in a systematic way using the results presented here. Figure [2](#page-2-0) illustrates the fragility of macroscopic quantum physics. It quantifies how decoherence modeled by displacement and frequency noise limits the scale of quantum delocalization that can be achieved with a nanoparticle. In this regard, since collapse models effectively induce displacement noise [\[9,18\]](#page-4-5), the loop protocol can also be directly used to falsify them by measuring a high state purity.

FIG. 2. Coherent expansion. The purity at the end of the protocol as a function of the total protocol time T and the decoherence rate (a) Γ_1 [(b) Γ_2]. The lower panel in (a) relates the decoherence rate Γ_1 to the displacement noise spectrum S_1 for $\omega_0 = 2\pi \times 10^5$ Hz and several typical particle masses m. The lower panel in (b) relates the decoherence rate Γ_2 to the frequency noise spectrum $S₂$ for two different frequencies (there is no mass dependence for frequency noise). As the initial state, we consider the ground state $\bar{n} = 0$ with $\mathcal{P} = 1$. We considered a silica nanoparticle of $R = 100$ nm and mass density $\rho = 2201 \text{ kg/m}^{-3}$.

Let us now show that the loop protocol also enhances static force sensing. Similarly to the experiment done with free falling nanoparticles in [\[11\],](#page-4-6) we consider the loop protocol in the presence of a static force F . That is, the Hamiltonians H_{\pm} in the protocol are modified to

$$
\hat{H}_{\pm}^{f} = \hat{H}_{\pm} + F\hat{X} = \hat{H}_{\pm} + \hbar \omega_0 f \hat{x}.
$$
 (5)

Here $f = Fx_0/(\hbar\omega_0)$ is the dimensionless static force to be detected. The state after the loop protocol will now depend on f, that is, $\hat{\rho}(T, f)$. Using standard methods in quantum metrology, we consider the quantum Fisher information [\[38\]](#page-5-9) of $\hat{\rho}(T, f)$, denoted as I, and the Cramér-Rao bound to calculate the minimal force $f_{\text{min}} \equiv 1/\sqrt{\mathcal{I}}$ that can be detected with the loop protocol. For a Gaussian state, I can be calculated from first and second order moments [\[30\]](#page-5-10) (see [\[28\]](#page-5-2)). In Fig. [3,](#page-3-0) we display f_{min} as a function of T. The solid, dashed, and dotted-dashed black lines (the same in both panels) are f_{min} in the absence of decoherence $(\Gamma_1 = \Gamma_2 = 0)$ for the loop protocol, free evolution $(\omega_0 = 0)$, and evolution in a constant inverted potential, respectively. These lines show the enhanced force sensing, as compared to free dynamics [\[11\],](#page-4-6) provided by the fast and

FIG. 3. Force sensing. Minimal detectable force as a function of the total time T. Results are shown in the complete absence of decoherence (black lines) and in color for various values of (a) Γ_1 with $\Gamma_2 = 0$ [(b) Γ_2 with $\Gamma_1 = 0$]. Solid lines show the performance of the loop protocol. Dashed (dash-dotted) lines show the performance of a particle that is evolving freely (in an inverted potential) for a time T . The right (top) axis shows the values in Newton (microseconds) for a silica nanoparticle of $R = 100$ nm and $\omega_0 = 2\pi \times 10^5$ Hz.

large expansion induced by an inverted potential. In [\[28\]](#page-5-2), we provide the analytical expressions of f_{min} in these three cases as well as the modified key equality [Eq. [\(2\)\]](#page-1-0) of the loop protocol in the presence of a static force. The colored lines in Fig. [3\(a\)](#page-3-0) [Fig. [3\(b\)](#page-3-0)] show f_{min} in the presence of displacement noise (frequency noise). Interestingly, displacement noise saturates f_{min} as a function of time. In contrast, frequency noise provides an optimal time where f_{min} is minimal. In both Figs. [2](#page-2-0) and [3](#page-3-0), one can see that, as expected from the form of the dissipators in Eq. [\(3\)](#page-1-1), frequency noise becomes dominant whenever $\Gamma_2 \eta^4 \gtrsim \Gamma_1 \eta^2$, that is, as soon as the nanoparticle expands to regimes $\eta \gtrsim \sqrt{\Gamma_1/\Gamma_2}$.

Let us also show how the loop protocol can enhance the entanglement generation via a weak static interaction of two nanoparticles. We consider two particles, one with position vector $(\hat{X}_1, 0, 0)$ and the other with $(\hat{X}_2, d, 0)$, interacting via the general central potential

$$
\hat{V}_{\text{int}} = \frac{b}{[(\hat{X}_1 - \hat{X}_2)^2 + d^2]^{a/2}}.
$$
\n(6)

We therefore assume that the motion of the two particles in their corresponding trapping potential, as described by the position $X_{1(2)}$ and momentum $P_{1(2)}$ for particle 1 (2), is parallel and separated by a constant distance d . The dimensional real parameter b and the dimensionless integer a determine the type of interaction (e.g., Coulomb [\[19\],](#page-4-7) Casimir [\[20\]](#page-4-8), gravitational [\[21,22\]](#page-4-9)). By assuming $d^2 \gg \langle (\hat{X}_1 - \hat{X}_2)^2 \rangle$, one can Taylor expand Eq. [\(6\)](#page-3-1) to get, together with Eq. [\(1\)](#page-1-2), the total quadratic Hamiltonian

$$
\frac{\hat{H}_{\pm}^{\text{two}}}{\hbar \tilde{\omega}_0} = \frac{1}{4} \sum_{j=1,2} (\hat{p}_j^2 \pm \hat{x}_j^2) + \frac{g}{\tilde{\omega}_0} \hat{x}_1 \hat{x}_2.
$$
 (7)

Here $\tilde{\omega}_0^2 \equiv \omega_0^2 + ba/(d^{a+2}m)$ is the shifted harmonic oscillator and $g \equiv ba/(d^{a+2}2m\tilde{\omega}_0)$ the coupling rate. The dimensionless position and momentum operators are defined with the zero-point motion that depends on the shifted frequency $\tilde{\omega}_0$. From Eq. [\(7\)](#page-3-2), one can realize that the coupling rate between nanoparticles in a center-of-mass state that is expanded by a factor η , as it is achieved in the loop protocol, will be enhanced to $\eta^2 g$. In the loop protocol, the state is in the expanded phase during an amount of time t_H . To generate entanglement one requires $\eta^2 gt_H > 1$, a condition that profits from the η^2 enhancement. To confirm this reasoning, let us assume that the initial state is the product state $\hat{\rho}(0) = |0\rangle_1\langle 0| \otimes |0\rangle_2\langle 0|$ and that the evolution during the loop protocol is coherent. In Fig. [4\(a\),](#page-3-3) we

FIG. 4. Entanglement via weak forces. (a) The purity at the end of the protocol as a function of the protocol time T and coupling strength g. In the absence of decoherence, the decrease of purity signals the creation of entanglement (starting from the ground state $\bar{n} = 0$ with initial $\mathcal{P} = 1$). (b) Coupling g as a function of the particle separation d/R for Coulomb, Casimir, and gravitational interaction for a silica nanoparticle of $R = 100$ nm and $\tilde{\omega}_0 = 2\pi \times 10^5$ Hz. In the case of Coulomb interaction, each particle is assumed to carry a single electron charge. See [\[28\]](#page-5-2) for further details.

show the purity $\mathcal{P} = \text{tr}[\hat{\rho}_1^2(T)]$ of the reduced state of particle 1, namely $\hat{\rho}_1(T) = \text{tr}_2[\hat{\rho}(T)]$ (here, $\text{tr}_2[\cdot]$ is the partial trace over particle 2). Since the initial state is a product of pure states and the evolution is coherent, the reduced purity at the end of the loop protocol is a measure of the entanglement generated between the two particles. As an example, if the loop can be done coherently up to $\eta = 10^3$ (see Fig. [2](#page-2-0) for the required decoherence rates), entanglement could be generated with a weak coupling rate of $g \approx 2\pi \times 0.1$ Hz in the $T \approx 25 \mu s$ that the loop protocol requires. As shown in Fig. [4\(b\),](#page-3-3) this coupling rate corresponds to the Coulomb interaction of two nanoparticles of $R = 100$ nm, each having a single charge, separated by a distance $d \approx 3.2 \mu$ m. Entangling two nanoparticles interacting with such a week coupling rate would be a proof-ofprinciple demonstration of the enhancement induced by large expansions. Entangling via Casimir interaction [\[20\]](#page-4-8) would offer a novel approach to measure these weak forces. Entangling via gravity would have fundamental implications regarding the quantum character of gravity [\[39\]](#page-5-11). However, as shown in Fig. [4\(b\)](#page-3-3), the gravitational interaction leads to a coupling rate of the order of $g/\tilde{\omega}_0 \sim 10^{-23}$, which shows that entangling via gravitational force is a formidable task.

In summary, we have proposed a loop protocol to delocalize the center-of-mass state of a nanoparticle over large distances and compress it back to the initial localized state. The state in the expanded phase is very fragile to its environment. The loop protocol can thus be used to detect the environment, either its noisy signals, the presence of a static weak force, or the coherent interaction with another nanoparticle. We have presented the results in a way that one can easily extract the required levels of decoherence and noise in order to explore macroscopic quantum physics. From the current state of the art in experiments, where ground-state cooling of nanoparticles has been recently achieved, our results provide a quantitative path to progressively venture into the exciting regime of macroscopic quantum physics.

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Note added.—As the authors were submitting this manuscript, the preprint arXiv:2012.07815 [\[40\]](#page-5-12) discussing related ideas was posted.

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Correction: Missing information in Ref. [40] has been inserted.