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Probing macroscopic quantum superpositions with nanorotors

Benjamin A Stickler¹, Birthe Papendell¹, Stefan Kuhn², Björn Schrinski¹, James Millen²,³, Markus Arndt¹ and Klaus Hornberger¹,⁴

¹ University of Duisburg-Essen, Faculty of Physics, Lotharstraße 1, D-47048 Duisburg, Germany
² University of Vienna, Faculty of Physics, VCQ, Boltzmannsgasse 5, A-1090, Vienna, Austria
³ King’s College London, Department of Physics, Strand, WC2R 2LS, London, United Kingdom
⁴ Author to whom any correspondence should be addressed.

E-mail: klaus.hornberger@uni-due.de

Abstract

Whether quantum physics is universally valid is an open question with far-reaching implications. Intense research is therefore invested into testing the quantum superposition principle with ever heavier and more complex objects. Here we propose a radically new, experimentally viable route towards studies at the quantum-to-classical borderline by probing the orientational quantum revivals of a nanoscale rigid rotor. The proposed interference experiment tests a macroscopic superposition of all possible orientations. It requires no diffraction grating, uses only a single levitated particle, and works with moderate motional temperatures under realistic environmental conditions. The first exploitation of quantum rotations of a massive object opens the door to new tests of quantum physics with submicron particles and to quantum gyroscopic torque sensors, holding the potential to improve state-of-the-art devices by many orders of magnitude.

1. Introduction

Various experiments have been carried out to test the validity of the quantum superposition principle [1] for ever heavier and more complex objects [2–10]. Such experiments aim at probing modifications of linear quantum physics, have far-reaching technological applications, and may eventually reveal quantum aspects of gravity [11, 12]. Most proposals for future experiments suggest observing the superposition of different motional states of micromechanical oscillators [13, 14] or the center-of-mass interference of free-flying nanoparticles [15–20]. Here we propose an entirely new way of testing the quantum superposition principle by exploiting the quantized orientational degree-of-freedom of a rigid object.

The proposed interference scheme is based on the fact that an initially tightly oriented quantum rotor rapidly disperses, while at multiples of a much longer quantum revival time the collective interference of all occupied angular momentum states leads to a complete re-appearance of the initial state [21, 22]. Surprisingly, we find that such orientational quantum revivals can be probed with modest initial temperatures, involving hundreds of thousands of total angular momentum quanta, and under realistic environmental conditions, taking into account all relevant sources of orientational decoherence [23, 24]. The proposed experiment enables the first test of macroscopic angular momentum quantization with massive objects. It requires no diffraction grating and uses only a single recyclable nanoparticle. The empirical measure of macroscopicity [25] (quantifying to which extent a superposition test falsifies a wide class of classicalizing modifications of quantum physics) is comparable to that of ambitious center-of-mass proposals.

We argue that macroscopic orientational quantum revivals can be observed using existing technology for optically manipulating the motion and alignment of levitated particles [26–29]. The proposed scheme requires cavity- or feedback-cooling of the nanoparticle rotation [30, 31] to below a Kelvin, while it is independent of its center-of-mass temperature. An observation of orientational quantum revivals with nanorotors would substantially advance macroscopic superposition tests, provide the first experimental test of the angular...
momentum quantization of massive objects, and enable quantum coherent gyroscopic torque sensing with the potential of improving state-of-the-art devices [29, 32] by many orders of magnitude.

2. Orientational quantum revival scheme

The proposed scheme consists of cycling the four consecutive steps displayed in figure 1: (a) alignment, (b) dispersion, (c) revival, and (d) recapture. To discuss each step in detail, we consider a nanoscale linear rigid rotor of length ℓ, mass M, and moment of inertia $I = ML^2/12$ levitated in high vacuum by an optical tweezer, consisting of two counter-propagating beams of power P and linear polarization direction $\varepsilon$, which form a standing wave of waist $w$. The tweezer is aligned with the gravitational field so that the released particle drops along the tweezer axis. Denoting the angle between the rotor symmetry axis and $\varepsilon$ by $\beta$, the optical potential of the nanoparticle orientation at the antinode is given by $V(\beta) = -V_0 \cos^2 \beta$. The potential depth is $V_0 = 4\Delta \alpha \ell / \pi c e \omega_0 w^2$ with the polarizability anisotropy $\Delta \alpha = \alpha_{||} - \alpha_\perp$.

2.1. Alignment

The orientation of the nanoparticle can be cavity or feedback cooled [30, 31] leading to a tight alignment of the rotor with the field polarization direction. To discuss each step in detail, consider a nanoscale linear rigid rotor of length $\ell$, mass $M$, and moment of inertia $I = ML^2/12$ levitated in high vacuum by an optical tweezer, consisting of two counter-propagating beams of power $P$ and linear polarization direction $\varepsilon$, which form a standing wave of waist $w$. The tweezer is aligned with the gravitational field so that the released particle drops along the tweezer axis. Denoting the angle between the rotor symmetry axis and $\varepsilon$ by $\beta$, the optical potential of the nanoparticle orientation at the antinode is given by $V(\beta) = -V_0 \cos^2 \beta$. The potential depth is $V_0 = 4\Delta \alpha \ell / \pi c e \omega_0 w^2$ with the polarizability anisotropy $\Delta \alpha = \alpha_{||} - \alpha_\perp$.

The quantum state of the rotor is then characterized by its librational temperature $T$ and the angular momentum quantum number can be approximated as (see appendix A)

$$\langle jm | \rho_0 | jm' \rangle \approx \frac{\delta_{mm'}}{Z} \frac{\ell_{ij-j'j'/2}}{2k_B T} \left[ \frac{V_0}{2k_B T} \left( 1 - \frac{4m^2}{(j + j' + 1)^2} \right) \right] \exp \left[ -\hbar^2 (j + j' + 1)^2 / 8k_B T \right]$$

for $j = j'$ even and $\langle jm | \rho_0 | jm' \rangle = 0$ otherwise. Here, $I_0(\cdot)$ denotes the modified Bessel function. The expectation value of the total angular momentum quantum number can be approximated as (see appendix B)

$$\langle j \rangle \approx \sqrt{\pi k_B T / 2 \hbar^2}$$

yielding a mean occupation of $\langle j \rangle \approx 2.6 \times 10^4$ for $\ell = 50$ nm, $M = 10^6$ amu, and $T = 1$ K.

Figure 1. Scheme to observe orientational quantum revivals of a nanoscale rotor. (a) The rotor is levitated in an optical tweezer formed by two counter-propagating linearly polarized laser beams. Cavity or feedback cooling to subkelvin temperatures tightly aligns it with the field polarization (indicated by the small arrows). (b) After switching off the trapping and cooling beam the quantum state of the rotor quickly disperses into a superposition of all possible orientations, while its center-of-mass drops in the gravitational field. (c) At integer multiples of the revival time $T_{rev}$, the initial state is recovered by constructive interference of all occupied rotation states. This orientational quantum revival is detected by the total light scattered from a plane-wave probe pulse, which collapses the rotor into a state of definite orientation. (d) To repeat this quantum measurement several times, the rotor is recaptured by the trapping lasers and then transferred back to step (a) by tuning their relative phase.
The initial orientational alignment can be quantified by the expectation value \( \langle \cos^2 \beta \rangle_0 = \text{tr}(\rho_0 \cos^2 \hat{\beta}) \), from now on referred to as the alignment. It is unity for a perfectly aligned particle and 1/3 for uniformly distributed orientations. For the initial state equation \( \hat{\rho} \) the alignment can be determined in leading order of \( k_\text{B} T / V_0 \) as (see appendix B)

\[
\langle \cos^2 \beta \rangle_0 = k_\text{B} T \frac{\partial}{\partial V_0} \ln Z \approx 1 - \frac{k_\text{B} T}{V_0}.
\]

(2)

This relation holds if many angular momentum quanta are occupied (but fails in the deep quantum regime where the alignment is limited by the uncertainty relation).

2.2. Rotation dynamics during free fall

Once the trapping laser is turned off, the orientation state evolves freely while its center-of-mass drops in the gravitational field along the tweezer axis. The ensuing delocalization of the orientation state is counteracted by orientational decoherence processes [23, 24] which potentially suppress the revivals. As in other matter-wave experiments [15, 17, 20, 34, 35], the dominant sources of environmental decoherence are the scattering of residual gas atoms and the thermal emission of photons.

Exactly solving the Markovian quantum master equation of orientational decoherence [23, 24] is challenging due to the vast number of occupied angular momentum states. As a conservative estimate, we assume that a single decoherence event suffices to completely destroy the alignment signal, by producing a state \( \rho \) with \( \langle \hat{\Omega} | \rho | \hat{\Omega} \rangle = 1/4\pi \) where \( \hat{\Omega} \) is the rotor orientation. The rotor dynamics can then be described by the master equation \( \partial_t \rho = -i[H_0, \rho] / \hbar + \Gamma (\rho - \rho) \), where \( \Gamma \) is the total rate of decoherence events. The alignment \( \langle \cos^2 \beta \rangle = \text{tr}[\rho(t) \cos^2 \hat{\beta}] \) at time \( t \) follows as

\[
\langle \cos^2 \beta \rangle = \langle \cos^2 \beta \rangle_0 e^{-\Gamma t} + \frac{1}{3} (1 - e^{-\Gamma t}),
\]

(3)

where \( \langle \cos^2 \beta \rangle_0 = \text{tr}[\rho_0(t) \cos^2 \hat{\beta}] \) denotes the alignment dynamics of the decoherence-free evolution (see appendix C), and

\[
\langle jm|\rho(t)\rangle \langle j' m' \rangle = \sum_{j, j'} \sum_{m = -j}^j \sum_{m' = -j'} \langle jm|\rho_0\rangle \langle j' m' \rangle \exp \left[ \frac{i\hbar}{2\Gamma} (j(j+1) - j'^{(j'+1)}) \right].
\]

(4)

The initially trapped orientation state quickly disperses during free fall due to its angular momentum spread, see figure 2. This rapid alignment decay can be approximated using the shearing dynamics associated with a flat orientation space (see appendix D)

\[
\langle \cos^2 \beta \rangle_u \approx \langle \cos^2 \beta \rangle_0 e^{-\kappa^2 \tau^2} + \frac{1}{2} (1 - e^{-\kappa^2 \tau^2}),
\]

(5)

with rate \( \kappa = \sqrt{2k_\text{B} T / I} \).

The corresponding classical dynamics exhibits the same alignment decay since \( \rho_0 \) is virtually indistinguishable from a classical thermal state for the considered temperatures. After this initial alignment reduction to a value of 1/2, the classically expected alignment shows no revivals at all. Rather, it decays as 1/3 + e^{-\Gamma t}/6, based on the same assumptions that lead to equation (3).

The alignment of the quantized rotor follows this classical prediction for most of the time. However, the initial orientation state equation (2) recurs at integer multiples of the revival time \( T_{\text{rev}} = 2\pi I / \hbar \), as follows directly from equation (4) and as displayed in figure 2. The width and height of the orientational quantum revival are determined by the initial librational temperature as described by equations (2) and (5). In general, quantum revivals can occur when the energy depends quadratically on the quantum number [36], such as for the quantum particle in a box or for photons in a Kerr nonlinear medium [37, 38].

2.3. Measuring the alignment

The instantaneous alignment at variable times can be measured by illuminating the rotor with a weak plane-wave probe pulse of nanosecond duration and collecting the scattered light, as demonstrated in [27]. Using a plane-wave laser ensures that light scattering is independent of the rotor center-of-mass position. Choosing the pulse polarization in the same direction as that of the trapping laser, the total light scattered from the probe laser pulse is proportional to the instantaneous alignment \( \langle \cos^2 \beta \rangle \) [27, 30]. The scattering of probe photons decoheres the rotor state to a definite orientation, implementing a projective quantum measurement of \( \cos^2 \hat{\beta} \). By probing the particle orientation after variable times \( t \) in repeated experimental runs one can thus test for the emergence of orientational revivals. Note that the optical torque exerted by the probe laser is irrelevant since the rotor is recaptured and recycled in the next step.
2.4. Recapture

In the final step of the scheme the rotor is recaptured by switching on the trapping laser when the particle traverses an antinode. Moderate laser powers of a few tens of Watts suffice for a nanoparticle of length $\ell = 50$ nm and mass $M = 10^6$ amu. For such particles the revival time is as short as $T_{\text{rev}} \approx 21$ ms so that the rotor drops only 2.1 mm and reaches the center-of-mass velocity $0.2$ m s$^{-1}$. For longer revival times, it can be beneficial to use the probe pulse for recapturing the rotor.

Note that the current proposal requires no diffraction grating, the position from which the rotor is released does not affect the alignment signal, and the particle can be recycled during the experiment. These advantages substantially reduce the requirements on nanoparticle fabrication and on source stability as compared to center-of-mass superposition tests.

3. Discussion

3.1. Carbon nanotubes (CNTs) and silicon nanorods (SNRs)

To demonstrate the viability of the proposed scheme, we discuss the experimental realization for two types of nanoscale rotors, semiconducting double-walled CNTs and SNRs. Both can be fabricated with a length of $\ell = 50$ nm and a mass $M = 10^6$ amu. For such particles the revival time is as short as $T_{\text{rev}} \approx 21$ ms so that the rotor drops only 2.1 mm and reaches the center-of-mass velocity $0.2$ m s$^{-1}$. For longer revival times, it can be beneficial to use the probe pulse for recapturing the rotor.

Note that the current proposal requires no diffraction grating, the position from which the rotor is released does not affect the alignment signal, and the particle can be recycled during the experiment. These advantages substantially reduce the requirements on nanoparticle fabrication and on source stability as compared to center-of-mass superposition tests.

![Figure 2](image_url)

**Figure 2.** (a) Orientational alignment signal $\langle \cos^2 \beta \rangle$ of carbon nanotubes ($M = 1.9 \times 10^5$ amu, $\ell = 50$ nm) as a function of time (blue solid line) for $T = 100 \mu$K. For most of the time, the alignment shows the classical behavior (dashed red line) of decaying exponentially with rate $\Gamma$ from $1/2$ towards $1/3$. However, the initial alignment recurs at integer multiples of the quantum revival time $T_{\text{rev}} = 2\pi \hbar / \ell \approx 3.8$ ms and approaches a minimum at half integer multiples of $T_{\text{rev}}$. (b) Initial alignment decay, half-revival and revival for three different initial temperatures. The width of the revival peaks is determined by the decay time of the initial state, $1/\kappa = \sqrt{T_{\text{rev}}T}$. For $T = 100 \mu$K and $T = 1$ mK the initially trapped state is numerically calculated by exact diagonalization of $H$. For SNRs it has been demonstrated in [17] that internal heating and photon emission is negligible for a wavelength of $1.55$ $\mu$m. For CNTs, where excitonic excitations play no role for wavelengths well above $2.5$ $\mu$m [40], the exact
position and width of the vibrational excitations depends on the structural details of the particle [41] and the optimal trapping wavelength can be determined experimentally.

Cavity or feedback cooling the rotation of the trapped particles to subkelvin temperatures is feasible [30, 31], but may require low mode volume cavities to enhance the nanoparticle-light interaction and detection efficiency. The deeply trapped particle librates harmonically, so that well established techniques of center-of-mass optical cooling can be adapted [42–45].

The total decoherence rate $\Gamma$ accounts for collisions with residual gas atoms and for the emission of thermal photons, $\Gamma = \Gamma_{\text{gas}} + \Gamma_{\text{em}}$. The rate at which thermal gas atoms of mass $m_g$, pressure $p_g$, and temperature $T_g$ scatter off a cylinder of length $\ell$ and effective diameter $d_{\text{eff}}$ can be estimated by integrating the mean particle flux into the surface over the particle shape

$$\Gamma_{\text{gas}} = \frac{\pi p_g d_{\text{eff}} \ell}{\sqrt{2\pi m_g k_B T_g}} \left(1 + \frac{d_{\text{eff}}}{2\ell}\right).$$

The rate of thermally emitted photons depends on the internal temperature and the material-specific spectral absorption cross section of the nanoparticle [17, 46]. The former is determined by the internal heating of the particle during the recycling and alignment step. The effect of heating can be minimized by choosing the infrared wavelength of the trapping laser between vibrational transitions, where the particle is practically transparent.

Figure 2 shows the expected alignment for CNTs as a function of the time delay between nanoparticle release and detection, as numerically calculated by propagating the initial state. The latter is determined by exact quantum revivals that contradict the predicted loss of orientational coherence [32, 47].

3.2. Macroscopicity

In order to assess the proposed superposition test we consider the empirical measure of macroscopicity $\mu$ as defined in [25]. It quantifies to what extent a successful superposition experiment serves to rule out a wide class of classicalizing modifications of quantum theory. The resulting dynamics can be solved exactly for planar rotations (see appendix F), which provides a lower bound for the macroscopicity of the $n$th linear rotor quantum revival

$$\mu \geq \log_{10} \left( \frac{n \theta_m}{\ln f} \left( \frac{M}{m_e} \right)^2 \frac{T_{\text{rev}}}{1 \text{ s}} \right),$$

where $f$ is the ratio of observed-to-expected signal visibility and $\theta_m \simeq 0.12$ is a numerical factor. Assuming $M = 10^6$ amu, $\ell' = 50$ nm, and $f = 0.8$ at the tenth revival yields a lower bound of $\mu \geq 17.5$, on a par with ambitious center-of-mass interference proposals. For comparison, a tobacco mosaic virus [47] with $\ell' \simeq 300$ nm, $d = 20$ nm, and $M \simeq 4 \times 10^7$ amu has $T_{\text{rev}} \simeq 30$ s, and observation of its first revival ($f = 0.8$) would imply $\mu \geq 22.8$.

3.3. Sensing applications

The presence of an external torque during free fall can have a strong influence on the orientational revival signal. By monitoring the alignment as a function of time for different initial orientations the magnitude and direction of an applied torque can thus be deduced. The torque sensitivity of the tenth revival of a CNT ($T = 100 \mu$K) is illustrated in figure 3, showing the alignment reduction due to an external torque of magnitude $N_{\text{ext}}$ orthogonal to the trapping laser polarization. The numerical simulations of the torque-induced dynamics, involving transitions between 360 000 angular momentum states (see Materials and Methods), show that torques on the order of $10^{-30}$ Nm are observable, eight orders of magnitude smaller than the levitated [28, 29] or solid-state-integrated [32] setups considered so far.

By attaching single elementary charges to the ends of the SNR discussed above one can measure electrostatic fields at values well below mV m$^{-1}$. The decay of the alignment signal when the rotor is exposed to an atomic beam or other controlled environments can be used for studying collisional decoherence and thermalization of quantum nanoscale rotors. Finally, objective collapse models could be tested by observing orientational quantum revivals that contradict the predicted loss of orientational coherence [48].
4. Conclusion

We presented a viable scheme for the first observation of orientational quantum revivals of nanoscale particles. The proposed experiment can be realized with upcoming technology, representing a macroscopic test of the superposition principle and opening the door to quantum enhanced torque sensing. The successful demonstration of orientational revivals may well be the starting point for interferometric manipulation methods of nanoscale rigid rotors, based on applying a sequence of optical potentials during the free evolution.

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Appendix A. Semiclassical evaluation of matrix elements

For completeness, we briefly summarize how to approximate quantum mechanical matrix elements of the linear rotor using Bohr–Sommerfeld quantization. The angles \( \alpha, \beta \) and their canonical angular momenta \( p_{\alpha}, p_{\beta} \) are related to the action-angle variables \( \alpha_m, \alpha_j, m, j \) via

\[
\begin{align*}
\alpha &= \alpha_m + \arctan(\zeta \tan \alpha_j) - \pi, \\
\cos \beta &= \cos \alpha_j \sqrt{1 - \zeta^2}, \\
\sin \beta &= \sqrt{\sin^2 \alpha_j + \zeta^2 \cos^2 \alpha_j}, \\
p_{\beta} &= \hbar \left( j + \frac{1}{2} \right) \frac{\sin \alpha_j \sqrt{1 - \zeta^2}}{\sqrt{\sin^2 \alpha_j + \zeta^2 \cos^2 \alpha_j}}, \\
\end{align*}
\]

with \( \zeta = m/(j + 1/2) \). Note that these relations imply \( p_{\beta}^2 / \sin^2 \beta + p_{\beta}^4 = \hbar^2 (j + 1/2)^2 \).

The matrix elements \( \langle jm|A|j'm'\rangle \) of an arbitrary operator \( A = A(\hat{\alpha}, \hat{\beta}, p_{\alpha}, p_{\beta}) \) can be semiclassically approximated by first replacing all arguments of \( A \) according to equation (A.1), yielding \( \tilde{A}(\alpha_m, \alpha_j, m, j) \). The matrix elements can then be obtained by calculating

\[
\langle jm|A|j'm'\rangle \approx \frac{1}{(2\pi \hbar)^2} \int_0^{2\pi} d\alpha_m \int_0^{2\pi} d\alpha_j e^{i\alpha_m(j-m')} e^{i\alpha_j(j'-j)} \tilde{A}(\alpha_m, \alpha_j, m + m', j + j'/2),
\]

Applying this to \( \rho = \exp(-H/k_B T)/Z \) yields equation (1).
Appendix B. Initial alignment

In order to estimate the initial alignment of a rotor in the potential \( V(\beta) = -V_0 \cos^2 \beta \) we calculate the expectation value \( \langle \cos^2 \beta \rangle_0 = \text{tr}(\rho_0 \cos^2 \beta) \). Inserting the initial state \( \rho_0 = \exp(-\mathbf{H}/k_B T)/Z \) with \( \mathbf{H} = \mathbf{J}^2/2I + V(\beta) \) shows that the expectation value can be expressed in terms of the partition function \( Z \),

\[
\langle \cos^2 \beta \rangle_0 = k_B T \frac{\partial}{\partial V_0} \ln Z. \tag{B.1}
\]

The latter can be calculated explicitly in the semiclassical limit, where the matrix elements of \( \rho_0 \) take the form equation (1).

\[
Z = \sum_{j=0}^{\infty} \sum_{m=-j}^{j} \exp \left[ -\frac{\hbar^2 (j + 1/2)^2}{2I k_B T} \right] I_0 \left[ \frac{V_0}{2k_B T} \left( 1 - \frac{m^2}{(j + 1/2)^2} \right) \right] \times \exp \left[ \frac{V_0}{2k_B T} \left( 1 - \frac{m^2}{(j + 1/2)^2} \right) \right]. \tag{B.2}
\]

One then replaces the sum over \( m \) by an integral from \(- (j + 1/2)\) to \(+ (j + 1/2)\) and the sum over \( j \) by an integral over \( j + 1/2 \). After the substitution \( u = m/(j + 1/2) \) we thus obtain

\[
Z \simeq \frac{k_B T}{\hbar^2} \int_{-1}^{1} du I_0 \left[ \frac{V_0}{2k_B T} (1 - u^2) \right] \exp \left[ \frac{V_0}{2k_B T} (1 - u^2) \right]. \tag{B.3}
\]

The integral can be evaluated for \( k_B T/V_0 \ll 1 \) by using the asymptotic expansion \( I_0(z) \sim e^z/\sqrt{2\pi z} \) as \( z \to \infty \),

\[
Z \simeq \frac{1}{k_B T} \exp \left( \frac{V_0}{k_B T} \right). \tag{B.4}
\]

Inserting this into equation (B.1) gives equation (2).

We remark that equation (2) can also be obtained classically by calculating \( \langle \cos^2 \beta \rangle_0 \) with the marginal Boltzmann distribution of the polar angle \( \beta \) in the trap \( f(\beta) = \sin \beta \exp(V_0 \cos^2 \beta/k_B T)/Z' \).

The total angular momentum expectation value of the initial distribution can be estimated with the semiclassical substitutions used above. In particular, carrying out the integral over \( m \) and subsuming the result into the new normalization \( Z' \) yields

\[
\langle j \rangle_0 \simeq \frac{1}{Z'} \int_{0}^{\infty} dj \, j^2 \exp \left( -\frac{\hbar^2 j^2}{2I k_B T} \right) = \sqrt{\frac{\pi I k_B T}{2\hbar^2}}. \tag{B.5}
\]

Appendix C. Decoherence free alignment dynamics

The time-dependent alignment \( \langle \cos^2 \beta \rangle_u \) due to the unitary dynamics is numerically calculated by carrying out the trace over the product of the time evolved state equation (4) and the operator-valued observable \( \cos^2 \beta \),

\[
\langle \cos^2 \beta \rangle_u = \sum_{j=0}^{\infty} \sum_{m=-j}^{j} \sum_{j'=-j}^{j} \sum_{m'=-j'}^{j'} \langle jm|\rho_u(t)|j'm'\rangle \langle j'm'\cos^2 \beta| jm \rangle, \tag{C.1}
\]

where the matrix elements of \( \rho_u(t) \) are given by equation (4) and

\[
\langle jm|\cos^2 \beta| j'm'\rangle = \frac{\delta_{mm'}}{3} \left[ (-1)^{m^{\prime}2} \sqrt{(2j'+1)(2j+1)} \left( \begin{array}{c} j^\prime \vspace{0.2cm} j \\ m \vspace{0.2cm} 0 \end{array} \right) \left( \begin{array}{c} j^\prime 2 \vspace{0.2cm} j \\ m 0 \vspace{0.2cm} -m \end{array} \right) + \delta_{jj'} \right]. \tag{C.2}
\]

Here, the angular brackets denote Wigner-3j symbols [49]. Due to their selection rules, equation (C.2) vanishes unless \( j = j' \), \( j' = \pm 2 \), which significantly simplifies the evaluation of the alignment (3).

Appendix D. Classical dispersion

We estimate the dispersion timescale by calculating the classical alignment loss of a Gaussian state released from the laser potential \( V_0 \) and approximating the rotor dynamics as flat. On a short timescale, the angle \( \beta \) evolves to \( \beta(t) \simeq \beta + p_\beta t/\hbar \), where \( p_\beta \) is the corresponding angular momentum. The marginal distribution of \( \beta \) and \( p_\beta \) follows from the Boltzmann distribution as \( g(\beta, p_\beta) = \sin \beta \exp(-p_\beta^2/2I k_B T + V_0 \cos^2 \beta/k_B T)/N \). With this
one obtains the expectation value

\[
\langle \cos^2 \beta \rangle_a \approx \int_0^\pi d\beta \int_{-\infty}^{\infty} dp_\beta \ g(\beta, p_\beta) \cos^2 \left( \beta + \frac{p_\beta t}{T} \right) = \langle \cos^2 \beta \rangle_0 e^{-\kappa^2 t^2} + \frac{1}{2} \left( 1 - e^{-\kappa^2 t^2} \right),
\]

with \( \kappa = \sqrt{2k_B T/\hbar} \).

Appendix E. Orientational revivals of SNRs

Figure E1 shows the alignment signal for SNRs as discussed in the main text, at a gas pressure of \( p_g = 5 \times 10^{-10} \) mbar. As in the case of CNTs, many revivals can be observed. The revival time is \( T_{rev} = 28 \) ms.

Appendix F. Macroscopicity

In order to estimate the macroscopicity of the superposition state reached in the proposed experiment one replaces the unitary time evolution of the rotation state \( \rho \) between particle release and revival by the dynamics described by the quantum Markovian master equation \( \partial_t \rho = -i[H, \rho] / \hbar + \mathcal{L}\rho \) discussed in [25]. In the present case, the superoperator \( \mathcal{L}\rho \) takes the form

\[
\mathcal{L}\rho = \frac{1}{\tau m_e^2} \left( \frac{1}{2\pi \sigma_q^2} \right)^{3/2} \int d^3q \exp \left( -\frac{q^2}{2\sigma_q^2} \right) \left[ \mathcal{B}[R^\dagger(\Omega) q] \rho \mathcal{B}^*[R^\dagger(\Omega) q] - \frac{1}{2} \{ \mathcal{B}[R^\dagger(\Omega) q], \rho \} \right],
\]

since it is permissible to neglect position displacements and to approximate the particle by a homogeneous mass density. Here, \( \tau \) gives the timescale on which the modification acts, \( m_e \) is the electron mass, \( \sigma_q \) is the width of the momentum-kick distribution, \( R(\Omega) \) is the operator-valued rotation matrix, and \( \mathcal{B}(q) \) is the Fourier transform of the mass density. This master equation is similar to that for rotational collapse dynamics in the model of continuous spontaneous localization [48]. For a homogeneous rod of mass \( M \), length \( \ell \), and whose symmetry axis points into direction \( m(\Omega) = R(\Omega) e_z \), one obtains

\[
\mathcal{B}[R^\dagger(\Omega) q] \simeq M \sin \left( \frac{\ell}{2\hbar} q \cdot m(\Omega) \right).
\]

The master equation can be solved exactly if the rotor is confined to a plane so that its orientation is characterized by a single angle \( \alpha \) and \( m(\Omega) = e_\beta(\alpha) \). The resulting macroscopicity of the planar rotor is then a lower bound to that of the linear top. The solution of the master equation can be calculated by using the discrete phase space of the orientation state [48]. If the rotor is initially perfectly aligned, \( \langle \cos^2 \alpha \rangle_0 = 1 \), the alignment at an integer multiple of the linear rotor revival time, \( t = nT_{rev} \), is reduced by the modification equation (F.1) according to

\[\text{Figure E1. Orientational alignment } \langle \cos^2 \beta \rangle \text{ of the silicon nanorod discussed in the main text, as a function of time at } T = 100 \, \mu K. \]
\[
\langle \cos^2 \alpha \rangle_n = \frac{1}{2} + \frac{1}{2} \exp \left[ -n \frac{T_{\text{rev}}}{\tau} \theta \left( \frac{\ell \sigma_0}{\hbar} \right) \left( \frac{M}{m} \right)^2 \right]
\]

where

\[
\theta(x) = \frac{1}{2} \int_0^\infty du \, u e^{-u^2/2} \int_0^{2\pi} d\varphi \frac{d\alpha}{2\pi} \int_0^{2\pi} \frac{d\alpha}{2\pi} \left( \sin \left( \frac{ux}{2} \cos \left( \alpha - \frac{\varphi}{2} \right) \right) - \sin \left( \frac{ux}{2} \cos \left( \alpha + \frac{\varphi}{2} \right) \right) \right)^2.
\]

The macroscopicity associated with the measured alignment signal \(\langle \cos^2 \alpha \rangle_n = (1 + f)/2\) is determined by inserting the maximum of equation (F.4), \(\theta_m \approx 0.12\), into equation (F.3), solving for \(\tau\) and taking the decadic logarithm. Here \(f = 2\langle \cos^2 \alpha \rangle_n - 1\) is the ratio of observed to expected signal visibility.

### Appendix G. Torque sensing

In order to quantitatively assess the sensing potential of the proposed revival experiment, the free time evolution (4) has to be replaced by that in presence of an external torque of magnitude \(N_{\text{ext}}\). For illustration, we consider a torque acting orthogonal to the polarization direction of the trapping laser, so that the operator-valued potential reads \(\Psi_{\text{ext}} = -N_{\text{ext}} \sin^2 \beta \cos^2 \alpha\) with the azimuth \(\alpha\).

The exact time evolution can in principle be obtained by numerically diagonalizing the rotational Hamiltonian \(H = J^2/2I + \Psi_{\text{ext}}\) to describe transitions between all relevant rotation states, which amounts to approximately 400 000 states for a \(T = 100 \mu K\) CNT. Since all resulting eigenvalues and eigenvectors must be stored, this method soon becomes numerically expensive, scaling as \(J^4\) with the maximal angular momentum quantum number \(J\). However, in the present case one can use that the considered torques \(N_{\text{ext}}\) are much smaller than the kinetic energy of the occupied states to apply degenerate perturbation theory.

The eigenergies of the Hamiltonian \(H\) can be obtained to first order in \(N_{\text{ext}}\) by diagonalizing for each \(j\) the matrices

\[
H_j = \frac{\hbar^2}{2I} \left[ (j + 1) I_j + V_j \right],
\]

where \(I_j\) is a \((2j + 1) \times (2j + 1)\) identity matrix and \((V_j) = \langle jm | \Psi_{\text{ext}} | jm' \rangle\). The resulting eigenenergies \(E_{jm}\) are generally non-degenerate and labeled by \(n = -j, \ldots, j\) for each \(j\). The eigenenergies are thus no longer an integer multiple of \(2\pi \hbar / T_{\text{rev}}\), leading to the accumulation of a phase, which causes the revival to diminish. Our simulations show that this phase accumulation is the dominant contribution to the revival decay, while the perturbation of the eigenstates can be ignored.

We thus replace the exact unitary time evolution by the \(j\)-conserving evolution operator \(U_j = \exp(-iH_j/\hbar)\), acting on the subspace of fixed \(j\), thereby reducing the above scaling to \(J^2\). Diagonalization of the initial state \(\rho_0 = \sum_k p_k |\psi_k\rangle \langle \psi_k|\), then yields the alignment signal in the numerically tractable (but still expensive) form

\[
\langle \cos^2 \beta \rangle = \sum_k p_k \sum_{j,j'} \langle \Psi_{jk} | C_{jj'} \rho U_j U_j^\dagger | \Psi_{jk}\rangle,
\]

where \(C_{jj'}\) is the \((2j + 1) \times (2j' + 1)\)-matrix with elements equation (C.2) and we defined the vector \(|\Psi_{jk}\rangle = \langle jm | \Psi_{jk}\rangle\). This expression can be simplified further by exploiting that \(C_{jj'}\) is only non-zero for \(j = j'\) and \(j = j' \pm 2\).

### ORCID iDs

Stefan Kuhn @ https://orcid.org/0000-0002-7974-1922
James Millen @ https://orcid.org/0000-0002-6950-3461
Markus Arndt @ https://orcid.org/0000-0002-9487-4985
Klaus Hornberger @ https://orcid.org/0000-0002-3145-1117

### References

[1] Arndt M and Hornberger K 2014 Nat. Phys. 10 271–7
[2] Arndt M, Nairz O, Vos-Andreae J, Keller C, Van der Zouw G and Zeilinger A 1999 Nature 401 680–2
[3] Friedman J R, Patel V, Chen W, Tolpygo S and Lukens E J 2000 Nature 406 43
[4] Kohstall C, Riedl S, Guajardo E S, Sidorenkov L, Denschlag J H and Grimm R 2011 New J. Phys. 13 065027
[5] Eibenberger S, Gerlich S, Arndt M, Mayor M and Tüxen J 2013 Phys. Chem. Chem. Phys. 15 14696–700
[6] Berrada T, van Frank S, Bücker R, Schumm T, Schaff J F and Schmiedmayer J 2013 Nat. Commun. 4 2077
[7] Kovachy T, Asenbaum P, Overstreet C, Donnelly C, Dickerson S, Sugarbaker A, Hogan J and Kasevich M 2015 Nature 528 530
[8] Robens C, Alt W, Meschede D, Emary C and Alberti 2015 Phys. Rev. X 5 011003
[9] Riedinger R, Wallucks A, Marinkovic I, Loschauera C, Aspelmeyer M, Hong S and Gröblacher S 2018 Nature 556 473
[10] Ockeloen-Korppi C, Damskaöe G, Pirkkalainen J M, Asjad M, Clerk A, Massel F, Woolley M and Sillanpää M 2018 Nature 556 478
[11] Bose S, Mazumdar A, Morley G W, Ulbricht H, Toros M, Paternostro M, Geraci A A, Barker P F, Kim M and Milburn G 2017 Phys. Rev. Lett. 119 240401
[12] Marletto C and Vedral V 2017 Phys. Rev. Lett. 119 240402
[13] Marshall W, Simon G, Penrose R and Bueno-meester D 2003 Phys. Rev. Lett. 91 130401
[14] Pikovski I, Vanner M R, Aspelmeyer M, Kim M and Brukner Č 2012 Nat. Phys. 8 393
[15] Romero-Isart O, Pflanzer A C, Blaser F, Kaltenbaek R, Kiesel N, Aspelmeyer M and Cirac I J 2011 Phys. Rev. Lett. 107 020405
[16] Scala M, Kim M S, Morley G W, Barker P F and Bose S 2013 Phys. Rev. Lett. 111 180403
[17] Bateman J, Nimmrichter S, Hornberger K and Ulbricht H 2014 Nat. Commun. 5 478
[18] Kaltenbaek R et al 2016 EPJ Quantum Technol. 3 5
[19] Wan C, Scala M, Morley G W, Rahman A A, Ulbricht H, Bateman J, Barker P F, Bose S and Kim M S 2016 Phys. Rev. Lett. 117 143003
[20] Pino H, Prat-Camps J, Sinha K, Venkatatesh B P and Romero-Isart O 2018 Quantum Sci. Technol. 3 025001
[21] Seideman T 1999 Phys. Rev. Lett. 83 4971
[22] Poulsen M D, Peronne E, Stappelfeldt H, Bisgaard C Z, Viftrup S S, Hamilton E and Seideman T 2004 J. Chem. Phys. 121 783–91
[23] Stickler B A, Papendell B and Hornberger K 2016 Phys. Rev. A 94 033828
[24] Zhong C and Robicheaux F 2016 Phys. Rev. A 94 052109
[25] Nimmrichter S and Hornberger K 2013 Phys. Rev. Lett. 110 160403
[26] Kane B 2010 Phys. Rev. B 82 115441
[27] Kuhn S, Asenbaum P, Kosloff A, Scafani M, Stickler B A, Nimmrichter S, Hornberger K, Cheshnovsky O, Patolsky F and Arndt M 2015 Nano Lett. 15 5604–9
[28] Hoang T M, Ma Y, Ahn J, Bang J, Robicheaux F, Yin Z Q and Li T 2016 Phys. Rev. Lett. 117 123604
[29] Kuhn S, Stickler B A, Kosloff A, Patolsky F, Hornberger K, Arndt M and Millo M 2017 Nat. Commun. 8 1670
[30] Stickler B A, Nimmrichter S, Martinetz L, Kuhn S, Arndt M and Hornberger K 2016 Phys. Rev. A 94 033818
[31] Zhong C and Robicheaux F 2017 Phys. Rev. A 95 053421
[32] Wu M, Wu N L Y, Firdous T, San I F, Losby J E, Freeman M R and Barclay P E 2017 Nat. Nanotechnol. 12 127
[33] Child M S 2014 Semiclassical Mechanics with Molecular Applications (Oxford: Oxford University Press)
[34] Hornberger K, Uttenthaler S, Brezger B, Hackermüller L, Arndt M and Zeilinger A 2003 Phys. Rev. Lett. 90 160401
[35] Hackermüller L, Hornberger K, Brezger B, Zeilinger A and Arndt M 2004 Nature 27 711
[36] Robins W R 2004 Phys. Rep. 392 1–119
[37] Yurke B and Stoler D 1986 Phys. Rev. Lett. 57 13–6
[38] Sanders B C and Milburn G J 1992 Phys. Rev. A 45 1919–23
[39] Kozinsky B and Marzari N 2006 Phys. Rev. Lett. 96 166801
[40] Liu K et al 2012 Nat. Nanotechnol. 7 325–9
[41] Kim U J, Liu X M, Furtado C A, Chen G, Saito R, Jiang J, Dresselhaus M S and Elend P C 2003 Phys. Rev. Lett. 95 157402
[42] Gieseler J, Deutsch B, Quidant R and Novotny L 2012 Phys. Rev. Lett. 109 103603
[43] Kiesel N, Blaser F, Deli E, Grass D, Kaltenbaek R and Aspelmeyer M 2013 Proc. Natl Acad. Sci. USA 110 14180–5
[44] Asenbaum P, Kuhn S, Nimmrichter S, Sezer U and Arndt M 2013 Nat. Commun. 4 2743
[45] Millen J, Fonseca P, Mavrogordatos T, Monteiro T and Barker P 2015 Phys. Rev. Lett. 114 123602
[46] Hansen K and Campbell E 1998 Phys. Rev. E 58 5477
[47] Bruckman M A and Steinmetz N F 2014 Chemical modification of the inner and outer surfaces of Tobacco Mosaic Virus (TMV) Virus Hybrids as Nanomaterials (Berlin: Springer) pp 173–85
[48] Schirski B, Stickler B A and Hornberger K 2017 J. Opt. Soc. Am. B 34 C1–7
[49] Brink D M and Satchler G 2002 Angular Momentum (Oxford: Clarendon)