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Optically Levitated Nanodumbbell Torsion Balance and GHz Nanomechanical Rotor

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Levitated optomechanics has great potentials in precision measurements, thermodynamics, macroscopic quantum mechanics and quantum sensing. Here we synthesize and optically levitate silica nanodumbbells in high vacuum. With a linearly polarized laser, we observe the torsional vibration of an optically levitated nanodumbbell. This levitated nanodumbbell torsion balance is a novel analog of the Cavendish torsion balance, and provides rare opportunities to observe the Casimir torque and probe the quantum nature of gravity as proposed recently. With a circularly-polarized laser, we drive a 170-nm-diameter nanodumbbell to rotate beyond 1 GHz, which is the fastest nanomechanical rotor realized to date. Smaller silica nanodumbbells can sustain higher rotation frequencies. Such ultrafast rotation may be used to study material properties and probe vacuum friction.

Levitated optomechanical systems provide a powerful platform for precision measurements with great isolation from the thermal environment [1–6]. Optically levitated nano- and microspheres have been used to demonstrate force sensing at the level of 10^{-21} N [7] and to search for interactions associated with dark energy [8]. Optically trapped nanoparticles can also be driven to rotate at high speed. Previously, a rotation frequency of about 10 MHz has been achieved [9–11]. It is desirable to increase the rotation frequency further for studying material properties under extreme conditions [12, 13] and probing vacuum friction [14–16].

Recently, a novel ultrasensitive torsion balance with an optically levitated nonspherical nanoparticle was proposed [17], utilizing the coupling between the spin angular momentum of photons and the mechanical motion of the nanoparticle [4, 17-20]. Torsion balances have enabled many breakthroughs in the history of modern physics [21–24]. For example, the Cavendish torsion balance (Fig. 1(a)) determined the gravitational constant and the density of the Earth [21]. An optically levitated nanoscale torsion balance can provide a rare opportunity to detect the Casimir torque [25–28], and test the quantum nature of gravity [29-31]. An essential step towards these goals is to optically trap a well-defined nonspherical nanoparticle in high vacuum. However, optically trapped nonspherical nanoparticles such as nanodiamonds and silicon nanorods in former experiments were lost at about 1 torr due to laser heating [10, 17, 32–34]. Additionally, several years ago, levitated nanodumbbells were theoretically proposed to study many-body phase transitions [19]. To our best knowledge, however, there was no report on optical levitation of a nanodumbbell in

vacuum prior to this work.

In this letter, we synthesize silica nanodumbbells with two different methods and optically trap them in high vacuum. With a linearly polarized laser (Fig. 1(b)), we observe the torsional vibration of a levitated nanodumbbell in high vacuum, which is an important step towards probing the Casimir torque [25–28] and the quantum nature of gravity [29–31]. With a circularly polarized laser, we drive the nanodumbbell to rotate beyond 1 GHz, which is the highest mechanical rotation frequency reported to date.

In a linearly polarized optical tweezer, the long axis of a nanodumbbell will tend to align with the polarization direction of the trapping laser (Fig. 1(b)). This is because the polarizability of the nanodumbbell along its long axis is larger than the polarizability perpendicular to its long axis. If the nanodumbbell is not aligned with the polarization direction of the optical tweezer, it will twist the polarization of the optical tweezer (Fig. 1(b)), as an analog of twisting the torsion wire by the lead spheres in the original Cavendish torsion balance (Fig. 1(a)). If the optical tweezer is circularly polarized, the nanodumbbell will be driven to rotate at high speed. The torsional vibration or rotation of the nanodumbbell can be detected by monitoring the change of the polarization of the trapping laser (Fig. 1(c)).

We have synthesized pure silica nanodumbbells using chemical and physical methods. The interpenetration of the two particles in a nanodumbbell is tunable. To synthesize nanodumbbells with diameter D and length L (inset of Fig. 1(c)), we first synthesize silica cores with a diameter of d = L - D by adding tetraethyl orthosilicate (TEOS) to a mixture of ammonia, pure water and

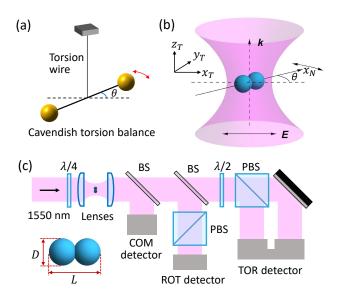


FIG. 1. (a) A simplified diagram of the original Cavendish torsion balance that has two lead spheres suspended by a copper silvered torsion wire. (b) A nanodumbbell levitated by a linearly polarized optical tweezer in vacuum. The linearly polarized optical tweezer provides the restoring torque that confines the orientation of the nanodumbbell. x_T, y_T, z_T are Cartesian coordinates of the trapping laser. x_T is parallel to the electric field **E** of the incoming linearly polarized laser, and z_T is parallel to the wave vector **k** of the laser. x_N is parallel to the long axis of the nanodumbbell. The angle between x_T and x_N is θ . (c) A simplified diagram for detecting the center-of-mass (COM) motion, the torsional (TOR) vibration, and the rotation (ROT) of a levitated nanodumbbell. The nanodumbbell is trapped at the focus of the lenses. The laser beam is initially linearly polarized. A quarter-wave $(\lambda/4)$ plate is used to control its polarization. BS: non-polarizing beam splitter; PBS: polarizing beam splitter; $\lambda/2$: half-wave plate. Inset: A nanodumbbell with diameter D and length Lcreated by attaching two identical nanospheres.

ethanol under stirring [35, 36]. Then acetone is added into the solution to induce aggregation. Next, a small amount of TEOS is added under stirring to grow the silica shells [35]. This chemical method can synthesize a large quantity of silica nanodumbbells with a tunable aspect ratio [35], but is demanding. So we also develop a physical method to assemble nanodumbbells. In this method, we first prepare a colloidal suspension of silica nanospheres in water. We then generate water microdroplets in air with an ultrasonic nebulizer [33]. By controlling the concentration of silica nanospheres, a fraction of water microdroplets ($\sim 5\mu m$ in diameter) contain 2 silica nanospheres in them. Two nanospheres in the same microdroplet will form a nanodumbbell as the water evaporates. Fig. 2(a), 2(b) show SEM images of our nanodumbbells with two different sizes. Their aspect ratio is between 1.9 and 2.

To optically levitate a silica nanodumbbell in vacuum, a 500 mW, 1550 nm laser is tightly focused with an NA

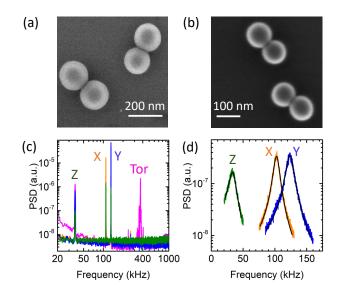


FIG. 2. (a), (b): SEM images of silica nanodumbbells with two different sizes. The scale bar is 200 nm in (a), and 100 nm in (b). (c) Measured power spectrum densities (PSD) of the torsional vibration (labeled by 'Tor') and the translational vibrations (labeled by 'X', 'Y', 'Z') of a 170 nm-diameter nanodumbbell optically levitated at 5×10^{-4} torr. (d) Measured PSD of the translational vibrations of the nanodumbbell levitated at 10 torr. The black curves are Lorentz fits.

= 0.85 objective lens in a vacuum chamber. The laser is initially linearly-polarized, and its polarization can be controlled with a quarter wave plate (Fig. 1(c)). Silica nanodumbbells are delivered into the optical trap at atmospheric pressure with an ultrasonic nebulizer [33]. Once a nanoparticle is trapped, we evacuate the vacuum chamber to below 0.01 torr, and then increase the pressure back to desired levels for measurements. This procedure removes extra nanoparticles in the chamber. To monitor the trapping process, a 532 nm laser is applied on the nanoparticle and the scattered light is viewed using a camera. We verify the trapped nanoparticle is a nanodumbbell by checking the ratios of damping coefficients for translational motions along different directions. The motion of the levitated nanoparticle changes the direction and polarization of the laser beam, which allows us to monitor the motion of a nanodumbbell with the same 1550 nm trapping laser (Fig. 1(c)) [17]. Fig. 2(c) shows the power spectrum density (PSD) of both torsional (TOR) vibration and center-of-mass (COM) vibration of a levitated 170-nm-diameter nanodumbbell in vacuum at 5×10^{-4} torr. This is an important step towards quantum ground state cooling of the torsional vibration and testing recent theoretical proposals [25, 29–31].

A levitated nanodumbbell will have anisotropic damping rates for translational motions in air if its orientation is fixed. We use the direct simulation Monte Carlo (DSMC) method to obtain the damping force and damping torque of a nanodumbbell in the free molecular flow

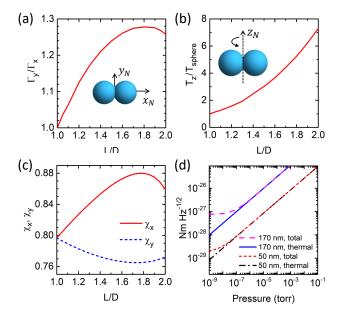


FIG. 3. (a) The ratio of air damping coefficients for translational motions perpendicular or parallel to its long axis $(x_N\text{-}\mathrm{axis})$ as a function of the aspect ratio (L/D) of a nanodumbbell. (b) Calculated normalized drag torque of the rotation of a levitated nanodumbbell around z_N axis as a function of the aspect ratio. (c) Effective susceptibilities of a silica nanodumbbell parallel (χ_x) or perpendicular (χ_y) to its long axis. (d) Calculated torque detection sensitivity of a levitated nanodumbbell with D=170 nm or D=50 nm as a function of air pressure. We assume L/D=1.9 in the calculations. The optical tweezer is assumed to be a focused 500 mW, 1550 nm laser with a waist of 820 nm.

regime [36-39]. In the simulation, molecules with uniform spatial distribution are launched from a spherical surface enclosing the nanodumbbell [36]. The speeds of these molecules satisfy a shifted Maxwell distribution to include the effect of the motion of the nanodumbbell [39]. Fig. 3(a) shows the calculated ratio of damping rates of a nanodumbbell moving along (Γ_x) and perpendicular (Γ_y) to its axial direction. The calculated ratio is $\Gamma_y/\Gamma_x = \Gamma_z/\Gamma_x = 1.276$ when L/D = 1.9, and $\Gamma_y/\Gamma_x = \Gamma_z/\Gamma_x = 1.258$ when L/D = 2. The measured ratios for the data of a 170 nm nanodumbbell shown in Fig. 2(d) are $\Gamma_y/\Gamma_x = 1.25 \pm 0.01$ and $\Gamma_z/\Gamma_x =$ 1.27 ± 0.02 , which agree excellently with our theoretical predictions. The DSMC method is also utilized to obtain the drag torque T_z on a nanodumbbell rotating at speed Ω . We then calculate the ratio T_z/T_{sphere} , where $T_{sphere} = \pi \mu D^4 \Omega/(11.976 \lambda_M)$ is the drag torque on a single sphere with diameter D rotating at the same speed Ω [39]. Here μ is the viscosity of air, and λ_M is the mean free path of air molecules which is inversely proportional to air pressure p_{air} . The calculated results are shown in Fig. 3(b). The damping rate of the rotation or torsional vibration around the z axis is $\Gamma_{\theta} = T_z/(I_z\Omega)$, where I_z

is its moment of inertia.

When the size of a silica nanodumbbell is much smaller than the wavelength of the trapping laser, the dipole approximation can be applied. The complex amplitude of the induced dipole of the nanodumbbell is $\mathbf{p} =$ $\alpha_x E_x \hat{x}_N + \alpha_y E_y \hat{y}_N + \alpha_z E_z \hat{z}_N$, where the complex amplitude of the electric field of the laser beam E is decomposed into components along the principle axes of the nanodumbbell. \hat{x}_N is in the direction along the long axis of the nanodumbbell. The components of the optical force F_i and the optical torque M_i acting on the nanodumbbell can be expressed as [40]: $F_j = \frac{1}{2} Re\{\mathbf{p}^* \cdot \partial_j \mathbf{E}\}$ and $M_j = \frac{1}{2} Re\{\mathbf{p}^* \times \mathbf{E}\}_j$. The quasi-static polarizability α_i^0 (j=x,y,z) of a nanodumbbell can be calculated assuming the electric field is static [41]. Fig. 3(c) shows the effective susceptibilities $\chi_x = \alpha_x^0/(\epsilon_0 V), \ \chi_y = \alpha_y^0/(\epsilon_0 V)$ of the nanodumbbell as a function of the aspect ratio. Here ϵ_0 is the permittivity of vacuum, and V is the volume of the nanodumbbell. $(\chi_x - \chi_y)/\chi_y = 0.14$ when L/D = 1.9. The dipole approximation can be improved by including the effects of radiation reaction due to the oscillation of the electric field in a laser beam. Then the polarizability is [40, 42, 43]: $\alpha_j = \alpha_j^0/[1 - ik_0^3 \alpha_j^0/(6\pi\epsilon_0)],$ where k_0 is the wave number. The real part of the polarizability $Re[\alpha_j] \approx \alpha_j^0$ is responsible for optical confinement and alignment, while the imaginary part $Im[\alpha_i]$ is important for optically rotating a nanodumbbell.

With calculated damping rates and polarizabilities, we can calculate the torque detection sensitivity of a levitated nanodumbbell. The minimum torque that it can measure limited by thermal noise is [44]: $M_{th} =$ $\sqrt{4k_BT_{env}I_z\Gamma_{\theta}/\Delta t}$, where T_{env} is the environmental temperature. In ultrahigh vacuum, the thermal noise from the residual air molecules becomes negligible and the minimum torque it can detect will be limited by the shot noise of the laser beam [25]: $M_{rad} = (\chi_x \chi_y k_0^2 V \hbar \sqrt{J_p/(3\pi\Delta t)}$. Here, $J_p = I_{laser}/(\hbar\omega_L)$ is the photon flux. I_{laser} is the laser intensity, and ω_L is the angular frequency of the laser. As shown in Fig. 3(d), at pressures below 10^{-8} torr, a nanodumbbell with D = 170 nm and D = 50 nm trapped in a 500 mW laser will have a torque detection sensitivity about $10^{-27} \text{Nm}/\sqrt{\text{Hz}}$ and $10^{-29} \text{Nm}/\sqrt{\text{Hz}}$, respectively. Remarkably, a levitated nanodumbbell at 10^{-4} torr is already much more sensitive than the current state-of-theart nanofabricated torsion balance, which has achieved a torque sensitivity of $10^{-22} \text{Nm}/\sqrt{\text{Hz}}$ at room temperature, and $10^{-24} \text{Nm}/\sqrt{\text{Hz}}$ at 25 mK in a dilution refrigerator [45]. The original Cavendish experiment measured a torque of about 10^{-7} Nm caused by gravitational attraction [21].

While a nanodumbbell levitated by a linearly-polarized optical tweezer can be an ultrasensitive nanoscale torsion balance, it will become an ultrafast nanomechanical rotor in a circularly-polarized optical tweezer (Fig. 4(a)). The frequency of the detected signal will be twice the rotation

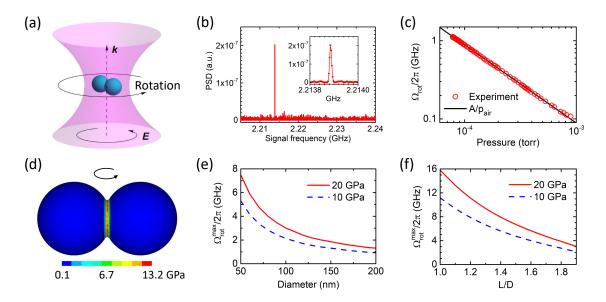


FIG. 4. (a) A nanodumbbell levitated by a circularly polarized laser will rotate. (b) Measured PSD of the rotation signal of a nanodumbbell. It has a sharp peak near 2.2 GHz. (c) Measured rotation frequency of the nanodumbbell as a function of pressure. (d) Calculated stress distribution of a nanodumbbell (D=170 nm, L/D=1.9) rotating at 1.2 GHz. (e) Calculated ultimate rotation frequency as a function of the diameter of the nanodumbbell when L/D=1.9. (f) Calculated ultimate rotation frequency as a function of the aspect ratio of the nanodumbbell when D=100 nm. The ultimate tensile strength is assumed be 20 GPa (red solid line) or 10 GPa (blue dashed line).

frequency of the nanodumbbell due to the symmetry of its shape. Fig. 4(b) shows a PSD of the rotation of a 170 nm-diameter nanodumbbell at 7.9×10^{-5} torr. The detected signal has a sharp peak near 2.2 GHz. This shows the nanodumbbell rotates at 1.1 GHz, which is much faster than that from former experiments [9-11]. Remarkably, the rotation is also very stable. The measured linewidth of the signal is less than 20 kHz, limited by the resolution of our spectrum analyzer, which leads to a rotating quality factor of larger than 10^5 . The steady-state rotation speed is determined by the balance between the optical torque M_z exerted on the nanodumbbell and the drag torque T_z from air molecules. Since drag torque is proportional to the air pressure, the steady state rotation speed Ω_{rot} is inversely proportional to the air pressure p_{air} . Fig. 4(c) shows the rotation speed as a function of the air pressure. The experimental data agrees with A/p_{air} , where A is a fitting parameter. As the air pressure decreases, the rotation speed increases. The maximum rotation frequency that we can measure is limited by the bandwidth of our photodetector.

As the rotation speed increases, eventually the nanodumbbell will fall apart due to the centrifugal force. The ultimate rotating frequency is determined by the ultimate tensile strength (UTS) of the material. Thus this experiment provides an opportunity to study material properties under extreme conditions. The stress distribution of a nanodumbbell rotating at high speed can be calculated by the finite element method. Figure 4(d) shows the stress distribution of a $D=170~\mathrm{nm}, L/D=1.9$

silica nanodumbbell rotating at 1.2 GHz. We assume the contacting point has a curvature radius of 5 nm. Remarkably, the maximum stress of the nanodumbbell under these conditions is about 13 GPa, which is 2 orders larger than the UTS of a bulk glass. This shows that our silica nanodumbbells is as strong as state-of-the-art silica nanowires [46]. The finite element method can be used to calculate the ultimate rotating frequency of silica nanodumbbells with different diameters and aspect ratios. The range of ultimate rotating frequency is calculated by setting the UTS to be in the range of 10-20 GPa, which agrees with our observation and the results for silica nanowires [46]. Figure 4(e) shows the calculated ultimate frequency is in the range of 1.1 - 1.6 GHz for D=170 nm, L/D=1.9 nanodumbbells. The ultimate frequency increases as the size of the nanodumbbell decreases. Figure 4(f) shows that for a given diameter D = 100 nm, the ultimate rotation frequency increases when the aspect ratio decreases. Thus nanodumbbells with smaller diameters and smaller aspect ratios can sustain rotation frequencies beyond 10 GHz.

In conclusion, we have synthesized and optically trapped nanodumbbells in high vacuum. When the laser is linearly polarized, the torsional vibration of the levitated nanodumbbell is observed, an important step towards detecting the Casimir torque [25] and developing a quantum Cavendish torsion balance for studying the quantum nature of gravity [29–31]. With a circularly polarized laser, 170 nm-diameter nanodumbbells were driven to rotate beyond 1 GHz, which is the highest re-

ported rotation frequency for a nanoparticle. Such high speed rotation may be used to study material properties and vacuum friction [12–16]. Based on Ref. [14, 16], a 100-nm-diameter SiC nanosphere rotating at 1 GHz near a SiC surface at 1000 K with a separation of 150 nm will experience a vacuum friction torque on the order of 10^{-28} Nm, which seems detectable at 10^{-8} torr.

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Note added. After completing this work, we became aware of a related work on fast rotation of a nanoparticle [47].

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