Absolute pressure and gas species identification with an optically levitated rotor

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ABSTRACT

The authors describe a novel variety of spinning-rotor vacuum gauge in which the rotor is a \( \approx 4.7 - \mu \text{m} \)-diameter silica microsphere, optically levitated. A rotating electrostatic field is used to apply torque to the permanent electric dipole moment of the silica microsphere and control its rotational degrees of freedom. When released from a driving field, the microsphere's angular velocity decays exponentially with a damping time inversely proportional to the residual gas pressure and dependent on gas composition. The gauge is calibrated by measuring the rotor mass with electrostatic co-levitation and assuming a spherical shape, confirmed separately, and uniform density. The gauge is cross-checked against a capacitance manometer by observing the torsional drag due to a number of different gas species. The techniques presented can be used to perform absolute vacuum measurements localized in space, owing to the small dimensions of the microsphere and the ability to translate the optical trap in three dimensions, as well as measurements in magnetic field environments. In addition, the dynamics of the microsphere, paired with a calibrated vacuum gauge, can be used to measure the effective molecular mass of a gas mixture without the need for ionization and at pressures up to approximately 1 mbar.

I. INTRODUCTION

Vacuum technology plays an integral role in science and technology. While a number of different technologies exist,1–3 under high vacuum conditions, absolutely calibrated pressure measurements are challenging. Some of the most sensitive pressure gauges ionize residual gas molecules and measure the resulting electrical current. Such ionization gauges require an empirical calibration, accounting for the efficiency of the ionization technique employed, which varies across molecular species and filament materials, and may change over time.4–8 Additionally, the production of cations and their associated electrons, inherent to ionization gauges, can have a detrimental effect on conditions within an experimental chamber. At higher pressures, the measurement of heat transport in a gas does not require ionization but still requires empirical and species-dependent calibrations.

Gauges based on mechanical measurements and the kinetic theory of gases are known to provide absolute measurements of pressure. For example, capacitance manometers measure the force on a membrane exposed to the residual gas. Force sensitivity limits the pressure accessible to such gauges to approximately \( 10^{-5} \) mbar.9 By comparison, spinning-rotor gauges measure the torsional drag induced by residual gas on a macroscopic rotor (an idea originally proposed by Maxwell9) which, in existing devices, is generally magnetically levitated. At high vacuum, in the molecular flow regime, such drag can be simply related to the pressure,10–15 resulting in an absolute calibration. The minimum measurable pressure, approximately \( 5 \times 10^{-7} \) mbar, is usually limited by systematic uncertainties or the required integration times.12,16,17

We have developed a spinning-rotor vacuum gauge based on an optically levitated, electrically driven microsphere (MS). The torsional drag on the MS can be measured by analyzing the rotational dynamics of the trapped MS. While silica MSs commonly used in optical levitation experiments are electrically neutralized,21–23 they have been shown to have residual electric dipole moments.24,25 Rotation can then be induced by applying torque with an electric field while being measured optically, as
described in Ref. 34. Similar techniques with an electric drive and optical readout have been demonstrated for graphene nanoplatelets in a Paul trap.36

In this scheme, the driving and interrogation mechanisms are not influenced by magnetic fields, and thus, the gauge can operate in conditions inaccessible to conventional spinning-rotor gauges. Another significant advantage of this technology, most applicable to experiments based on optically levitated particles, is that the measurement of pressure is relative to the environment in the immediate vicinity of the particle itself. A translation of the optical trap, or an array of optical traps each filled with a single rotating microsphere, would thus allow for mapping of pressure gradients with a spatial resolution limited only by the dimensions of the trapped particle(s).

II. EXPERIMENTAL APPARATUS

The apparatus is identical to that presented in Refs. 33–35. Briefly, a vertically oriented optical tweezer formed in vacuum by two identical aspheric lenses is surrounded by six independently biased electrodes, used to shield or manipulate the MS. A schematic of the inner region of the electrodes is shown in Fig. 1, together with a photograph of a trapped microsphere within the same electrode structure. One electrode in the foreground of both the schematic and the photo has been removed to facilitate viewing. A field-programmable gate array serves as a multi-channel direct-digital-synthesis waveform generator which drives the six electrodes to produce arbitrary electric fields inside the trapping region. Importantly, this system is capable of generating electric fields that are constant in magnitude, but rotating in direction, in order to drive the MS’s rotational degrees of freedom (DOFs) through their permanent electric dipole moment.

An optical system, detailed in Refs. 30 and 33, is used to measure the MS’s three translational DOFs and provide stabilizing feedback under high vacuum conditions. Polarization-sensitive optics, described in Ref. 34, measure the power of light in the polarization orthogonal to the trapping beam. Rotating, birefringent particles, such as the silica MSs used here, couple some linearly-polarized incident light into the orthogonal polarization at twice the frequency of their rotation, as demonstrated in Ref. 34. The phase of this power modulation can be used to deduce the rotation angle of the MS.

The vacuum system is similar to that in Ref. 34 with the addition of a residual gas analyzer (RGA), and a manifold providing He, N2, Ar, Kr, Xe, and SF6 gases to the chamber. The manifold can be evacuated with a dedicated scroll pump. Capacitance manometers, a Pirani gauge, and the RGA, serving as an ionization gauge, together determine pressures in the range of $\sim 10^{-6}$ – $10^3$ mbar and, with reduced accuracy, down to $\sim 10^{-8}$ mbar. This system allows for control of the species and pressure of gas present in the experimental chamber.

III. TORQUE NOISE AND TORSIONAL DRAG

Spinning-rotor vacuum gauges operate in the molecular flow regime, where they make use of the proportionality between gas pressure and the torsional drag induced by the gas in order to measure the chamber pressure. From the fluctuation dissipation theorem, the torsional drag coefficient $\beta_{\text{rot}}$ is proportional to the single-sided torque noise power spectral density $S_{\text{SN}}$,

$$S_{\text{SN}} = 4k_B T \beta_{\text{rot}}, \quad (1)$$

where $k_B = 1.381 \times 10^{-23}$ m² kg s⁻² K⁻¹ is the Boltzmann constant and $T$ is the temperature of the residual gas. $S_{\text{SN}}$ can also be computed by considering the force noise spectral density, $S_{\text{F}_r}$, imparted parallel to a surface element of the MS by successive gas molecule collisions and adsorption, followed by thermalization and re-emission, the common way to treat diffusive scattering. $S_{\text{F}_r}$ can be integrated across the surface of the MS to compute $S_{\text{SN}}$, which is then combined with Eq. (1) to obtain the relation

$$\frac{\beta_{\text{rot}} \times \nu_h}{P} = \pi r^4 \left( \frac{32}{9\pi} \right)^{1/2}, \quad (2)$$

FIG. 1. Schematic of the inner region of the shielding electrodes with a photograph of the same structure and a trapped microsphere, visible through a quartz vacuum window. The foreground electrode has been removed from the schematic in order to clearly show the trapping beam, as well as from the actual apparatus in order to capture this photograph and show the trapped microsphere. The trapping beam shown has a numerical aperture of 0.12 and a Rayleigh range of $\sim 25$ μm. For all measurements discussed, all six electrodes are in place.
where \( v_{th} = \sqrt{\frac{k_B T}{m_0}} \) is the characteristic thermal velocity of residual gas molecules of mass \( m_0 \) at temperature \( T \), \( P \) is the gas pressure, and \( r \) is the MS radius. This is identical to the results in Refs. 13 and 15 and allows an absolute measurement of the pressure \( P \) via \( \beta_{rot} \). The right hand side of Eq. (2) is a geometric factor that depends only on \( r \), the radius of the MS itself. Thus, if the gas species (i.e., \( v_{th} \)) is known, a measurement of the torsional drag constant \( \beta_{rot} \) can be used to directly infer the gas pressure from Eq. (2).

Two phenomena can be used to measure the torsional drag \( \beta_{rot} \) produced on a spinning MS. (1) When released from a driving field, the angular momentum of the MS will decay exponentially with a time constant inversely proportional to the torsional drag coefficient. This is the primary method used to measure pressure with a spinning-rotor gauge and depends only on the knowledge of the spherical rotor’s radius and moment of inertia. (2) When driven with an electric field, the torque exerted on the MS is proportional to the sine of the angle between the rotating field and the MS’s permanent dipole moment. Gas surrounding the MS produces a drag torque which determines the equilibrium angle between the driving field and the MS’s dipole moment. This angle can be measured as a phase difference between the driving field and the power modulation produced in the cross-polarized light.

It is distinctly possible that translational damping on resonance could also be used to infer the vacuum pressure. However, optically trapped MSs in the described apparatus exhibit instabilities at high vacuum, requiring active feedback in order to retain them for long periods. The active feedback is designed primarily with derivative gain to mimic the damping provided by residual gas in lower vacuum environments where the trap is stable. It is difficult to deconvolve this feedback with the damping provided by actual residual gas in the vicinity of the MS. At first order, the rotational degrees of freedom are unaffected by the stabilizing feedback.

**IV. SPINDOWN PRESSURE MEASUREMENTS**

From Eq. (2), the pressure can be written in terms of a constant \( \kappa = \kappa(T, r) \), the gas particle mass \( m_0 \), and the torsional drag coefficient \( \beta_{rot} \).

\[
P = \frac{\kappa}{\sqrt{m_0}} \beta_{rot},
\]

where \( \kappa = \frac{1}{r^4} \sqrt{\frac{9}{2}} k_B T / 32 \pi \) is determined by measuring the MS radius \( r \) and the experimental chamber temperature \( T \). The MS radius is determined indirectly by measuring the mass of the MS via electrostatic co-levitation and assuming the same uniform density found in Ref. 35, as the MSs are derived from the same lot. The drag coefficient \( \beta_{rot} \) is measured by observing the spindown time of the MS released from a driving field. If torsional drag due to residual gas is the only torque present, then the angular velocity of the MS will decay exponentially with time constant \( \tau = 1 / \beta_{rot} \), where \( I \) is the MS moment of inertia.

In fact, there is a roughly constant residual torque, likely optical in nature and generated from a small ellipticity in the trapping beam, which couples to the residual birefringence of the MS, as discussed in Refs. 31 and 34. Under these conditions, the solution of motion for the MS angular momentum and the solution for the frequency of rotation are given by

\[
\frac{dL}{dt} = N_{opt} \beta_{rot} \frac{L}{I}(t),
\]

\[
\Rightarrow f(t) = f_{rot} e^{-(t-t_0)/\tau} + f_{opt}(1 - e^{-(t-t_0)/\tau}),
\]

where \( L(t) \) is the angular momentum of the MS, \( f_{rot} \) is the initial rotation frequency at time \( t = t_0 \), \( \tau = 1 / \beta_{rot} \) is the decay time due to gas drag, and \( f_{opt} = N_{opt} / (2\pi\beta_{opt}) \) is the terminal rotation frequency of the MS due to a constant optical torque \( N_{opt} \). The quantity \( f_{opt} \) can be measured by allowing the MS to reach a steady state in the absence of any electrical driving torques and observing the terminal rotation frequency. Three different MSs were used in this work, with generally consistent results. For MS No. 1, the terminal rotation frequency was measured to be \( f_{opt} = (8315 \pm 559) \) Hz, where the uncertainty is the maximum observed deviation from the central value over approximately 6 h of successive measurements, where for each 2-s integration, the terminal rotation can be measured to within \( \pm 1 \) Hz uncertainty. This type of systematic effect is similar to the "offset correction" necessary for magnetically levitated spinning-rotor gauges.

To measure \( \tau \), the MS is released from an \( f_0 = 110 \) kHz driving field, and the frequency of power modulation of cross-polarized light produced by the rotation of the MS is observed. Successive 2-s integrations, separated by approximately one second, are first bandpass-filtered and then Hilbert-transformed to recover the instantaneous frequency of power modulation as a function of time. The median time of the \( i \)-th integration, \( t_i \), relative to the field turning off at \( t_0 \), and the arithmetic mean of the instantaneous frequency measured during this integration, \( f_i \), are used to calculate a value for \( \tau \) from Eq. (5),

\[
\tau_i = \frac{(t_i - t_0)}{\log\left(\frac{f_i - f_{opt}}{f_0 - f_{opt}}\right)}.
\]

The calculated value of \( \tau_i \) is plotted as a function of median integration time in Fig. 2 for four example measurements.
performed with MS No. 1. The shaded bands represent the quadrature sum of systematic and statistical uncertainties and are dominated by a systematic uncertainty in the assumed value of the terminal rotation frequency. For a single measurement, the estimation of $\tau$ from Eq. (6) is self-consistent for approximately the first 500-s. For longer times, the effect of a slowly changing optical torque exerts a torque roughly six orders of magnitude smaller than the drag torque at the initial rotation velocity. Anomalous damping may also arise from electric field noise on the, nominally grounded, driving electrodes. Results in plasma physics suggest that fluctuating fields in the frequency band around the rotation frequency of the MS could also tend to increase the angular velocity of the MS through stochastic acceleration, resulting in an apparent pressure difference across the experimental chamber. With the current apparatus, there is a substantial pumping impedance between the trapping region and the RGA filament, as the trap is confined within the six electrodes which form a cage.

While the presence of anomalous sources of dissipation cannot be excluded, two possible sources are shown to be too small to account for the pressure difference observed. The image charge induced on the 6 electrodes by the electric dipole moment would exert a torque roughly six orders of magnitudes smaller than the drag torque at the initial rotation velocity. Anomalous damping may also arise from electric field noise on the, nominally grounded, driving electrodes. Results in plasma physics suggest that fluctuating fields in the frequency band around the rotation frequency of the MS could also tend to increase the angular velocity of the MS through stochastic acceleration, resulting in an apparent pressure that would be lower instead of higher.

To test the second phenomenon, the equation of motion given by Eq. (4) was numerically integrated, including 100 distinct implementations of random torque fluctuations given by Eq. (1), and distinct realizations of anomalous torque from the electric field noise produced by the driving electronics when the field is nominally off. As this anomalous torque is incoherent with the decaying angular momentum of the MS, its effect on the apparent damping time was found to be less than 0.1% and cannot explain the higher value of the measured pressure.

The apparent pressure difference may also be due to an elevated temperature of the MS, which is a notoriously difficult quantity to measure, often yielding only an upper bound. The derivation of Eq. (2) assumes that the MS itself is in thermal equilibrium with the surrounding gas, so that incident and re-emitted gas particles have the same Maxwellian thermal velocity. If this assumption is removed, then the right-hand side of Eq. (3) would have a multiplicative factor $[2T_{\text{gas}}/(T_{\text{gas}} + T_{\text{MS}})]$, with $T_{\text{gas}}$ being the temperature of incident residual gas particles assumed to be in equilibrium with the chamber itself and $T_{\text{MS}}$ being the elevated temperature of the MS assumed to be in equilibrium with outgoing residual gas molecules. If the apparent pressure difference between the RGA and the location of the trap is due entirely to an elevated MS temperature, this implies $T_{\text{MS}} \sim 1500$ K.

**V. EQUILIBRIUM PHASE LAG**

The second method proposed to measure $\beta_{\text{rot}}$ can be substantially faster, but in our apparatus has only sensitivity for moderate vacuum. Since this is also the regime in which absolutely calibrated capacitance manometers operate, we use this method for a cross-

**TABLE I. Results of spindown pressure measurements, taken with MS No. 1. Statistical and systematic uncertainties are propagated independently from uncertainties on $r$, $l$, and $r$.**

| Pressure Measurement | Pressure (10^{-6} mbar) |
|----------------------|--------------------------|
| $P_0$                | 3.53 ± 0.17 (stat.) ± 0.30 (sys.) |
| $P_1$                | 3.56 ± 0.17 (stat.) ± 0.31 (sys.) |
| $P_2$                | 3.49 ± 0.17 (stat.) ± 0.30 (sys.) |
| $P_3$                | 3.43 ± 0.16 (stat.) ± 0.29 (sys.) |

**FIG. 3.** Exponential decay of MS No. 1 angular velocity due to torsional drag from residual gas after a 110-kHz driving field has been turned off. Data are shown for the same four measurements detailed in Fig. 2. Dotted lines indicated realizations of Eq. (5) with $\tau = r_j$ for measurements $j = 0, 1, 2, 3$, where $r_j$ is the average of the values plotted in Fig. 2. The structure in the residuals, with amplitude ~2%, is likely the result of a slowly fluctuating optical torque.

**J. Vac. Sci. Technol. B 38(2) Mar/Apr 2020; doi: 10.1116/1.5139638**

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check of the technique. Consider an MS rotating at a fixed frequency under the influence of a rotating electric field. In the rotating reference frame, the equation of motion for the phase \( \phi \) between the driving field and the orientation of dipole moment has an equilibrium solution developed in Ref. 34 and given by

\[
\phi_{eq} = -\arcsin\left(\frac{\beta_{tot, rot}}{Ed}\right) = -\arcsin\left(\frac{P}{P_{\text{max}}}\right),
\]

where \( \omega_0 \) is the angular driving frequency, \( E \) is the electric field magnitude, and \( d \) is the permanent electric dipole moment of the MS. The expression is written in terms of a single parameter \( P_{\text{max}} \), the maximum pressure that has a valid equilibrium solution given particular values of \( E, d, \) and \( \omega_0 \). Above this pressure, the driving field can no longer provide sufficient torque to maintain the MS’s rotation. Using Eq. (2),

\[
P_{\text{max}} = \frac{Ed}{\omega_0} \frac{P}{\omega_0} = \frac{Ed}{\omega_0} \frac{\kappa}{\sqrt{\gamma_0}},
\]

where \( \kappa \) is already defined. The prefactor \( (Ed/\omega_0) \) is related to the driving torque and can be set by the experimenter, whereas \( \kappa \) is a constant across gas species, as the MS radius is unchanged throughout the measurements.

Thus, a gauge cross-check consists of measuring \( P_{\text{max}} \), which is inversely proportional to the damping constant, as a function of \( m_i \), the particle mass of a residual gas species, while maintaining a constant driving torque and chamber temperature. \( P_{\text{max}} \) can be measured by linearly ramping the pressure, monitored with a capacitance manometer, while continuously measuring \( \phi_{eq} \) until unlocking, at which point the phase lag \( \phi \) becomes random, as shown in Ref. 34.

This effect is demonstrated for He, Ar, and SF\(_6\) in Fig. 4. Due to both the librational motion of the MS and torque noise from the increasing pressure of residual gas, the MS rotation becomes unlocked from the driving field at a pressure slightly below \( P_{\text{max}} \). Hence, \( P_{\text{max}} \) is determined by extrapolating the end point of the arcsine relationship in Eq. (7), as seen in Fig. 4.

The measurement of \( \phi_{eq} \) is used to validate Eq. (2) and cross-check the MS as a spinning-rotor gauge against the capacitance manometer. By comparison, magnetically levitated spinning-rotors often make use of static expansion (wherein a defined amount of gas is allowed to fill a chamber of known volume) in order to provide cross-checks with ion gauges. Practical limitations of the current apparatus, primarily the bandwidth of the driving electronics, place a lower bound on the pressure observable with this method at \(~10^{-5}\) mbar, as at lower pressures \( \phi_{eq} \) is too small to measure. Rotation velocities greater than the \(~100\) kHz achievable in the current system may allow for the extension of the method to lower pressures.

Silicon nanorods in counterpropagating beam traps have also been used to measure equilibrium phase lag induced by gas drag, but with the rotation driven optically by modulating the polarization (from linear to circular) of the trapping beams at a fixed frequency.\(^{14}\) The phase lag between the optical drive and the induced rotation of the silicon rotor was demonstrated to accurately predict gas pressures in the range of \(~4\)–\(~10\) mbar.

**VI. GAS COMPOSITION AND CHANGES IN DIPOLE MOMENT**

In many vacuum chambers, the ultimate base pressure achieved is limited by a single residual gas species, such as water vapor or hydrogen, and Eq. (8) is directly applicable. In this work, requiring the deliberate introduction of various gas species, a small correction needs to be applied to account for contamination, primarily by water. When there are multiple species, each contributes independently to the total torsional drag. Appealing to the analysis in Refs. 13 and 15, we can add the torque noise spectra of multiple gases, each with partial pressure \( P_i = \chi_i P_{\text{tot}} \) and particle mass \( m_{ij} \),

\[
S_{N,\text{tot}} = 4k_B T \beta_{\text{rot}} = 4k_B T \frac{P_{\text{rot}}}{\kappa \sqrt{m_{\text{eff}}}},
\]

where we have written the expression in terms of an effective mass, \( m_{\text{eff}} = \left(\sum \chi_i m_{ij}\right)^2 \). The mole fractions, \( \chi_i \), can be measured with an RGA, appropriately accounting for differences in ionization probability for different gas species. For He, N\(_2\), Ar, and SF\(_6\), the preparation of the manifold results in an apparent purity of \( \approx99.9\% \), limited by systematic uncertainties in RGA ionization probabilities for different gas species. In the cases of Kr and Xe, the apparent purity is \( \approx99\% \), limited by water contamination in the manifold.

The process of characterizing the residual gas leaked into the experimental chamber makes use of an RGA on the experimental chamber itself, which tends to charge the trapped MS with an excess of electrons. The MS is returned to zero charge, prior to spinning up and performing the drag measurements, by repeated exposure to ultraviolet photons from an Xe flashlamp.\(^{23,25}\) The process of charging and discharging appears to change the MS electric dipole moment.

![Fig. 4. Equilibrium phase lag between the orientation of the MS dipole moment and a rotating electric field, vs the residual gas pressure determined by two capacitance manometers. Three distinct measurements are shown with He, Ar, and SF\(_6\) residual gas. As the pressure is increased, the phase lags according to Eq. (7), and \( P_{\text{max}} \) is measured from a fit to this expression. As \( \phi_{eq} \) approaches \(-\pi/2\), the MS rotation becomes unlocked from the driving field (\( \phi \) becomes random) at a pressure slightly below \( P_{\text{max}} \), due to librational motion of the MS and torque fluctuations from the increasing pressure of the surrounding gas.](image-url)
In order to measure \( \kappa \) as defined in Eq. (8), the term \((Ed/\omega_0)\) must be measured precisely. The quantities \( E \) and \( \omega_0 \) are controlled and measured precisely, and via methods presented in Ref. 34, it is possible to characterize the MS permanent dipole moment \( d \). This is done by analyzing the librational motion about the instantaneous direction of the electric field, together with a calculation of the moment of inertia, another derived quantity.

Over the course of all measurements presented here, the dipole moment assumed values between approximately 95 and 120 \( \varepsilon \)-\mu m for MS No. 1, between 30 and 55 \( \varepsilon \)-\mu m for MS No. 2, and between 80 and 110 \( \varepsilon \)-\mu m for MS No. 3. Without the RGA filament on, the dipole moment was measured to be constant, within measurement uncertainties, over the course of a day.

### VII. CAPACITANCE MANOMETER CROSS-CHECK

The MS spinning-rotor vacuum gauge is cross-checked via the following procedure: first, the gas manifold is prepared with a particular species and \( m_{\text{eff}} \) is characterized by leaking a sample of the gas into the experimental chamber where the RGA is present; second, the chamber is pumped to its base pressure, the MS is calculated, whereas the precision of the measurement of \( \kappa \) is limited by uncertainties in the measured dipole moment and \( P_{\text{max}} \).

| MS     | \( m_{\text{MS}} \) [pg] | Theory    | Experiment |
|--------|-----------------|-----------|-------------|
| No. 1  | 84.3 \pm 0.2 (stat.) \pm 1.5 (sys.) | 6.3 \pm 0.3 (stat.) \pm 0.5 (sys.) | 6.47 \pm 0.06 (stat.) \pm 0.25 (sys.) |
| No. 2  | 84.2 \pm 0.4 (stat.) \pm 1.4 (sys.) | 6.3 \pm 0.3 (stat.) \pm 0.5 (sys.) | 6.21 \pm 0.06 (stat.) \pm 0.24 (sys.) |
| No. 3  | 85.0 \pm 0.6 (stat.) \pm 1.5 (sys.) | 6.2 \pm 0.3 (stat.) \pm 0.4 (sys.) | 6.48 \pm 0.09 (stat.) \pm 0.17 (sys.) |
properties, as well as a tendency for nanoscale particles to be highly non-spherical (see scanning electron microscopy images in Refs. 44–49).

The consistency between the measured and computed values of $\kappa$ is also an indirect validation of the work presented in Ref. 35, which makes no assumptions about MS temperature or thermal equilibrium and demonstrates independence across vacuum pressures $P \approx 10^{-6} - 10^{-3}$ mbar. If the density computed there was incorrect, we would expect quantities derived from this density, such as $\kappa$, to be inconsistent with independent measurements of those quantities.

VIII. NON-IONIZING GAS ANALYZER

If the rotational dynamics of an MS with a known value of $\kappa$ are analyzed while the pressure is measured with a calibrated, species-independent vacuum gauge such as a capacitance manometer, the combination can be used as a non-ionizing gas analyzer, operable directly in moderate vacuum. In particular, this system would excel as a binary gas analyzer, comparing the concentrations of two gases, such as one might encounter in nanofabrication with dopants in a carrier gas, or in chemistry with a particular stoichiometric ratio of reagent gases.

From Eq. (3), we can solve for the effective residual gas particle mass to find

$$m_{\text{eff}} = \left( \frac{\kappa \beta_{\text{rot}}}{P} \right)^2 = \left( \sum_i \chi_i \sqrt{m_{\text{di}}} \right)^2,$$

where $\beta_{\text{rot}}$ can be determined from a spindown measurement, $\beta_{\text{rot}} = 1/\tau$, or from an equilibrium phase lag measurement, $\beta_{\text{eq}} = Ed \sin(-\phi_{\text{eq}})/\omega_0$, where $\phi_{\text{eq}}$ is strictly negative. The latter quantity can be measured continuously. Such a system has an immediate advantage over ionizing RGAs, as it can operate directly in the pressure regime of $P = 10^{-6} - 10^{-3}$ mbar, whereas an ionizing RGA requires $P < 10^{-4}$ mbar and is often connected to a vacuum chamber of interest by a leak valve, which can itself change the relative concentrations of gases. Indeed, the total absence of ionization offers its own advantage.

IX. CONCLUSION

We have demonstrated the use of an optically levitated silica microsphere as a microscopic spinning-rotor vacuum gauge. The gauge operates on the principle that the torsional drag is proportional to the gas pressure in the vicinity of the microsphere. A spinning microsphere driven by a rotating electric field is released from the driving field, and the subsequent decay of angular momentum induced by torsional gas drag is observed. The residual gas pressure is then inferred from the decay time and the calculated moment of inertia.

Once a microsphere has been calibrated by measuring its mass, pressures within the range of $10^{-6} - 10^{-3}$ mbar can be determined within 10 s with a precision of $\leq 7%$ under the given assumptions and limited by the uncertainty in the calibration $\kappa$, which, in turn, depends on uncertainty in the assumed microsphere density. The minimum measurable pressure could easily be reduced by using a faster initial rotation velocity, which is limited only by the bandwidth of the driving electronics in the current apparatus.

A second method measures the equilibrium phase lag of the electric dipole moment relative to a driving field, which is induced by the gas drag. This method is used to cross-check the microsphere spinning-rotor gauge against a capacitance manometer. The same bandwidth limitations in the driving electronics limit the minimum measurable pressure of this method to $\geq 10^{-5}$ mbar.

A geometric calibration factor $\kappa$ related to the microsphere radius is independently determined by measuring the torsional drag as a function of gas species, since the drag depends on the momentum imparted by gas particle collisions and, thus, the molecular mass. The torsional drag $\beta_{\text{rot}}$ has been shown to scale inversely to the thermal velocity and, thus, $\beta_{\text{rot}} \propto \rho_{\text{mol}}^{-1/2}$, with $\rho_{\text{mol}}$ being the molecular mass, validating the rotational dynamics presented here and in Refs. 13 and 15, and opening the possibility to measure the effective molecular mass of a mixture of gases without ionization and directly in moderate vacuum.

ACKNOWLEDGMENTS

The authors would like to thank the Moore group (Yale) for general discussions related to trapping microspheres. This work was supported, in part, by the National Science Foundation (NSF) under Grant No. PHY1802952, the Office of Naval Research (ONR) under Grant No. N00014-18-1-2409, and the Heising-Simons Foundation. A.K. acknowledges the partial support of William M. and Jane D. Fairbank Postdoctoral Fellowship of Stanford University. N.P. acknowledges the partial support of the Koret Foundation.

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