

SPECIAL ISSUE ARTICLE

Large colloidal probes for atomic force microscopy: Fabrication and calibration issues

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Abstract

Atomic force microscopy (AFM) is a powerful tool to investigate interaction forces at the micro and nanoscale. Cantilever stiffness, dimensions and geometry of the tip can be chosen according to the requirements of the specific application, in terms of spatial resolution and force sensitivity. Colloidal probes (CPs), obtained by attaching a spherical particle to a tipless (TL) cantilever, offer several advantages for accurate force measurements: tunable and well-characterisable radius; higher averaging capabilities (at the expense of spatial resolution) and sensitivity to weak interactions; a well-defined interaction geometry (sphere on flat), which allows accurate and reliable data fitting by means of analytical models. The dynamics of standard AFM probes has been widely investigated, and protocols have been developed for the calibration of the cantilever spring constant. Nevertheless, the dynamics of CPs, and in particular of large CPs, with radius well above 10 μm and mass comparable, or larger, than the cantilever mass, is at present still poorly characterized. Here we describe the fabrication and calibration of (large) CPs. We describe and discuss the peculiar dynamical behaviour of CPs, and present an alternative protocol for the accurate calibration of the spring constant.

KEYWORDS

atomic force microscopy, calibration, colloidal probes, deflection sensitivity (invOLS), spring constant

1 | INTRODUCTION

Atomic force microscopy (AFM) is a powerful tool to investigate molecular interactions at bio-interfaces, as well as their mechanical properties, with nanometric spatial resolution and 1 to 10 pN force resolution.^{1,2}

The force sensing capability of AFM is provided by the use of probes consisting of an elastic microlever (cantilever) with an interacting tip at its apex. A deflection Δz of the cantilever as small as 1/10 of a nanometre can be detected by means of an optical beam deflection,³ and translated into a tip-sample force F according to Hook's law: $F = k_{\text{app}}\Delta z$. The value of the spring constant of the cantilever can be selected in the range 0.01 to 100 N/m. Here an apparent spring constant k_{app} is used because due to the cantilever tilt θ (usually $\theta = 10^\circ\text{--}15^\circ$, see Figure 1) both the force F and the measured

deflection Δz are perpendicular to the sample surface, rather than to the cantilever axis. The intrinsic spring constant k in turn relates the deflection and the force normal to the cantilever axis.

While sharp AFM tips are mandatory when high spatial resolution is necessary, large spherical probes, also known as colloidal probes (CPs),^{4–7} can be advantageous when one or more of the following requirements are important: a well-defined interaction geometry (sphere on flat) and reduced stress and strain in mechanical tests; a good spatial averaging; a broader and smoother surface for probe functionalization. The well-defined interaction geometry makes CPs a suitable tools for the investigation of mechanical properties of soft or biological samples^{8–11}; the larger contact area allows to detect weak interaction forces^{12,13} and generally CPs provide an easier interpretation of data in terms of theoretical models.^{8,11,14,15}

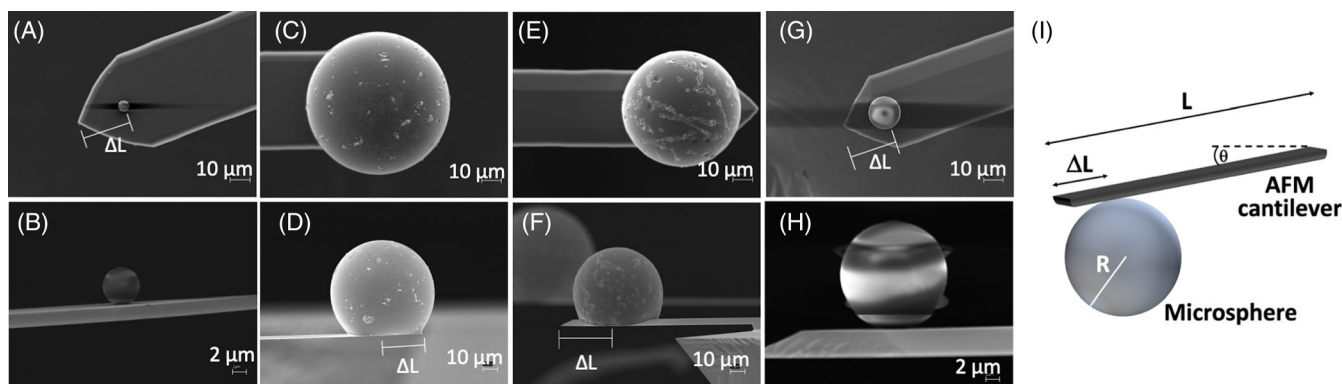


FIGURE 1 SEM images of some CPs fabricated (see Table 1). A, CM1, borosilcate; B, CM1 at higher magnification; C and D, CM4, soda-lime; E and F, TM4, soda-lime; G and H, TM3, borosilcate. I, Schematics of a CP, with the relevant parameters highlighted

The obvious price to pay when using CPs is the dramatic reduction of the spatial resolution, although the optimal spatial resolution clearly depends on the typical length scale of the physical phenomena under investigation. For example, as long as one is interested in characterizing the average rigidity of mesoscopic cellular regions (cell body vs periphery), tissues or extracellular matrices, CPs provide enough spatial resolution despite their micrometric radii.^{9,16–18}

Large CPs, where the radius R can be far larger than $10\ \mu\text{m}$, are very useful to study extremely weak interaction forces, or the mechanics of samples over a wide range of contact area, averaging nanoscale inhomogeneities and local variations of the sample (eg, in the case of the mechanical response of tissues). Large CPs allow to better compare the results obtained by AFM to those obtained by other indenters^{19–22} and in principle to bridge nano- and microscale mechanics to macroscale mechanics.^{23,24}

Despite the many advantages of CPs, some issues related to the calibration of these probes must be addressed in order to exploit their full potential, especially in the case of large CPs.²⁵ In particular, calibration of AFM probes require the determination of the intrinsic and apparent spring constants k and k_{app} and of the static deflection sensitivity S_z (in nm/V), also known as inverse optical lever sensitivity (invOLS), which converts the raw output ΔV of the photodetector (in Volts) into a deflection Δz (in nm) normal to the sample surface: $\Delta z = S_z \Delta V$.²⁶ The force F is then calculated as:

$$F = k_{\text{app}} S_z \Delta V \quad (1)$$

The calibration of both k (k_{app}) and S_z for CPs presents issues, which become more important as the probe radius increases.

The issues related to the calibration of CPs arise when the radius and mass of the attached sphere are comparable to the length and mass of the cantilever, respectively. In order to describe the effect of the attached sphere on the cantilever dynamics, two dimensionless parameters can be introduced (see also Section 2)²⁷: the reduced mass \tilde{m} is the ratio of the mass of the sphere m_s to the mass of the cantilever m_c ; the reduced gyration radius of the sphere \tilde{r} is proportional to the ratio of the radius R of the sphere to the length L of the cantilever.

The basic cantilever calibration parameters, such as the resonant frequency ν_0 , the quality factor Q , the deflection sensitivity S_z , the spring constant k , depend on \tilde{m} and \tilde{r} .

In this report, we discuss some critical aspects of CP production and calibration, and present methods to minimize their impact on the accuracy of AFM measurements.

2 | METHODS

2.1 | Fabrication of the CPs

We have produced eight monolithic glass probes following an established custom protocol²⁸ (see Table 1).

We used borosilicate glass spheres (Thermo Fisher Scientific, series 9000 glass particles) with radius $R = 2, 5, 10\ \mu\text{m}$ (probes CM1-3 and TM1-3), and soda-lime spheres (Polyscience, glass beads Amine, N° 23 584-10) with nominal radius $R = 30\ \mu\text{m}$ (probes CM4 and TM4); we could not find on the market borosilicate glass spheres with radius above $10\ \mu\text{m}$. These latter spheres are of far better quality than soda-lime glass spheres, in terms of surface roughness and morphology. Soda-lime glass spheres present extended surface defects (including craters, protrusions; see Figure 1C-F).

In order to span a wide range of reduced masses and gyration radii, we used two different kinds of silicon tipless (TL) cantilevers: tapping mode (TM; TL-NCH-50, Nanosensor) and contact mode (CM; TL-CONT, Nanosensor).

After mounting the cantilever into the AFM (Bioscope Catalyst, Bruker) tip holder, the beads were captured using the XY motorized stage of the AFM. The AFM is integrated in an optical inverted microscope (Olympus X71), which helps locating and capturing the beads. Depending on the size of the spheres, environmental capillary adhesion or Vaseline is exploited to attach the sphere to the tipless cantilever. The cantilevers with the attached beads are then transferred into a high-temperature oven (PXZ series, Fuji Electric) already at the target temperature, and heated for 2 hours. We used different temperatures for the different materials of the sphere: 780°C for borosilicate

TABLE 1 Properties of the CPs fabricated for this study

Probe	Bead radius (nm)	Cantilever length (μm)	Cantilever width (μm)	Cantilever thickness (μm)	\tilde{m}	\tilde{r}	$\Delta L/L$	β
CM1	2189	450	50	2	9.76×10^{-4}	5.7×10^{-3}	0.048	0.817
CM2	4462	450	50	2	8.18×10^{-3}	1.16×10^{-2}	0.053	0.826
CM3	7696	450	50	2	4.24×10^{-2}	2.02×10^{-2}	0	0.849
CM4	36 130	125	30	2	7.02	9.5×10^{-2}	0.088	1.018
TM1	1767	125	30	4	1.54×10^{-3}	1.67×10^{-2}	0	0.817
TM2	4292	125	30	4	2.21×10^{-2}	4.06×10^{-2}	0.112	0.836
TM3	7690	125	30	4	1.27×10^{-1}	7.2×10^{-2}	0.175	0.8768
TM4	31 752	125	30	4	9.62	3×10^{-1}	0.501	1.111

Note: These probes were fabricated using two different kind of tipless cantilevers: Tapping mode (TM) and contact mode (CM).

and 700°C for soda-lime glass, respectively. These temperatures are slightly below the softening point of the materials, which is qualitatively defined as the temperature at which a solid object begins to collapse under its own weight. After 2 hours, the microsphere is covalently attached to the cantilever; the glass locally melts at the sphere-cantilever interface, forming a neck, which is more or less extended depending on curing time and temperature.

Besides being much smoother than the soda-lime spheres and less dispersed in radius, borosilicate glass spheres present typically a lower deformation after the heating procedure (compare Figure 1B,H and Figure 1D,F). Moreover, the radius of soda-lime glass spheres can differ significantly from the nominal one.

2.2 | Calibration of CPs

2.2.1 | Radius

1. $R \leq 10 \mu\text{m}$. The characterization of the radius of the CP is obtained by an AFM reverse imaging of the probe,^{5,29,30} as described in Reference 28. The CP is used to scan a spiked calibration grating³¹ (TGT1, Tips Nano). Spikes are separated by approximately $3 \mu\text{m}$, therefore a $20 \mu\text{m} \times 20 \mu\text{m}$ image typically contains a hundred independent replicas of the probe apical region. Upon collecting several such images, the volume, projected area and height of each replica are measured. By fitting a spherical cap model to the experimental data, in the form, for example, of a volume vs height curve: $V = \frac{\pi}{3}h^2(3R-h)$, it is possible to determine the value of the sphere radius R , with an accuracy as good as 1%. While the spherical cap model adopted here allows using all data points from the three-dimensional reverse AFM image, deconvolution methods³² are limited by the fact that only so-called certainty regions can be determined, corresponding to those points where the tip apex is effectively in contact with the surface of the calibration sample, and analytical formulas are limited to the study of single profiles.³³ Since the spikes are relatively close, the sphere can penetrate through them by only a few hundred nanometres, at best. For this reason, as

long as gratings with sparser, and taller spikes are not available, the application of this method is limited to relatively small spheres.

2. $R = 30 \mu\text{m}$. CPs with $R > 10 \mu\text{m}$ are too large for the calibration by reverse AFM imaging. The radius of these probes is measured optically, using a calibrated metallographic optical microscope equipped with a calibrated 50x objective (AXIO, Zeiss). The error associated to the optical characterization of the probe radius is $1 \mu\text{m}$ (standard deviation). Since the nominal radius of the soda-lime spheres attached to the tipless cantilevers is $R = 30 \mu\text{m}$, the relative error of its characterization is approximately 3%. Alternatively, it is possible to follow an AFM-based reverse imaging approach based on single profiles, by scanning the large CP across a calibration grating with well separated ($>10 \mu\text{m}$) and tall ($>1.5 \mu\text{m}$) sharp steps. The radius R can be obtained by fitting an arc of circumference to the AFM profile. For a better characterization of R , we suggest to rotate the calibration grating in order to obtain different nominally equivalent profiles of the sphere, and average the different values.

2.2.2 | Nominal and reduced masses

We calculated the mass of the spheres m_s using the nominal densities of borosilicate glass $\rho_{\text{BG}} = 2.23 \text{ g/cm}^3$ and soda-lime glass $\rho_{\text{SLG}} = 2.52 \text{ g/cm}^3$ and the calibrated radius R as: $m_s = \rho_{\text{BG/SLG}} \frac{4}{3} \pi R^3$.

The mass of the cantilever m_c was evaluated assuming a rectangular section, using the nominal dimensions (length L , width w , thickness t) and the silicon density $\rho_{\text{Si}} = 2.33 \text{ g/cm}^3$ as: $m_c = \rho_{\text{Si}} Lwt$. We ignored the fact that some cantilevers have a trapezoidal cross section.

The total mass m_p of the probe was evaluated by summing the cantilever and the sphere masses: $m_p = m_c + m_s$.

The reduced mass \tilde{m} and the reduced gyration radius \tilde{r} of the sphere are defined as²⁷:

$$\tilde{m} = \frac{m_s}{m_c}, \tilde{r} = \sqrt{\frac{7R}{5L}} \quad (2)$$

We evaluated the effective mass m^* of the cantilever, before and after the attachment of the sphere, as the effective mass of a SHO,

using the standard formula: $m^* = k/\omega_0^2$, where k is the spring constant and ω_0 is the angular frequency of the oscillator.

2.2.3 | Deflection sensitivity

The calibration of S_z is performed by engaging the AFM tip on a very rigid surface and collecting a series of raw deflection vs z -piezo-displacement curves. Assuming that neither the tip nor the surface is deformed, the deflection sensitivity S_z is calculated as the inverse of the slope of the curves.²⁶ This sensitivity is referred to as the static deflection sensitivity, and differs from the dynamic deflection sensitivity $S_{z,dyn}$ ^{34,35}; the latter sensitivity must be used when the cantilever oscillates instead of being statically bent due to an end loading, as in the application of the thermal noise method for the calibration of the cantilever spring constant.

It must be noted that S_z strongly depends on the loading configuration.³⁶⁻³⁸ In particular, the static sensitivity of a tilted tipless cantilever differs from that of a standard tilted cantilever, with a sharp tip, or a sphere, attached to its end.

In the case of a standard probe, the force is not applied at the cantilever end, but rather at the tip apex, which is some distance apart, therefore a torque is generated. Moreover, because of the tilt angle θ , during the standard operation of an AFM the interaction force is perpendicular to the sample surface, instead of the cantilever axis, while during the thermally driven oscillation of the cantilever the effective driving force is always perpendicular to the cantilever axis. The corresponding cantilever displacement is therefore different from that obtained by the application of the same force at the cantilever end.

Corrections for the deflection sensitivity S_z are necessary, to account for the differences in the loading configuration of tipless and tipped cantilevers in the different experimental conditions. Correction factors are therefore used to determine both the intrinsic spring constant k using the thermal noise method, and the apparent spring constant k_{app} used to rescale the raw deflection signal into a force (Equation 1). While the latter correction factor is discussed in Section 2.2.5, the first factor is introduced in the following paragraphs.

As mentioned, during the thermal noise calibration, the cantilever is driven by a distributed thermal force, which can be considered as effectively located at the cantilever end, where it acts perpendicularly to the cantilever axis. Assuming that the deflection sensitivity depends on the parameters θ , R , L , ΔL introduced in Figure 11, the correction factor is therefore $\hat{s} = S_z^{0,L,0} / S_z^{\theta,R,L,\Delta L}$, where $S_z^{\theta,R,L,\Delta L}$ represents the deflection sensitivity measured in a particular loading configuration and $S_z^{0,L,0}$ is the sensitivity in the thermal loading configuration. According to Reference 37 (Equation 3 and other equations therein):

$$1/\hat{s}^2 = \left(1 - \frac{\Delta L}{L}\right)^2 \left[\frac{1 - \frac{3}{2} \frac{R/L}{1 - \frac{\Delta L}{L}} \tan \theta}{1 - 2 \frac{R/L}{1 - \frac{\Delta L}{L}} \tan \theta} \right]^2 \cos^2(\theta) \quad (3)$$

The $\cos^2(\theta)$ term in Equation (3) accounts for the tilt θ of the cantilever³⁹; the term in square brackets accounts for the torque applied to the cantilever because of the finite height of the tip; the first term $(1 - \Delta L/L)^2$

accounts for the fact that when the static deflection sensitivity is calibrated, the deflection at the loading point, rather than at the end of the cantilever, is measured.^{37,38} A special case of Equation (3) was first proposed by Hutter³⁸ under the hypothesis $\Delta L = 0$; in this case, Equation (3) simplifies to:

$$1/\hat{s}^2 = \left[\frac{1 - \frac{3}{2} \frac{R}{L} \tan(\theta)}{1 - 2 \frac{R}{L} \tan(\theta)} \right]^2 \cos^2(\theta) \quad (4)$$

For standard AFM tips, the radius R of the sphere in Equations (3) and (4) must be replaced by the tip height H .³⁷

2.2.4 | Thermal noise and CPs

Several approaches have been developed to characterize the spring constant of a cantilever,^{6,40-45} among which, the thermal noise method^{40,41,46,47} is based on the equipartition theorem.

Under the assumption that the mass m_t of the tip is negligible with respect to the mass m_c of the cantilever, the probe (cantilever + tip) can be modelled as a point body with effective mass m^* , bound to an ideal spring with stiffness k , in thermal equilibrium with its environment.

Assuming that the tip oscillates only along the direction perpendicular to the cantilever, and being z_{\perp} its displacement relative to the equilibrium position, its average potential energy is $U = \frac{1}{2}k \langle z_{\perp}^2 \rangle$, where $\sqrt{\langle z_{\perp}^2 \rangle}$ is the thermally driven root mean square oscillation amplitude of the free end of the cantilever, where the tip is located, ranging from approximately 5 to 500 pm. According to the equipartition theorem, $U = \frac{1}{2}k_B T$, where k_B is the Boltzmann constant and T is the local absolute temperature. It then follows:

$$1/2k \langle z_{\perp}^2 \rangle = 1/2k_B T \quad (5)$$

If $\langle z_{\perp}^2 \rangle$ is measured, the cantilever spring constant k is then calculated from Equation (5) as:

$$k = \frac{k_B T}{\langle z_{\perp}^2 \rangle} \quad (6)$$

When the optical beam deflection method is used to measure the cantilever deflection, with the laser aligned at the end of the cantilever, Equation (6) is replaced by Equation (7)^{37,38,48}:

$$k = \frac{\hat{\alpha} k_B T}{\chi^2 (\hat{s} S_z)^2 P_V} \quad (7)$$

In Equation (7), $P_V = \langle \Delta V^2 \rangle$ is the area below the first resonant peak in the power spectral density (PSD) of the raw deflection signal (in units of V^2/Hz).

The factor $\hat{\alpha}$ in Equation (7) takes into account the fraction of the total oscillation amplitude stored in the first normal mode of the cantilever, when one goes beyond the single harmonic oscillator (SHO)

approximation.⁴⁶ For the first normal mode of a tipless rectangular cantilever, $\hat{\alpha} \equiv \hat{\alpha}_1 = 0.971$.

The factor χ ^{34,35,48} corrects for the differences between the static (which is typically measured) and dynamic deflection sensitivities: $\chi = S_{z,dyn}/S_z$.^{46,49} Indeed, as pointed out by Butt and Jaschke in their seminal work,⁴⁶ the cantilever inclination, rather than its vertical displacement, is measured using the optical beam deflection apparatus; the slope of the free end of the statically loaded cantilever differs from those of the modal shapes of the oscillating cantilever.^{46,48,50} For the first normal mode of a tipless rectangular cantilever, assuming a negligibly small laser spot aligned at the end of the cantilever, $\chi \equiv \chi_1 = 1.09$. For the general case, the reader is referred to References 37 and 50 for the calculation of χ .

The factor \hat{s} represents the correction to the deflection sensitivity due to the change in the loading condition of the cantilever (Equations 3 and 4).

In principle, the thermal noise method could be applied to higher normal modes, in which case Equation (7) should be rewritten in terms of the parameters $P_{V,i}$, $\hat{\alpha}_i$, χ_i , where the index $i = 1, 2, 3, \dots$ refers to the i -th normal mode. In practice, due to the finite laser spot size, which causes an averaging of different cantilever inclinations across a finite area, the first normal mode has to be preferred³⁵; moreover, the first normal mode typically provides the largest oscillation amplitude, that is, the largest signal to noise ratio.

The factors $\hat{\alpha}$ and $1/\chi^2$ can be merged in a single parameter $\beta = \hat{\alpha}/\chi^2$. For the first normal mode of an ideal rectangular cantilevers, $\beta = \beta_0 = 0.817$.

The implementation of the thermal noise model for AFM cantilevers assumes that the mass of the cantilever is uniformly distributed. For this statement to be valid, the mass of the tip must be negligible compared to the mass of the cantilever. When large spheres are attached at the end of a cantilever, this hypothesis is not always satisfied. Since the mass of the microsphere scales up with the cube of radius R , it can easily reach values comparable to the mass of the cantilever. This typically happens for a radius $R > 10 \mu\text{m}$, for a $200 \mu\text{m}$ long, silicon nitride cantilever, and a glass sphere.

2.2.5 | Loading point and spring constant

For tipless cantilevers, the loading point is located at the free end of the lever. For sharp tips, the loading point corresponds to the location of the tip apex along the cantilever. For CPs, the loading point is located at the sphere apex, that is, approximately a distance ΔL from the cantilever end (Figure 11).

The attachment of the sphere to the tipless cantilever is not expected to change the intrinsic spring constant of the probe. Since the determination of the deflection sensitivity S_z can be critical for CPs (see discussion in Section 3.6), it can be convenient to calibrate the intrinsic spring constant k_{TL} of the tipless cantilever, before the attachment of the sphere, and then correct its value to determine the intrinsic spring constant k_{TL}^{LP} of the tipless cantilever at the loading point

of the CP; the latter spring constant can be supposed to be equal to the spring constant of the CP at the loading point, that is, $k_{TL}^{LP} = k_{CP}^{LP}$.

The spring constant k_{TL}^{LP} of the tipless cantilever corresponding to the loading point of the CP is calculated as^{51,52}:

$$k_{TL}^{LP} = \left(\frac{L}{L - \Delta L} \right)^3 k_{TL} \quad (8)$$

In Equation (8), ΔL is the distance of the loading point of the CP from the cantilever free end. In the case of rectangular, axisymmetric and relatively long cantilevers, Equation (8) is valid for arbitrary offset ΔL , not only in the limit $\Delta L/L \ll 1$.⁵²

Alternatively, the thermal noise method can be applied directly to the CP, provided the measured spring constant k_{CP} is corrected first by Equation (8) (to obtain k_{CP}^{LP}), and then by Equation (11), to take into account the peculiar dynamics of the CP, as described in the Results.

When the optical beam deflection method is used to measure the cantilever deflection, in order to rescale deflections into forces perpendicular to the sample surface, an apparent spring constant k_{app} must be used, instead of the intrinsic spring constant k .³⁷ The relation between the apparent and the intrinsic spring constants, assuming the sphere is located at the cantilever end, is:

$$k_{app} = \left\{ \left[1 - \frac{3R}{2L} \tan\theta \right] \cos^2(\theta) \right\}^{-1} k \quad (9)$$

where the correction factor accounts for the tilt of the cantilever θ and the torque applied on the cantilever because of the finite tip height; it was proposed by Heim et al,⁴⁰ and later corrected by Hutter³⁸ and generalized by Edwards et al³⁷ and Wang.⁵³ For standard AFM tips, the radius R of the sphere in Equation (9) must be replaced by the tip height (and not by half that value), as reported in Reference 37.

Following the approach of Edwards et al,³⁷ for CPs obtained from tipless cantilevers, assuming a sphere on cantilever configuration and considering a back-shift ΔL of the loading point, Equations (8) and (9) can be combined as:

$$\begin{aligned} k_{CP,app}^{LP} &= \left\{ \left[1 - \frac{3R/L}{2(1-\Delta L/L)} \tan\theta \right] \cos^2(\theta) \right\}^{-1} \left(\frac{L}{L - \Delta L} \right)^3 k_{TL} \\ &= \left\{ \left[1 - \frac{3R/L}{2(1-\Delta L/L)} \tan\theta \right] \cos^2(\theta) \right\}^{-1} k_{CP}^{LP} \end{aligned} \quad (10)$$

Equation (10) allows to calculate the apparent spring constant, at the loading point, of the CP from the intrinsic spring constant of the previously calibrated tipless cantilever.

The intrinsic spring constants are calibrated using the thermal noise method.^{41,46} Some details of the implementation of the thermal noise method for CPs are discussed in Section 3. The cantilever deflection signal is extracted from the AFM using a signal access module (intermodulation products) and sampled using an external ADC board (NI-DAQ PCI-6115). Signal acquisition and processing is

TABLE 2 Descriptions and abbreviations of the spring constants considered in this study

Spring constant name	Description
k_{TL}	Measured in the tipless configuration and related to the free end of the cantilever
k_{TL}^{LP}	Measured in the tipless configuration and related to the selected loading point of the probe
k_{CP}^{LP}	Measured with the attached sphere and related to the loading point of the probe corresponding to the position of the centre of the sphere
k_{app}	Apparent spring constant due to the tilt and torque of the cantilever in the standard experimental configuration

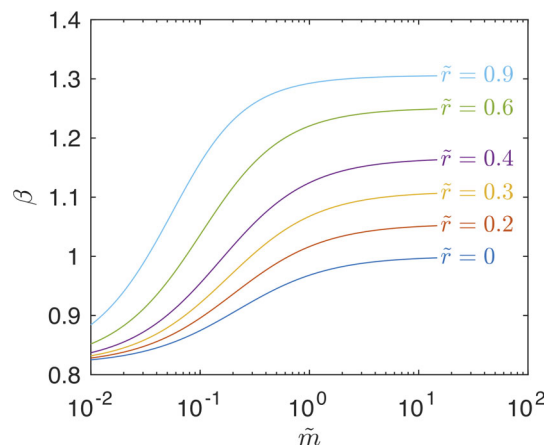
performed using custom software (LabView, National Instruments; see Reference 48 for practical indications). Alternatively, the Sader method can be used to calibrate the intrinsic spring constant of the cantilevers.^{45,54} The spring constant of a probe can be measured in different configurations (tipless cantilever vs CP) and it is always related to the loading point of the probe. For CPs, the loading point (LP) is located at the point where the tip/probe is attached to the cantilever; for TL cantilever, the LP is located at the cantilever free end. In Table 2 we summarize all the possible spring constant evaluated in this work, with the corresponding abbreviations.

3 | RESULTS

3.1 | Impact of the added mass on the dynamics of CPs

The laser doppler vibrometry (LDV) and in general interferometric techniques^{47,55-57} permit to measure with great accuracy the cantilever deflection instead of inclination, also in the case of oscillating cantilevers, including for the calibration of the spring constant by the thermal noise method. Interferometric techniques offer the advantage of not requiring the calibration of deflection sensitivity.

Using interferometric approaches it was recently demonstrated that the cantilever oscillation dynamics changes if a mass is attached to the free end.^{27,58} In particular Laurent et al²⁷ have measured the shape of first five eigenmodes along the profile of a rectangular cantilever, as a function of the reduced mass of the sphere \tilde{m} , and of the reduced gyration radius of the sphere \tilde{r} , introduced in Section 2. Apart from causing a decrease of the resonant frequency, the large attached mass of the sphere induces a shift of the normal mode nodes towards the free end of the cantilever, with the progressive shift of the centre of mass towards the cantilever free end and the increase of the contribution of the first normal mode to the total oscillation amplitude. As a consequence, both factors $\hat{\alpha}$ and χ are expected to be influenced by the presence of a large sphere at the end of the cantilever. It turned out indeed that the factor β_0 calculated for an ideal tipless cantilever

**FIGURE 2** Scaling of the factor β (see Section 2.2.4) as function of \tilde{m} , for different \tilde{r} , calculated according to the procedure described in Reference 27 based on the modified modal shape functions for a sphere-loaded cantilever

is no longer accurate, and leads to an underestimation of the spring constant measured by the thermal noise method.²⁷

Figure 2 shows the values of the factor β calculated according to the procedure described in Reference 27 based on the modified modal shape functions for a sphere-loaded cantilever, for both \tilde{m} and \tilde{r} varying over a broad range, considerably extending the previously reported results.

Once the spring constant $k_{CP,0}$ of the CP has been calibrated by the thermal noise method using the standard prefactor β_0 , the correct spring constant k_{CP} is calculated according to the formula:

$$k_{CP} = \frac{\beta}{\beta_0} k_{CP,0} \quad (11)$$

Representative curves $\beta \equiv \beta(\tilde{m})$ for different values of \tilde{r} are reported in Figure 2. In order to facilitate the application of Equation (11), Table 3 reports the values of the correction factor β for typical values of \tilde{m} and \tilde{r} .

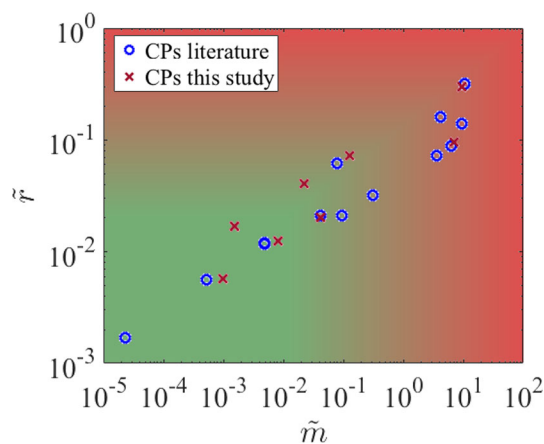
3.2 | A survey of CPs from the literature and the present study

CPs are fabricated with different cantilever and microsphere-relative dimensions and masses. Figure 3 reports the results of a literature survey about the values of \tilde{m} and \tilde{r} of CPs used in the experiments. Most of the CPs have relatively small spheres attached to the cantilevers, resulting in a reduced mass \tilde{m} on the order of 0.01 or even below.^{10,59-61} Nevertheless, very large CPs are also used,^{10,62-65} with \tilde{m} values as large as 5, up to 10.

In order to study how \tilde{m} and \tilde{r} of CPs affect the cantilever dynamics, we fabricated several CPs with different dimensions in order to cover the whole range shown in Figure 3. In particular, we produced eight probes, using both shorter and rigid tapping mode (TM), and longer and soft contact mode (CM) cantilevers. The properties of the CPs

TABLE 3 Values of the correction factor β (Equation 11) for different values of \tilde{m} and \tilde{r}

		\tilde{r}										
		0	0.1	0.2	0.3	0.4	0.5	0.6	0.7	0.8	0.9	1
\tilde{m}	0	0.817	0.817	0.817	0.817	0.817	0.817	0.817	0.817	0.817	0.817	0.817
	0.001	0.818	0.818	0.818	0.819	0.819	0.820	0.821	0.822	0.823	0.824	0.826
	0.002	0.819	0.819	0.820	0.820	0.821	0.823	0.825	0.827	0.829	0.832	0.834
	0.005	0.821	0.822	0.823	0.825	0.827	0.831	0.835	0.840	0.846	0.852	0.859
	0.01	0.825	0.826	0.828	0.832	0.837	0.844	0.852	0.861	0.872	0.884	0.897
	0.02	0.832	0.834	0.838	0.845	0.855	0.868	0.883	0.900	0.920	0.941	0.963
	0.05	0.851	0.854	0.864	0.879	0.901	0.927	0.957	0.990	1.024	1.059	1.093
	0.1	0.875	0.880	0.896	0.921	0.954	0.994	1.036	1.079	1.121	1.158	1.190
	0.2	0.905	0.913	0.936	0.972	1.018	1.068	1.117	1.161	1.199	1.230	1.254
	0.5	0.945	0.956	0.988	1.035	1.090	1.144	1.191	1.228	1.257	1.278	1.292
	1	0.968	0.981	1.016	1.068	1.124	1.177	1.220	1.253	1.276	1.292	1.304
	2	0.983	0.996	1.034	1.087	1.144	1.195	1.236	1.265	1.285	1.299	1.309
5	0.993	1.007	1.046	1.100	1.157	1.207	1.245	1.272	1.291	1.303	1.312	
10	0.996	1.011	1.050	1.105	1.162	1.211	1.248	1.275	1.293	1.305	1.313	

**FIGURE 3** CPs used by the authors of publications as from a literature survey (blue circles) and CPs fabricated during this study (red crosses) represented according to their reduced mass \tilde{m} and gyration radius \tilde{r} . The background colour highlights the region where corrections to the spring constant according to Equation (11) are negligible (green), and the region where the influence of the added mass becomes relevant (deviation larger than approximately 10%)

produced for this study are reported in Table 1 and also in Figure 3 (red crosses). We characterized the spring constant and the dynamics of our probes before and after the attachment of the microsphere.

3.3 | Shift of the resonant frequency of CPs

Figure 4 shows the thermal noise spectrum, that is, the raw PSD P_V in units of V^2/Hz , of the first resonant peak of the original tipless cantilevers, and of the CPs obtained upon attachment of the glass bead. Not surprisingly, as a consequence of the added mass, we observe a

decrease of the resonant frequency. These results confirm that a shift in the resonant frequency ν_0 is present even if the reduced mass is as small as $\tilde{m} = 9.76 \times 10^{-4}$ (Figure 5). This drop of the resonant frequency reaches a factor of 5 for largest masses (Figure 4D,H). These trends are in agreement with previous works.^{66,67}

Figure 4 shows that the increase of the full width at half maximum (FWHM) and of the oscillation amplitude are associated to the shift of the resonant peaks towards lower frequencies. For large CPs (Figure 4D,H), the peak is 30 times higher and 20 times narrower compared to the corresponding tipless cantilever case. These observations may be somewhat unexpected for probes whose mass is significantly increased. For instance, a reduction of the oscillation amplitude and a widening of the resonant peak is typically observed when the thermal noise spectrum is acquired in liquid, another case where the effective mass of the cantilever is increased significantly, due to the liquid viscosity that causes the cantilever to drag a finite volume of liquid during oscillation. As we will discuss later, this observation can be explained in terms of a combination of effects due to the increase of the inertial mass of probe⁶⁷ and the storage of a greater fraction of total oscillation in the first mode.²⁷

3.4 | Quality factor

The quality factor Q represents the ratio between the energy stored in the oscillation and the energy dissipated per cycle because of the viscous damping. For a single-mode oscillator, Q is defined as:

$$Q = \frac{m^* \omega_0}{\gamma} = 2\pi \frac{m^* \nu_0}{\gamma} \quad (12)$$

In Equation (12), γ is the damping coefficient, that is, the proportionality factor between the tip velocity and the viscous force. Q is related to

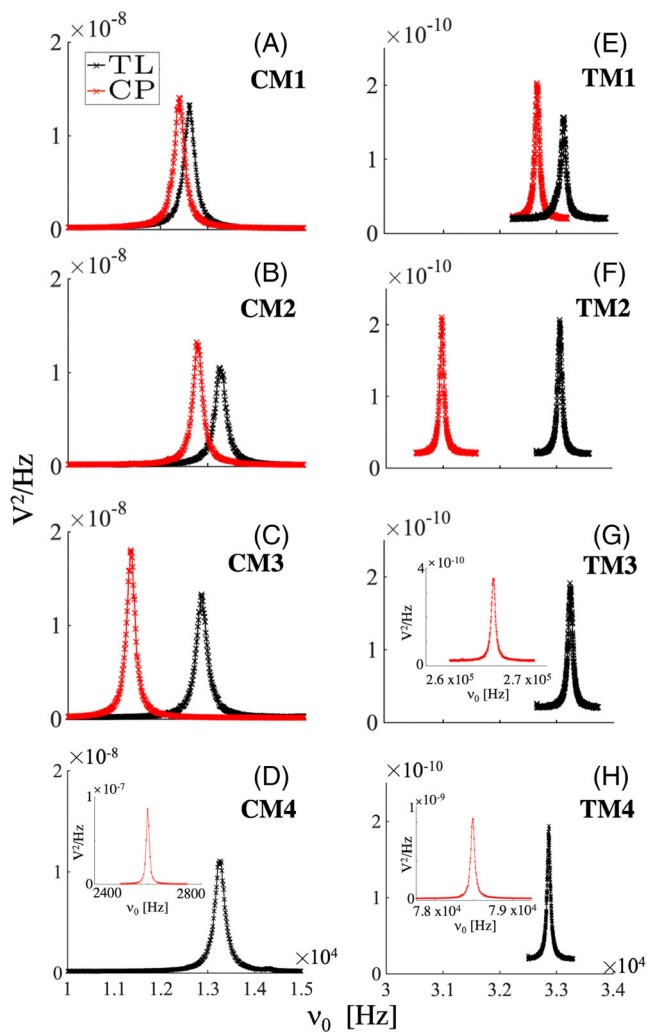


FIGURE 4 The measured PSD P_V at the first normal mode of the cantilever before (black) and after (red) the attachment of the microsphere for contact mode CPs. A-D, CMs; E-H, TMs. The mass of the attached sphere increases from top (CM1 (A)-TM1 (E)) to bottom (CM4 (D)-TM4 (H))

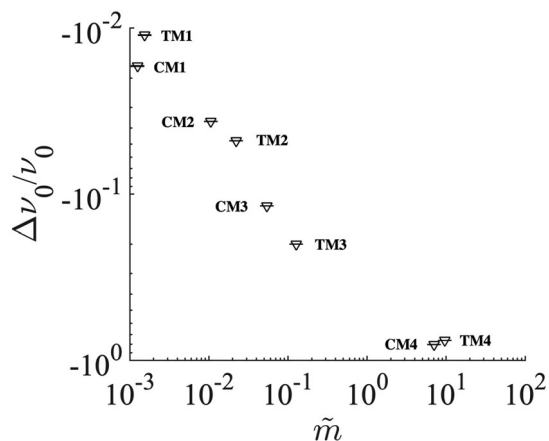


FIGURE 5 The relative shift of the measured resonant frequency ν_0 vs the reduced mass \tilde{m} . As shown in Figure 4, ν_0 decreases as \tilde{m} increases

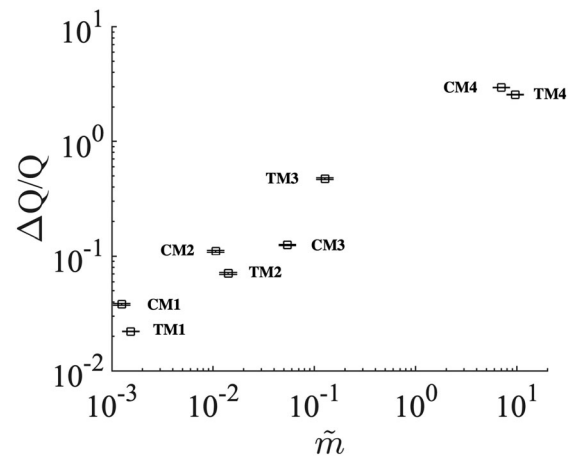


FIGURE 6 Relative shift of the measured quality factor Q vs the reduced mass \tilde{m}

the width of the resonant peak: at fixed mass m^* and frequency ν_0 , the higher is Q , the smaller is the FWHM, since $\text{FWHM} \approx \nu_0/Q$.

Figure 6 shows the relative increase of the quality factor Q of the CPs compared to the one measured in the tipless configuration. It is evident that the value of Q of CPs increases, rather than decreasing, with respect to the tipless cantilevers, and that the relative increase is larger for higher values of \tilde{m} .

To better understand the behaviour of Q upon the attachment of the sphere to the tipless cantilever, we calculated the effective mass $m^* = k/\omega_0^2$ for each probe, before and after the attachment of the bead. The ratio between the measured effective mass m^* and the nominal mass of the probe m_p (for both tipless probes and CPs) is shown in Figure 7A. The measured ratio m^*/m_p for the tipless cantilevers is 0.24 ± 0.02 (mean + standard deviation of the mean), in agreement with the theoretical value of 0.25 for the rectangular cantilever.⁴² A similar value is measured for those CPs with the smallest reduced mass, where the contribution of the cantilever mass is dominant (see Figure 7A). As \tilde{m} increases, the effective mass m_c^* becomes larger than $0.25m_c$. For large CPs, m^* becomes similar to m_p , which is in turn very similar to m_s , the mass of the sphere; in this limit, the mass of the sphere is dominant.

Equation (13) represents a simple model for the effective mass m_{CP}^* of a CP; the concentrated mass of the sphere, located at the free end of the cantilever, is added to the effective mass of the tipless cantilever²⁶:

$$m_{CP}^* = m_c^* + m_s \quad (13)$$

Equation (13) clearly highlights that the cantilever and the sphere contribute differently to the inertia of the probe. The cantilever mass, which is uniformly distributed along the cantilever length, is strongly underestimated in the sum, being weighted by a factor of $1/4$; the mass of the sphere, instead, being truly concentrated at the free end of the cantilever, is fully represented with weight 1. Figure 7B shows the correlation between the measured m^* and the values of the effective

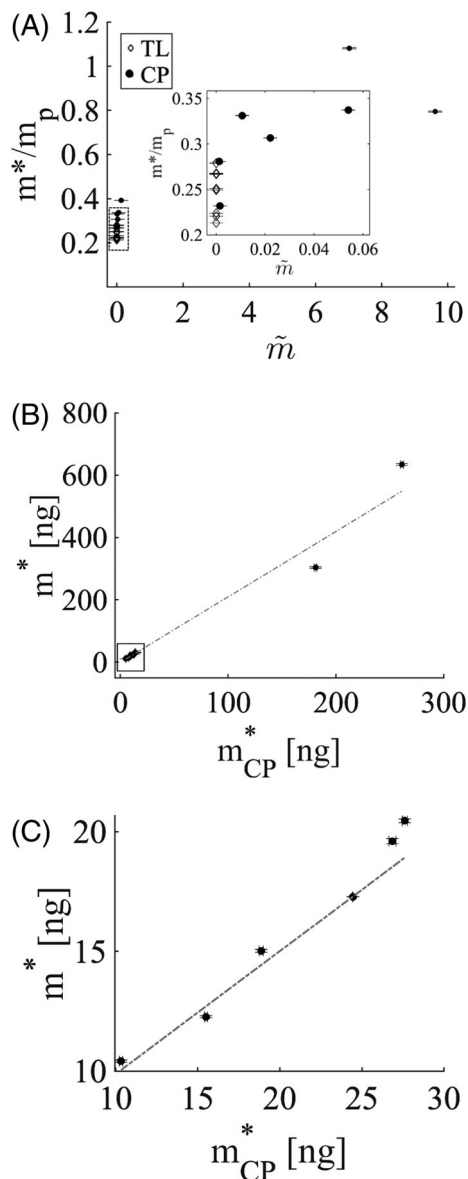


FIGURE 7 A Ratio of the effective mass m^* (calculated using Equation 12) to the nominal mass m_p of the fabricated probe as a function of the reduced mass \tilde{m} for tipless cantilevers (diamonds) and CPs (filled circles). B, Effective mass m^* of the CPs (calculated as $m^* = k/\omega_0^2$) as a function of m_{CP}^* (defined in Equation 13), with linear fit ($y = mx + b$, $m = 1.052 \pm 0.005$, $b = -1.14 \times 10^{-12} \pm 1.27 \times 10^{-13}$). C, Zoom on the data presented in (B) for $m_{CP}^* < 30$ ng highlighted in the drawn box, with the linear fit ($y = mx + b$, $m = 1.027 \pm 0.008$, $b = -1.14 \times 10^{-12} \pm 5.37 \times 10^{-13}$)

mass predicted by Equation (13). The slope of the straight line fitting the experimental data in Figure 7B is 1.052 ± 0.005 . In the case of the first six probes (data highlighted in Figure 7C), a better correlation is found, the slope being 1.027 ± 0.008 .

Figure 7B shows that Equation (13) represents a good model for the effective mass of CPs, which in turn confirms the accuracy of the description of (large) CPs as effective SHOs with the total mass concentrated at the loading point. Indeed, as the radius R of the sphere

increases, the mass of the sphere increases rapidly, scaling as R^3 , and the CP becomes more and more equivalent to an SHO with a concentrated mass equal to the sphere mass. Deviations for the largest CPs can be attributed to uncertainties in the determination of the sphere mass, due both to the uncertainties in the sphere density or in the sphere volume. SEM images (Figure 1) show indeed that soda-lime glass spheres possess many defects, which could also be present in the inner volume, as voids, or segregated phases with different density. Moreover, upon softening in the oven, the spheres deform to some extent in the region close to the cantilever. This process is supposed to conserve the volume of the sphere, but could alter the measurement of the radius, which is inferred from the transversal section of the sphere; as a consequence, errors in the volume estimation are likely.

The increase of the quality factor is in turn explained by Equation (12). Q is proportional indeed to the resonant frequency ν_0 and the effective mass m^* . Since ν_0 drops by a factor of 5 (Figure 5), while m^* increases by a factor of 40 (it passes from $0.24 m_c$ to approximately $10 m_c$), the latter change dominates and Q increases (considerably).

Eventually, Figure 7B,C and Figure 6 explain why the amplitude of the resonant peak of the first normal mode is enhanced, as shown in Figure 4. Indeed, large CPs are well described by a single-mode harmonic oscillator model, because the higher-order modes provide a less and less important contribution to the tip motion, as their last node approaches the free cantilever end²⁷; as the quality factor Q increases significantly, the constraint of energy conservation causes the amplitude to increase.

In conclusion, Equations (12) and (13) and the observed trends of both resonant frequency and quality factor explain the peculiarity of the thermal noise spectra of CPs with a large microsphere attached to their free end, in particular the fact that these probes, despite a marked reduction of the resonant frequency, have extremely sharp resonant peaks, with large Q and amplitude. The apparent anomaly of CPs can be attributed to the fact that, in the AFM community, an increase of the effective mass and a drop in the resonant frequency of the cantilever are typically observed when operating a soft cantilever in liquid,⁶⁸⁻⁷⁰ and they are associated to a marked drop of the quality factor Q due to the viscous damping. With (large) CPs in air, however, the factor γ in Equation (12) does not change significantly, while the dramatic increase of the mass is not counterbalanced by the reduction of the resonant frequency. A recent study confirms that the added mass mechanism described in this work dominates the effects related to energy dissipation and viscous drag also in the case of large CPs in liquid, where the attachment of large beads to the cantilever is exploited to enhance the quality factor and the force sensitivity of the probes.⁶⁷

3.5 | Spring constant of CPs

Figure 8A shows a comparison of the spring constants k_{TL}^{LP} , calibrated for the tipless cantilevers before the attachment of the glass beads, and then corrected for the shift of the loading point (Equation (8)), with the spring constants k_{CP}^{LP} of the CPs calibrated after the

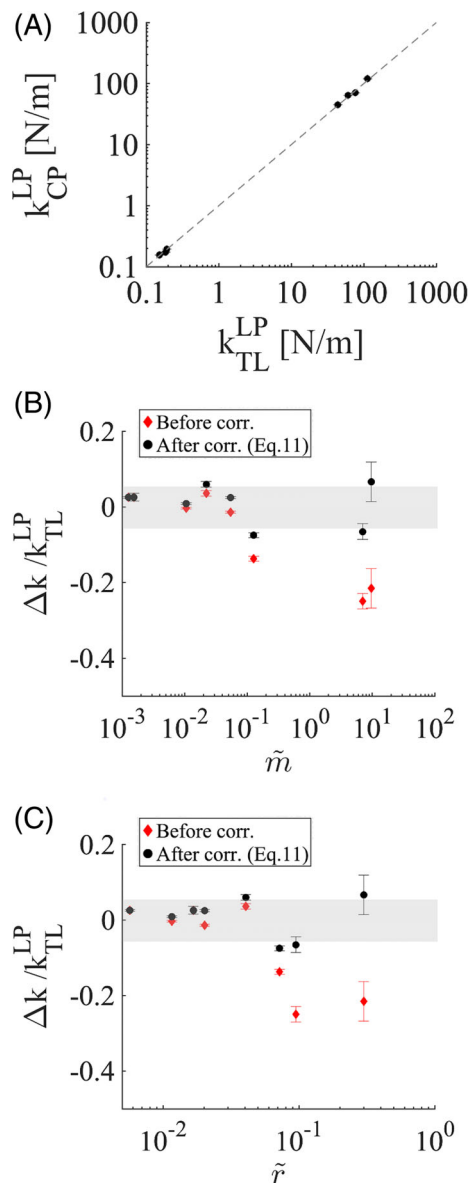


FIGURE 8 A, Spring constant k_{CP}^{LP} of CPs corrected according to Equations (8) and (11) vs the reference spring constant k_{TL}^{LP} of the tipless cantilever calculated at the loading point (Equation 8), measured before the sphere attachment. The dashed line represents a linear regression of the data, with slope 1.0090 ± 0.0006 . B and C, The relative difference between k_{CP}^{LP} and k_{TL}^{LP} as a function of the reduced mass and radius of the CP, respectively, without (red filled diamonds) and with (black filled circles) the correction by Equation (11). The shaded area represents the $\pm 5\%$ interval

attachment of the spheres, corrected for the loading point shift, and eventually for the added mass effect (Equation 11). A linear regression of the data provides a slope of 1.0090 ± 0.0006 , which confirms the effectiveness of the correction procedures.

Figure 8B,C shows the relative difference between the spring constant k_{TL}^{LP} and the spring constant k_{CP}^{LP} as a function of \tilde{m} and \tilde{r} , respectively. The spring constant values are shown before (red) and after (black) the β -factor correction according to Equation (11).

As shown in Reference 27, the effect of the added concentrated mass of the sphere is to induce an underestimation of the spring constant, if the standard thermal noise formula (Equation 7) is used, with $\beta = \beta_0 = 0.817$. When \tilde{m} and \tilde{r} are larger than approximately 0.1, the spring constant starts to decrease, reaching a maximum relative deviation from k_{TL}^{LP} of about 25% (Figure 8B,C, red filled diamonds); these deviations are effectively corrected by Equation (11) (Figure 8B,C, black filled circles). For $\tilde{m} < 0.1$, the correction is not very relevant, and falls below the typical noise level.

The small differences between k_{TL}^{LP} and k_{CP}^{LP} reported in Figure 8 could be due, among other factors, to errors in the determination of the deflection sensitivity of the CPs, as discussed below in Section 3.6, in the determination of the loading point, as well as in the possible rigidification effect of the cantilever, due to the attachment of the sphere, which could influence the dynamics of the last part of the cantilever.

3.6 | Deflection sensitivity and CPs

When big spheres are attached to the cantilever, the standard approach for the calibration of the deflection sensitivity S_z (described in Section 2) can be inaccurate, due to phenomena occurring during the loading phase of the force curve.

In particular, the cantilever tilt angle θ towards the surface may determine a tangential force component, which is due to friction, between the sphere and the substrate. As in the case of normal forces,^{37,38} also tangential friction forces applied at the loading point of the cantilever induce a torque^{53,65,66,71-73}; this torque in turn induces a modification of the cantilever curvature that impacts on the measurement of S_z .⁷²⁻⁷⁴

The friction-related effect is stronger in the high-friction limit, when the probe is pinned on the surface instead of sliding.⁷⁵ The friction, and therefore the torque, is stronger when the effective tip height is larger, as in the case of large CPs, and when the normal force is higher, as in the case stiffer cantilevers are used. Noticeably, for mechanical measurements stiffer cantilevers can be used, in combination with large spheres, in order to achieve reasonably high indentations.

Since the AFM optical lever detection scheme is sensitive to the inclination of the cantilever rather than to its deflection, it turns out that the measurement of S_z can be particularly inaccurate for large CPs, especially in conditions of strong adhesion and friction (as in air). In addition, larger adhesion related to the larger radius of CPs in general makes the acquisition of force curves in air, but also in liquid, critical.⁷⁶

The direct calibration of the deflection sensitivity of large CPs by the standard method based on the acquisition of force curves can therefore be problematic. In this case, alternative approaches should be adopted. One possibility is to consider the mean of the slope the loading and unloading portion of the force curve to effectively subtract the deflection artefact, and then inverting the mean slope to obtain the invOLS, as proposed by Chung et al.⁷⁶ The same authors proposed an electrostatic method to calibrate the deflection

sensitivity, which however requires to build a dedicated micro-device and the metallization of the CP.⁷⁶ In the following sections, we will propose an alternative method to overcome these limitations.

4 | BEST PRACTICES FOR THE CALIBRATION OF (LARGE) CPS

In summary, the thermal noise method can be applied directly to the CP, according to the following rules:

1. S_z is measured carefully, neglecting the initial region of the FC after the contact point, as the inverse of the mean of the slopes of the loading and unloading curves, to minimize friction-related effects, as proposed by Chung et al.⁷⁶
2. The deflection signal of the cantilever is sampled over time with sufficient resolution and number of points, in order to obtain, when applying the thermal noise method, a well-defined peak in the Fourier space; this will allow to accurately estimate the power P_V , despite the typical peak sharpness in air. Alternatively, the thermal noise calibration can be carried out in water, in order to obtain a wider resonance peak. In this case, also the deflection sensitivity must be characterized in water.
3. The β -factor correction (Equation 11) must be applied to the intrinsic spring constant calculated by Equation (7). To this purpose, the sphere radius, the cantilever dimensions, and therefore the reduced mass \tilde{m} and the reduced gyration radius \tilde{r} , must be accurately determined.
4. The corrected intrinsic spring constant k_{CP} is transformed into the apparent spring constant, including also the effect of the shift of the loading point, using Equation (10) (with k_{CP} in place of k_{TL}).

According to the results reported in the present study, we suggest an alternative accurate and reliable approach for the calibration of CPs, and especially of large CPs. This approach can be applied when the CP is produced in house, using tipless cantilevers. In this case, the following procedure can be followed:

1. The intrinsic spring constant k_{TL} of the tipless cantilever, before the attachment of the sphere, is calibrated by the thermal noise method, measuring the deflection sensitivity in the standard way, that is, from a FC acquired on a hard substrate, or using a deflection sensitivity-free approach, like LDV (and thermal noise)⁵⁶ or the Sader method.^{45,54,77}
2. Once the sphere is attached, measure the position of the loading point using an optical microscope and correct the intrinsic spring constant k_{TL} into the apparent $k_{CP,app}^{LP}$, using Equation (10).

It must be considered that, while the interferometric approaches can be used, together with Equation (11), to calibrate CPs (in fact, LDV is a technique of choice for calibrating the spring constant of

AFM probes, but at present it relies on expensive instrumentation), the applicability of the global calibration initiative launched by Sader⁴⁵ should be checked. This approach is based indeed on the statistical assessment of the hydrodynamic function for a given cantilever geometry and may not account for the contribution of the large attached sphere. The general Sader method for the calibration of cantilever of arbitrary shape and dimensions⁵⁴ could in turn work well, provided the calibration of the hydrodynamic function is performed in house.

Once the intrinsic spring constant k_{CP} of a CP has been accurately calibrated, the deflection sensitivity S_z of the current experiment can be calculated assuming that the thermal noise method, through Equation (7), must provide the known value of k_{CP} (see References 78 and 79 and the SNAP procedure⁸).

5 | CONCLUSIONS

Colloidal probes are used in a wide range of experimental conditions. Their dynamical behaviour presents peculiar aspects,²⁷ which also pose issues for their calibration, since the modal shapes are significantly affected.

In this work, we have demonstrated that the addition of a large mass to a tipless cantilever concentrated at its end, besides significantly decreasing the resonant frequency, boosts the quality factor and the amplitude of the first resonant peak, so that a large CP mimics a single-mode mass-on-a-spring system better than a standard AFM probe. These findings may have an impact on the application of CPs in spectroscopic dynamical applications.

Despite the relevant modifications of the cantilever dynamics of CPs, the accuracy of the calibration of CPs is not severely affected until the added mass of the sphere becomes comparable to that of the cantilever, which typically happens for diameters of the glass sphere of about 10 μm .

In the case of large CPs, it is possible to accurately calibrate the spring constant by applying a correction factor (Equation 10) to the value obtained according to the equipartition theorem for an ideal tipless rectangular cantilever. We present two alternative detailed procedures, including one allowing to minimize the impact of the inaccurate determination of the deflection sensitivity.

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CONFLICT OF INTEREST

The authors declare there are no conflicts of interests.

AUTHOR CONTRIBUTIONS

All authors have read and approved the final manuscript. Alessandro Podestà and Ludovic Bellon designed the research plan and contributed essential reagents, materials and tools. Luca Puricelli and Matteo Chighizola carried out the experimental activity and analysed the data with the support of Alessandro Podestà and Ludovic Bellon. Alessandro Podestà, Matteo Chighizola and Ludovic Bellon wrote and revised the paper.

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REFERENCES

- Binnig G, Quate CF, Gerber C. *Atomic force microscope*. *Phys Rev Lett*. 1986;56(9):930-933. <https://doi.org/10.1103/PhysRevLett.56.930>.
- Martin Y, Williams CC, Wickramasinghe HK. *Atomic force microscope-force mapping and profiling on a sub 100-Å scale*. *J Appl Phys*. 1987;61(10):4723-4729. <https://doi.org/10.1063/1.338807>.
- Gould SAC, Drake B, Prater CB, et al. *The atomic force microscope: a tool for science and industry*. *Ultramicroscopy*. 1990;33(2):93-98. [https://doi.org/10.1016/0304-3991\(90\)90011-A](https://doi.org/10.1016/0304-3991(90)90011-A).
- Ducker WA, Senden TJ, Pashley RM. *Direct measurement of colloidal forces using an atomic force microscope*. *Nature*. 1991;353(6341):239-241. <https://doi.org/10.1038/353239a0>.
- Neto C, Craig VSJ. *Colloid probe characterization: radius and roughness determination*. *Langmuir*. 2001;17(7):2097-2099. <https://doi.org/10.1021/la001506y>.
- Craig VSJ, Neto C. *In situ calibration of colloid probe cantilevers in force microscopy: hydrodynamic drag on a sphere approaching a wall*. *Langmuir*. 2001;17(19):6018-6022. <https://doi.org/10.1021/la010424m>.
- Domke J, Radmacher M. *Measuring the elastic properties of thin polymer films with the atomic force microscope*. *Langmuir*. 1998;14(12):3320-3325. <https://doi.org/10.1021/la9713006>.
- Schillers H, Rianna C, Schäpe J, et al. *Standardized nanomechanical atomic force microscopy procedure (SNAP) for measuring soft and biological samples*. *Sci Rep*. 2017;7:5117. <https://doi.org/10.1038/s41598-017-05383-0>.
- Puricelli L, Galluzzi M, Schulte C, Podestà A, Milani P. *Nanomechanical and topographical imaging of living cells by atomic force microscopy with colloidal probes*. *Rev Sci Instrum*. 2015;86(3):33705. <https://doi.org/10.1063/1.4915896>.
- Carl P, Schillers H. *Elasticity measurement of living cells with an atomic force microscope: data acquisition and processing*. *Pflügers Arch Eur J Physiol*. 2008;457(2):551-559. <https://doi.org/10.1007/s00424-008-0524-3>.
- Guz N, Dokukin M, Kalparathi V, Sokolov I. *If cell mechanics can be described by elastic modulus: study of different models and probes used in indentation experiments*. *Biophys J*. 2014;107(3):564-575. <https://doi.org/10.1016/j.bpj.2014.06.033>.
- Sokolov I, Iyer S, Subba-Rao V, Gaikwad RM, Woodworth CD. *Detection of surface brush on biological cells in vitro with atomic force microscopy*. *Appl Phys Lett*. 2007;91(2):23902. <https://doi.org/10.1063/1.2757104>.
- Drelich J, Long J, Xu Z, et al. *AFM colloidal forces measured between microscopic probes and flat substrates in nanoparticle suspensions*. *J Colloid Interface Sci*. 2006;301(2):511-522. <https://doi.org/10.1016/j.jcis.2006.05.044>.
- Fischer-Cripps AC. *The Hertzian contact surface*. *J Mater Sci*. 1999;34(1):129-137. <https://doi.org/10.1023/A:1004490230078>.
- Harding JW, Sneddon IN. *The elastic stresses produced by the indentation of the plane surface of a semi-infinite elastic solid by a rigid punch*. *Math Proc Cambridge Philos Soc*. 1945;41(1):16-26. <https://doi.org/10.1017/S0305004100022325>.
- Nebuloni M, Albarello L, Andolfo A, et al. *Insight on colorectal carcinoma infiltration by studying perilesional extracellular matrix*. *Sci Rep*. 2016;6:22522. <https://doi.org/10.1038/srep22522>.
- Alcaraz J, Buscemi L, Grabulosa M, et al. *Microrheology of human lung epithelial cells measured by atomic force microscopy*. *Biophys J*. 2003;84(3):2071-2079. [https://doi.org/10.1016/S0006-3495\(03\)75014-0](https://doi.org/10.1016/S0006-3495(03)75014-0).
- Luque T, Melo E, Garreta E, et al. *Local micromechanical properties of decellularized lung scaffolds measured with atomic force microscopy*. *Acta Biomater*. 2013;9(6):6852-6859. <https://doi.org/10.1016/j.actbio.2013.02.044>.
- Beekmans SV, Iannuzzi D. *A metrological approach for the calibration of force transducers with interferometric readout*. *Surf Topogr Metrol Prop*. 2015;3(2):25004. <https://doi.org/10.1088/2051-672X/3/2/025004>.
- Chavan D, van de Watering TC, Gruca G, et al. *Ferrule-top nanoindenter: an optomechanical fiber sensor for nanoindentation*. *Rev Sci Instrum*. 2012;83(11):115110. <https://doi.org/10.1063/1.4766959>.
- Hoon H, Kurniawan N, Koenderink G, Iannuzzi D. *Local dynamic mechanical analysis for heterogeneous soft matter using ferrule-top indentation*. *Soft Matter*. 2016;12(12):3066-3073. <https://doi.org/10.1039/C6SM00300A>.
- Chighizola M, Puricelli L, Osteroom LV, Antonovaite N, Podestà A. *The measurement of the Young's modulus of elasticity of gels and soft elastomers using atomic force microscopy and a dedicated nanoindenter: a comparative study*. In preparation; 2020.
- Swadener JG, George EP, Pharr GM. *The correlation of the indentation size effect measured with indenters of various shapes*. *J Mech Phys Solids*. 2002;50(4):681-694. [https://doi.org/10.1016/S0022-5096\(01\)00103-X](https://doi.org/10.1016/S0022-5096(01)00103-X).
- Alcalá J, Giannakopoulos AE, Suresh S. *Continuous measurements of load-penetration curves with spherical microindenters and the estimation of mechanical properties*. *J Mater Res*. 1998;13(5):1390-1400. <https://doi.org/10.1557/JMR.1998.0197>.
- McBride SP, Law BM. *Improved in situ spring constant calibration for colloidal probe atomic force microscopy*. *Rev Sci Instrum*. 2010;81(11):113703. <https://doi.org/10.1063/1.3502460>.
- Butt H-J, Cappella B, Kappl M. *Force measurements with the atomic force microscope: technique, interpretation and applications*. *Surf Sci Rep*. 2005;59(1-6):1-152. <https://doi.org/10.1016/j.surfrep.2005.08.003>.
- Laurent J, Steinberger A, Bellon L. *Functionalized AFM probes for force spectroscopy: eigenmodes shape and stiffness calibration through thermal noise measurements*. *Nanotechnology*. 2013;24(22):225504. <https://doi.org/10.1088/0957-4484/24/22/225504>.
- Indrieri M, Podestà A, Bongiorno G, Marchesi D, Milani P. *Adhesive-free colloidal probes for nanoscale force measurements: production and characterization*. *Rev Sci Instrum*. 2011;82(2):23708. <https://doi.org/10.1063/1.3553499>.
- Markiewicz P, Goh MC. *Atomic force microscope tip deconvolution using calibration arrays*. *Rev Sci Instrum*. 1995;66(5):3186-3190. <https://doi.org/10.1063/1.1145549>.
- Montelius L, Tegenfeldt JO. *Direct observation of the tip shape in scanning probe microscopy*. *Appl Phys Lett*. 1993;62(21):2628-2630. <https://doi.org/10.1063/1.109267>.
- Bykov V, Gologanov A, Shevyakov V. *Test structure for SPM tip shape deconvolution*. *Appl Phys A Mater Sci Process*. 1998;66(5):499-502. <https://doi.org/10.1007/s003390050703>.

32. Villarrubia JS. Morphological estimation of tip geometry for scanned probe microscopy. *Surf Sci.* 1994;321:287-300. [https://doi.org/10.1016/0039-6028\(94\)90194-5](https://doi.org/10.1016/0039-6028(94)90194-5).
33. Keller D. Reconstruction of STM and AFM images distorted by finite-size tips. *Surf Sci.* 1991;253(1-3):353-364. [https://doi.org/10.1016/0039-6028\(91\)90606-S](https://doi.org/10.1016/0039-6028(91)90606-S).
34. Sader JE, Lu J, Mulvaney P. Effect of cantilever geometry on the optical lever sensitivities and thermal noise method of the atomic force microscope. *Rev Sci Instrum.* 2014;85(11):113702. <https://doi.org/10.1063/1.4900864>.
35. Schäffer TE. Calculation of thermal noise in an atomic force microscope with a finite optical spot size. *Nanotechnology.* 2005;16(6):664-670. <https://doi.org/10.1088/0957-4484/16/6/007>.
36. Evans DR, Craig VSJ. Sensing cantilever beam bending by the optical lever technique and its application to surface stress. *J Phys Chem B.* 2006;110(11):5450-5461. <https://doi.org/10.1021/jp0536807>.
37. Edwards SA, Ducker WA, Sader JE. Influence of atomic force microscope cantilever tilt and induced torque on force measurements. *J Appl Phys.* 2008;103(6):64513. <https://doi.org/10.1063/1.2885734>.
38. Hutter JL. Comment on tilt of atomic force microscope cantilevers: effect on spring constant and adhesion measurements. *Langmuir.* 2005;21(6):2630-2632. <https://doi.org/10.1021/la047670t>.
39. Gates RS. Experimental confirmation of the atomic force microscope cantilever stiffness tilt correction. *Rev Sci Instrum.* 2017;88(12):123710. <https://doi.org/10.1063/1.4986201>.
40. Heim L-O, Rodrigues TS, Bonaccorso E. Direct thermal noise calibration of colloidal probe cantilevers. *Colloids Surfaces A Physicochem Eng Asp.* 2014;443:377-383. <https://doi.org/10.1016/j.colsurfa.2013.11.018>.
41. Hutter JL, Bechhoefer J. Calibration of atomic-force microscope tips. *Rev Sci Instrum.* 1993;64(7):1868-1873. <https://doi.org/10.1063/1.1143970>.
42. Cleveland JP, Manne S, Bocek D, Hansma PK. A nondestructive method for determining the spring constant of cantilevers for scanning force microscopy. *Rev Sci Instrum.* 1993;64(2):403-405. <https://doi.org/10.1063/1.1144209>.
43. Reitsma MG. Lateral force microscope calibration using a modified atomic force microscope cantilever. *Rev Sci Instrum.* 2007;78(10):106102. <https://doi.org/10.1063/1.2789653>.
44. Kim M-S, Choi J-H, Kim J-H, Park Y-K. SI-traceable determination of spring constants of various atomic force microscope cantilevers with a small uncertainty of 1%. *Meas Sci Technol.* 2007;18(11):3351-3358. <https://doi.org/10.1088/0957-0233/18/11/014>.
45. Sader JE, Borgani R, Gibson CT, et al. A virtual instrument to standardise the calibration of atomic force microscope cantilevers. *Rev Sci Instrum.* 2016;87(9):93711. <https://doi.org/10.1063/1.4962866>.
46. Butt H-J, Jaschke M. Calculation of thermal noise in atomic force microscopy. *Nanotechnology.* 1995;6(1):1-7. <https://doi.org/10.1088/0957-4484/6/1/001>.
47. Gates RS, Pratt JR. Accurate and precise calibration of AFM cantilever spring constants using laser Doppler vibrometry. *Nanotechnology.* 2012;23(37):375702. <https://doi.org/10.1088/0957-4484/23/37/375702>.
48. Cook SM, Lang KM, Chynoweth KM, Wigton M, Simmonds RW, Schäffer TE. Practical implementation of dynamic methods for measuring atomic force microscope cantilever spring constants. *Nanotechnology.* 2006;17(9):2135-2145. <https://doi.org/10.1088/0957-4484/17/9/010>.
49. Proksch R, Schäffer TE, Cleveland JP, Callahan RC, Viani MB. Finite optical spot size and position corrections in thermal spring constant calibration. *Nanotechnology.* 2004;15(9):1344-1350. <https://doi.org/10.1088/0957-4484/15/9/039>.
50. Schäffer TE, Fuchs H. Optimized detection of normal vibration modes of atomic force microscope cantilevers with the optical beam deflection method. *J Appl Phys.* 2005;97(8):83524. <https://doi.org/10.1063/1.1872202>.
51. Sader JE, White L. Theoretical analysis of the static deflection of plates for atomic force microscope applications. *J Appl Phys.* 1993;74(1):1-9. <https://doi.org/10.1063/1.354137>.
52. Sader JE, Larson I, Mulvaney P, White LR. Method for the calibration of atomic force microscope cantilevers. *Rev Sci Instrum.* 1995;66(7):3789-3798. <https://doi.org/10.1063/1.1145439>.
53. Wang F. Comment on "influence of atomic force microscope cantilever tilt and induced torque on force measurements" [*J. Appl. Phys.* 103, 064513 (2008)]. *J Appl Phys.* 2009;106(9):96103. <https://doi.org/10.1063/1.3257268>.
54. Sader JE, Sanelli JA, Adamson BD, et al. Spring constant calibration of atomic force microscope cantilevers of arbitrary shape. *Review of Scientific Instruments.* 2012;83:103705. <https://doi.org/10.1063/1.4757398>.
55. Gates RS, Osborn WA, Shaw GA. Accurate flexural spring constant calibration of colloid probe cantilevers using scanning laser Doppler vibrometry. *Nanotechnology.* 2015;26(23):235704. <https://doi.org/10.1088/0957-4484/26/23/235704>.
56. Ohler B. Cantilever spring constant calibration using laser Doppler vibrometry. *Rev Sci Instrum.* 2007;78(6):63701. <https://doi.org/10.1063/1.2743272>.
57. Paolino P, Aguilar Sandoval FA, Bellon L. Quadrature phase interferometer for high resolution force spectroscopy. *Rev Sci Instrum.* 2013;84(9):95001. <https://doi.org/10.1063/1.4819743>.
58. Gates RS, Osborn WA, Pratt JR. Experimental determination of mode correction factors for thermal method spring constant calibration of AFM cantilevers using laser Doppler vibrometry. *Nanotechnology.* 2013;24(25):255706. <https://doi.org/10.1088/0957-4484/24/25/255706>.
59. Mescola A, Vella S, Scotto M, et al. Probing cytoskeleton organisation of neuroblastoma cells with single-cell force spectroscopy. *J Mol Recognit JMR.* 2012;25(5):270-277. <https://doi.org/10.1002/jmr.2173>.
60. Beicker K, O'Brien ET, Falvo MR, Superfine R. Vertical light sheet enhanced side-view imaging for AFM cell mechanics studies. *Sci Rep.* 2018;8(1):1-12. <https://doi.org/10.1038/s41598-018-19791-3>.
61. Marsh G, Waugh RE. Quantifying the mechanical properties of the endothelial Glycocalyx with atomic force microscopy. *JoVE J Vis Exp.* 2013;(72):e50163. <https://doi.org/10.3791/50163>.
62. Kauppi A, Andersson KM, Bergström L. Probing the effect of superplasticizer adsorption on the surface forces using the colloidal probe AFM technique. *Cem Concr Res.* 2005;35(1):133-140. <https://doi.org/10.1016/j.cemconres.2004.07.008>.
63. Weadock N, Varongchayakul N, Wan J, Lee S, Seog J, Hu L. Determination of mechanical properties of the SEI in sodium ion batteries via colloidal probe microscopy. *Nano Energy.* 2013;2(5):713-719. <https://doi.org/10.1016/j.nanoen.2013.08.005>.
64. Delcea M, Schmidt S, Palankar R, et al. Mechanobiology: correlation between mechanical stability of microcapsules studied by AFM and impact of cell-induced stresses. *Small.* 2010;6(24):2858-2862. <https://doi.org/10.1002/sml.201001478>.
65. Chung K-H, Pratt JR, Reitsma MG. Lateral force calibration: accurate procedures for colloidal probe friction measurements in atomic force microscopy. *Langmuir.* 2010;26(2):1386-1394. <https://doi.org/10.1021/la902488r>.
66. Dohn S, Svendsen W, Boisen A, Hansen O. Mass and position determination of attached particles on cantilever based mass sensors. *Rev Sci Instrum.* 2007;78(10):103303. <https://doi.org/10.1063/1.2804074>.
67. Hoof S, Nand Gosvami N, Hoogenboom BW. Enhanced quality factors and force sensitivity by attaching magnetic beads to cantilevers for atomic force microscopy in liquid. *J Appl Phys.* 2012;112(11):114324. <https://doi.org/10.1063/1.4768713>.
68. Basak S, Raman A, Garimella SV. Hydrodynamic loading of micro-cantilevers vibrating in viscous fluids. *J Appl Phys.* 2006;99(11):114906. <https://doi.org/10.1063/1.2202232>.
69. Maali A, Hurth C, Boisgard R, Jai C, Cohen-Bouhacina T, Aimé J-P. Hydrodynamics of oscillating atomic force microscopy cantilevers in

- viscous fluids. *J Appl Phys.* 2005;97(7):74907. <https://doi.org/10.1063/1.1873060>.
70. Janovjak H, Struckmeier J, Müller DJ. *Hydrodynamic effects in fast AFM single-molecule force measurements.* *Eur Biophys J.* 2005;34(1):91-96. <https://doi.org/10.1007/s00249-004-0430-3>.
71. Warmack RJ, Zheng X-Y, Thundat T, Allison DP. *Friction effects in the deflection of atomic force microscope cantilevers.* *Rev Sci Instrum.* 1994;65(2):394-399. <https://doi.org/10.1063/1.1145144>.
72. Stiernstedt J, Rutland MW, Attard P. *A novel technique for the in situ calibration and measurement of friction with the atomic force microscope.* *Rev Sci Instrum.* 2005;76(8):83710. <https://doi.org/10.1063/1.2006407>.
73. Stiernstedt J, Rutland MW, Attard P. *Erratum: "A novel technique for the in situ calibration and measurement of friction with the atomic force microscope" [Rev. Sci. Instrum. 76, 083710 (2005)].* *Rev Sci Instrum.* 2006;77(1):19901. <https://doi.org/10.1063/1.2162429>.
74. Attard P, Carambassis A, Rutland MW. *Dynamic surface force measurement. 2. Friction and the atomic force microscope.* *Langmuir.* 1999;15(2):553-563. <https://doi.org/10.1021/la980848p>.
75. Weafer PP, McGarry JP, van Es MH, et al. *Stability enhancement of an atomic force microscope for long-term force measurement including cantilever modification for whole cell deformation.* *Rev Sci Instrum.* 2012;83(9):93709. <https://doi.org/10.1063/1.4752023>.
76. Chung K-H, Shaw GA, Pratt JR. *Accurate noncontact calibration of colloidal probe sensitivities in atomic force microscopy.* *Rev Sci Instrum.* 2009;80(6):65107. <https://doi.org/10.1063/1.3152335>.
77. Sader JE, Chon JWM, Mulvaney P. *Calibration of rectangular atomic force microscope cantilevers.* *Rev Sci Instrum.* 1999;70(10):3967-3969. <https://doi.org/10.1063/1.1150021>.
78. Higgins MJ, Proksch R, Sader JE, et al. *Noninvasive determination of optical lever sensitivity in atomic force microscopy.* *Rev Sci Instrum.* 2006;77(1):1-5. <https://doi.org/10.1063/1.2162455>.
79. Rico F, Sumbul F, RODRIGUEZ RAMOS J, Hassanpour N. *One-step calibration of AFM in liquid.* *Front Phys.* 2020;8:301. <https://doi.org/10.3389/FPHY.2020.00301>.

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