Nanotechnology 24 (2013) 225504 (13pp)

Functionalized AFM probes for force spectroscopy: eigenmode shapes and stiffness calibration through thermal noise measurements

Justine Laurent, Audrey Steinberger and Ludovic Bellon

Université de Lyon, Laboratoire de Physique, École Normale Supérieure de Lyon, CNRS, 46 allée d'Italie, F-69007, Lyon, France

E-mail: Ludovic.Bellon@ens-lyon.fr

Received 23 January 2013, in final form 6 March 2013 Published 3 May 2013 Online at stacks.iop.org/Nano/24/225504

Abstract

The functionalization of an atomic force microscope (AFM) cantilever with a colloidal bead is a widely used technique when the geometry between the probe and the sample must be controlled, particularly in force spectroscopy. But some questions remain: how does a bead glued at the end of a cantilever influence its mechanical response? And more importantly for quantitative measurements, can we still determine the stiffness of the AFM probe with traditional techniques?

In this paper, the influence of the colloidal mass loading on the eigenmode shape and resonant frequency is investigated by measuring the thermal noise on rectangular AFM microcantilevers with and without beads attached at their extremities. The experiments are performed with a home-made ultra-sensitive AFM, based on differential interferometry. The focused beam from the interferometer probes the cantilever at different positions and the spatial shapes of the modes are determined up to the fifth resonance, without external excitation. The results clearly demonstrate that the first eigenmode is almost unchanged by mass loading. However the oscillation behavior of higher resonances presents a marked difference: with a particle glued at its extremity, the nodes of the modes are displaced towards the free end of the cantilever. These results are compared to an analytical model taking into account the mass and inertial moment of the load in an Euler–Bernoulli framework, where the normalization of the eigenmodes is explicitly worked out in order to allow a quantitative prediction of the thermal noise amplitude of each mode. A good agreement between the experimental results and the analytical model is demonstrated, allowing a clean calibration of the probe stiffness.

S Online supplementary data available from stacks.iop.org/Nano/24/225504/mmedia

(Some figures may appear in colour only in the online journal)

1. Introduction

Atomic force microscopes (AFMs) are currently used in a great variety of studies in various disciplines to measure small forces by detecting the deflection of a microcantilever [1]. In biophysics, for example, it has been applied to the

unfolding of proteins [2, 3], probing the structure of biological membranes [4], and monitoring the mechanical response of living cells [5, 6]. Also in nanotechnology, micro-scale levers find applications in micro-electro-mechanical systems (MEMS) and other nanotechnological devices [7]. In material, surface or nanosciences in general, AFM probes appear

as a cornerstone for quantitative studies of nanoscale properties [8].

All these applications exploit the great accuracy in measuring the cantilever deflection offered by AFM, converting this measurement into units of force by assuming the cantilever behaves like a spring of known stiffness. Manufacturers often specify the spring constant of their cantilevers with a wide range of values, mainly because of the great uncertainties in the dimensions, particularly the thickness, resulting from the fabrication process. To overcome this problem several techniques have been proposed to calibrate the cantilever spring constant [1, 9–12]. The reader is referred to the work of Burnham *et al* [13] and the references therein for a comparative summary of the different techniques.

One of the first and still most commonly used calibration methods is the so-called thermal calibration method, based on the measurement of the vibration amplitude of the free end of a cantilever excited due to thermal noise [11]. The first peak of the thermal noise spectrum is related back to the spring constant of the cantilever modeled as an harmonic oscillator. In a more accurate model, Butt and Jaschke [14] introduced a correction factor deduced from the Euler–Bernoulli description of the flexural dynamics of a free-clamped beam. In a previous work, we demonstrated that measuring this thermal noise for the first resonant mode of the cantilever provides an excellent benchmark to probe the mechanical response of the cantilever and compare it to a simple mechanical model [15].

In this paper, we extend this method to the case of functionalized AFM cantilevers. Indeed, when the geometry between the probe and the sample must be controlled, it is common to use a colloidal bead fixed at the free end of the lever. The radius of curvature of the 'tip' is then controlled and stable, and offers a clean sphere–plane geometry to study interactions at nanometric distances. These modified probes (whether home-made or commercially available) are commonly used in force spectroscopy, in particular for the measurement of the nanorheology of confined fluids [16, 17] or of the Casimir interaction [18]. How does this loaded mass influence the mechanical response of the AFM cantilever? Can the common techniques (thermal noise calibration in particular) to determine the stiffness still be used?

In this work, we measure the thermal noise spectra of the cantilever deflection on its whole surface and compare the rms amplitudes obtained with and without a bead loaded at its free extremity. Furthermore, we compare the results to a simple mass model that modifies only the boundary conditions of the classical rectangular beam theory [19]. A good agreement between experimental data and this analytical model will be demonstrated, showing that thermal noise calibration of the probe stiffness is still perfectly pertinent for such cantilevers.

The paper is organized as follows. Section 2 describes the theoretical approach, with a special emphasis on eigenmode normalization to allow the prediction of the thermal noise amplitude of each mode. Section 3 details the experimental results in the light of this model, for two cantilevers probing various mass ratios between the cantilever and the colloidal



Figure 1. Schematics of the experiment and analytical model. The cantilever (length *L*, width *W*, thickness *H*) is modeled in the Euler–Bernoulli framework by its neutral axis subject to a deflection Z(x, t). The bead glued at its free end is modeled as a point mass m_b located at the extremity. r_g is the equivalent gyration radius of the bead and accounts for the inertia in rotation of the bead at the cantilever end. In the experiment, the deflection can be measured at any position *x* and *y* (along and transverse to the cantilever axis) with a very low noise differential interferometer [15, 21, 22], sensing the optical path difference between the two laser beams represented in the figure.

bead. A discussion and conclusions are given in section 4, with a specific focus on how our conclusions can be applied to the classic angular deflection measurement technique.

2. Analytical description of thermal noise

2.1. Flexural eigenmodes of a clamped cantilever

In the Euler-Bernoulli framework to describe the micrometersized mechanical beam, we assume that the cantilever length L is much larger than its width W, which itself is much larger than its thickness T (see figure 1). The flexural modes of the cantilever are supposed to be only perpendicular to its length and uniform across its width. The deformations can thus be described by the deflection Z(x, t), with t the time and x the spatial coordinate along the beam normalized to its length L. The bead is supposed to be nondeformable, and thus described as a rigid mass $m_{\rm b}$ at the free end of the cantilever. It may be offset from the neutral axis and carry a non-negligible inertia: these effects are taken into account with an inertial moment $m_{\rm b}r_{\rm g}^2$, $r_{\rm g}$ being the equivalent gyration radius computed at the free end of the cantilever. For a sphere of radius r, r_g is equal to $\sqrt{7/5}r$. We neglect in our analysis any offset along the cantilever axis and any coupling with torsion: precise gluing of the bead should make those effects negligible in a first approximation. In order to reach a simple analytical solution, we also neglect the actual triangular shape of the cantilever at its free end: we will consider that the lever has an effective length smaller than its total length to approximate the real geometry. This last approximation will prove to be reasonable to describe our measurements. Figure 1 sketches the experiment and the applied model. Following [19], we will include the effect of the bead in the boundary conditions of the cantilever dynamics.

The equation of motion for the cantilever, once the variables in time and space are separated, can be written as

$$\frac{k}{3}\frac{\mathrm{d}^4 z}{\mathrm{d}x^4} = m_\mathrm{c}\omega^2 z \tag{1}$$

with k the static stiffness of the cantilever, m_c its mass, $Z(x, t) = z(x)e^{i\omega t}$ the deflection, and ω the angular frequency. This equation can be rewritten as

$$z^{(4)} = \alpha^4 z \tag{2}$$

where $^{(n)}$ is the spatial derivative of order *n*, and α is given by the dispersion relation:

$$\alpha^4 = \frac{3m_c\omega^2}{k}.$$
 (3)

The generic solution to this equation is

$$z(x) = a\cos(\alpha x) + b\sin(\alpha x) - c\cosh(\alpha x) - d\sinh(\alpha x).$$
(4)

The boundary conditions in x = 0 correspond to a clamped end, implying z(0) = 0 and $z^{(1)}(0) = 0$, hence a = c and b = d. Defining R = b/a, expression (4) can thus be written as

$$z(x) = a(\cos(\alpha x) - \cosh(\alpha x) + R[\sin(\alpha x) - \sinh(\alpha x)]).$$
(5)

For x = 1, corresponding to the free end of the cantilever where the bead is glued, the conditions on the force and torque are linked to the inertia in translation and rotation of the bead [19]:

$$z^{(3)}(1) = -\alpha^4 \tilde{m} z(1) \tag{6}$$

$$z^{(2)}(1) = \alpha^4 \tilde{m} \tilde{r}^2 z^{(1)}(1) \tag{7}$$

where $\tilde{m} = m_b/m_c$ is the mass of the bead m_b normalized to that of the cantilever m_c , and $\tilde{r} = r_g/L$ is the gyration radius of the bead r_g normalized to the cantilever length *L*. Expressing these boundary conditions with expression (5) leads to

$$R = \frac{\sin \alpha - \sinh \alpha + \alpha \tilde{m}(\cos \alpha - \cosh \alpha)}{\cos \alpha + \cosh \alpha - \alpha \tilde{m}(\sin \alpha - \sinh \alpha)}$$
(8)

$$= -\frac{\cos\alpha + \cosh\alpha - \alpha^3 \tilde{m}\tilde{r}^2(\sin\alpha + \sinh\alpha)}{\sin\alpha + \sinh\alpha + \alpha^3 \tilde{m}\tilde{r}^2(\cos\alpha - \cosh\alpha)}.$$
 (9)

The values of α allowing this equality are quantified, and correspond to the spatial eigenvalues $\alpha_n(\tilde{m}, \tilde{r})$ of the resonant modes of the cantilever. They can be numerically computed. For $\tilde{m} = 0$ (no bead), the last equation simplifies to the usual condition $1 + \cos \alpha \cosh \alpha = 0$, leading to the common



Figure 2. Plot showing the first five normal modes $\phi_n(x)$ for various \tilde{m} and \tilde{r} . The amplitude of the first mode decreases and the nodes of the higher order modes get closer to the free end when either \tilde{m} or \tilde{r} increases.

tabulated eigenvalues of a clamped–free Euler–Bernoulli mechanical beam. The $\alpha_n(\tilde{m}, \tilde{r})$ values are reported for the first five modes, for $0 \le \tilde{m} \le 2$ and $0 \le \tilde{r} \le 0.1$, in tables I–V in the supplementary data (available at stacks.iop.org/Nano/24/225504/mmedia). The corresponding shapes of the eigenmodes are plotted in figure 2.

The length of the cantilever *L* used in the normalization of *x*, and thus impacting the spatial eigenvalues α_n , is sometimes experimentally ill-defined due to the triangular shape of the cantilever end. The direct comparison of the experimental values of α_n with the theoretical ones is thus hampered by this incertitude. However, their ratio is free from this bias and can be used to check analytical predictions. In the appendix, we

plot such ratios, useful to extract the values of \tilde{m} and \tilde{r} from the experimental observations.

2.2. Orthogonality relations and normalization

The eigenmodes $z_n(x)$ are given by equation (5), with a dependence of the three parameters a, R and α on mode number n and on \tilde{m} and \tilde{r} (the dependence in \tilde{m} and \tilde{r} will be implicit in our notation):

$$z_n(x) = a_n \zeta_n(x)$$
(10)
= $a_n (\cos(\alpha_n x) - \cosh(\alpha_n x))$

$$+ R_n \left[\sin(\alpha_n x) - \sinh(\alpha_n x) \right] \right). \tag{11}$$

However, if we only consider this expression, then the orthogonality between two modes z_n and z_m does not hold: it is easy to show that for $n \neq m$,

$$\int_{0}^{1} z_{n}(x)z_{m}(x)dx = -\tilde{m}z_{n}(1)z_{m}(1) - \tilde{m}\tilde{r}^{2}z_{n}^{(1)}(1)z_{m}^{(1)}(1).$$
(12)

We therefore define the normal modes by

$$\phi_n(x) = z_n(x) + z_n(1)\sqrt{\tilde{m}\delta(x-1)} + z_n^{(1)}(1)\tilde{r}\sqrt{\tilde{m}\delta(x-1+\epsilon)}$$
(13)

where $\delta(x - 1)$ and $\delta(x - 1 + \epsilon)$ are the Dirac distributions centered on x = 1 and $1 - \epsilon$, with ϵ being an arbitrarily small quantity ensuring only that there is no overlap between the $z_n(1)$ and $z_n^{(1)}(1)$ terms. With such a definition, it is straightforward to prove the orthogonality of the ϕ_n basis. Moreover, as $\phi_n(x) = z_n(x)$ for any $x \in [0, 1 - \epsilon[, \phi_n(x)$ thus obeys the initial differential equation (2) and the boundary conditions in the limit x = 0 and $x \to 1$ (in the limit $\epsilon \to$ 0). Eventually, the normalization of ϕ_n is easily obtained by imposing

$$\int_0^1 \phi_n^2(x) \, \mathrm{d}x = 1 \tag{14}$$

$$a_n^2 \left(\int_0^1 \zeta_n^2(x) \, \mathrm{d}x + \tilde{m}\zeta_n^2(1) + \tilde{m}\tilde{r}^2 \zeta_n^{\prime 2}(1) \right) = 1 \qquad (15)$$

where $\zeta_n(x)$ is defined in equation (10). This last equation thus imposes the values of a_n to construct the orthonormal basis ϕ_n . The result of this process is illustrated in figure 2 for a few values of \tilde{m} and \tilde{r} .

Let us now give an energetic meaning to the Dirac term, in the case $\tilde{r} = 0$. We first compute the kinetic energy of a mode ϕ_n with amplitude A: $Z(x, t) = A\phi_n(x) \cos \omega_n t$. The speed of a mass element $m_c dx$ is $-A\phi_n(x)\omega_n \sin \omega_n t$, thus the total kinetic energy is

$$E_{\rm c} = \int_0^1 \frac{1}{2} m_{\rm c} \, \mathrm{d}x \, A^2 \phi_n^2(x) \omega_n^2 \sin^2 \omega_n t \tag{16}$$

$$= \frac{1}{2}m_{\rm c}\left(\int_0^1 \phi_n^2(x) \,\mathrm{d}x\right) A^2 \omega_n^2 \sin^2 \omega_n t \qquad (17)$$

$$=\frac{1}{2}m_{\rm c}A^2\omega_n^2\sin^2\omega_n t.$$
 (18)

This is the kinetic energy of a harmonic oscillator of mass m_c , resonant angular frequency ω_n and amplitude *A*. Notice that the amplitude of this oscillator is different from the deflection at the free end of the cantilever: $A_c = A\phi_n(x \rightarrow 1) = Az_n(1) = Aa_n\zeta_n(1)$. We may also rewrite the integral on $\phi_n^2(x)$ in equation (17) using equation (15):

$$E_{\rm c} = \frac{1}{2} m_{\rm c} a_n^2 \left(\int_0^1 \zeta_n^2(x) \, \mathrm{d}x + \tilde{m} \zeta_n^2(1) \right) A^2 \omega_n^2 \sin^2 \omega_n t \quad (19)$$

=
$$\int_0^1 \frac{1}{2} m_{\rm c} \, \mathrm{d}x \, A^2 z_n^2(x) \omega_n^2 \sin^2 \omega_n t + \frac{1}{2} m_{\rm b} A_{\rm c}^2 \omega_n^2 \sin^2 \omega_n t. \quad (20)$$

We can easily identify here the sum of two terms: the kinetic energy of the mode *n* of the cantilever itself, subject to a sinusoidal motion with an amplitude A_c at its free end, and the kinetic energy of a point mass m_b , subject to a sinusoidal motion with the same amplitude A_c . The additional Dirac term in ϕ_n thus takes into account the bead motion in the total energy of the equivalent harmonic oscillator. The amplitude of the latter is not equal to the amplitude at the free end of the cantilever (which is also the case without the added mass since $|\phi_n(x = 1)| = 2$ for $\tilde{m} = 0$).

In the case where $\tilde{r} \neq 0$, we can extend this energetic approach to include the kinetic energy due to the rotation of the bead, and retrieve the normalization of the modes. A similar approach has been used by Oguamanam [19] to ensure the orthonormalization of the normal modes in a more general framework including the coupling of flexural and torsional modes.

2.3. Thermal noise amplitude of resonant modes

We compute the thermal noise of each resonant mode following [14, 20]: let us project the thermal noise driven deflection on the orthonormal basis $\phi_n(x)$:

$$Z(x,t) = \sum_{n=1}^{\infty} Z_n(t)\phi_n(x).$$
 (21)

Under the hypothesis of uncoupled modes, we have for each degree of freedom

$$\frac{1}{2}k_n \langle Z_n^2(t) \rangle = \frac{1}{2}k_{\rm B}T \tag{22}$$

where $k_{\rm B}$ is Boltzmann's constant, *T* the temperature of the cantilever, and k_n the stiffness of the mode defined by:

$$k_n = \frac{k}{3}\alpha_n^4 = m_c\omega_n^2.$$
(23)

The mean quadratic deflection measured in x should thus be

7

$$\langle Z^2(x,t)\rangle = \sum_{n=1}^{\infty} \langle Z_n \rangle^2(t) |\phi_n(x)|^2$$
(24)

$$= \frac{k_{\rm B}T}{k} \sum_{n=1}^{\infty} 3 \frac{|\phi_n(x)|^2}{\alpha_n^4}$$
$$= \frac{k_{\rm B}T}{k} \sum_{n=1}^{\infty} \eta_n(x, \tilde{m}, \tilde{r}).$$
(25)



Figure 3. Thermal noise expected for a cantilever of static stiffness $k = 0.1 \text{ N m}^{-1}$ at 300 K for a few values of \tilde{m} and \tilde{r} . The rms deflection of the first five modes is plotted as a function of the position along the cantilever *x*. The first mode is weakly impacted by the bead. When \tilde{m} increases, the amplitudes of the higher order modes vanish close to the free end, and the thermal energy is reported towards the antinodes, the amplitudes of which increase.

In figure 3, we plot the expected rms thermal noise at 300 K along a cantilever of static stiffness k = 0.1 N m⁻¹ for the first five eigenmodes, for a few values of \tilde{m} and \tilde{r} . Note that the normalization of the ϕ_n basis is a crucial step in applying the energy equipartition theorem in a quantitative manner to this analysis. Our approach also allows us to estimate the repartition of energy between the different modes. For example, if we perform the measurement at the free end of the cantilever (x = 1), the first mode accounts for $\eta_1(x = 1, \tilde{m} = 0, \tilde{r} = 0) = 97\%$ of the total thermal fluctuations for $\tilde{m} = \tilde{r} = 0$, and $\eta_1(x = 1, \tilde{m} = 1, \tilde{r} = 0) = 99.8\%$ for $\tilde{m} = 1$ and $\tilde{r} = 0$. In figure 4, we plot the contribution of each mode to



Figure 4. Fraction $\eta_n(x = 1, \tilde{m}, \tilde{r} = 0)$ at the free end of the cantilever of the mean quadratic deflection driven by thermal noise for the first five modes as a function of the normalized bead mass \tilde{m} . The amplitude of the first mode (plotted as $1 - \eta_1$ since $\eta_1 \rightarrow 1$) quickly overcomes the contribution of all the other modes, for which the inertia of the mass implies a node close to the free end of the cantilever.



Figure 5. Pictures from an optical microscope of beads glued at the apex of cantilevers A and B.

the mean quadratic deflection measured at its extremity when \tilde{m} changes (for $\tilde{r} = 0$): the larger the mass, the stronger is the contribution of the first mode.

3. Experimental methodology and results

3.1. Experiment description

Manufacturer specifications of our two samples (A and B) are given in table 1. Both present a 'rectangular' geometry, close to the model used in our analytical approach. However, the triangular end (see figure 5) departs from the model, and impedes a proper definition of their length L. We measure the deflection solely driven by thermal noise over the surface of these two different commercial cantilevers, first when they are still bare, then again after a glass bead has been glued at their free end with a two-part epoxy adhesive (Araldite). The bead of cantilever A is a fused borosilicate glass microsphere from Potters (Sphericel 110P8), whereas the bead of cantilever B is a borosilicate sphere from Cospheric (BSGMS 45–53 m). The radius r of the bead is reported in table 1 for each sample.

As already mentioned, due to the manufacturing process, the uncertainty in the thickness of the lever is large, resulting in a large uncertainty in the computation of its mass. The case is even worse for cantilever B, where the gold coating can significantly change the total mass due to the high density of gold. In addition, the quantity of glue cannot be measured precisely from the images of the cantilever.

Table 1. Manufacturer specifications for cantilevers A and B. The last line corresponds to the glass bead that we glued at the free end of the cantilever, as illustrated in figure 5.

	Cantilever A	Cantilever B		
Manufacturer	BudgetSensors	Nano-World		
Reference	AIO, lever A	Arrow-TL8Au		
Material	Silicon	Silicon		
Tip height	$(17 \pm 2) \ \mu m$	Tipless		
Coating	None	Ti 5 nm + Au 30 nm		
Resonant frequency	$(15 \pm 5) \text{ kHz}$	6 kHz		
1 2	. ,	(3–14) kHz		
Force constant k	$0.2 \text{ N} \text{ m}^{-1}$	$0.03 \text{ N} \text{ m}^{-1}$		
	(0.04-0.7) N m ⁻¹	(0.004-0.54) N m ⁻¹		
Length L	$(500 \pm 10) \mu m$	$(500 \pm 5) \mu m$		
Width W	$(30 \pm 5) \mu m$	$(100 \pm 5) \mu m$		
Thickness T	$(2.7 \pm 1.0) \ \mu m$	$1.0 \mu \mathrm{m}$		
		$(0.5-2.5) \ \mu m$		
Bead radius r	$(7.8\pm0.2)~\mu\mathrm{m}$	$(25.8 \pm 0.5) \ \mu m$		

The geometric calculation of \tilde{m} can thus only give a rough estimation of the actual value. We estimate $\tilde{m}_A \approx 0.1$ (range 0.04–0.3) for cantilever A, and $\tilde{m}_B \approx 1.2$ (range 0.5–2.6) for cantilever B. The geometric estimation of \tilde{r} is less hampered by the uncertainty of the cantilever geometry, but still suffers from the uncontrolled repartition of the glue. We estimate $\tilde{r}_A = 0.02 \pm 0.01$ for cantilever A, and $\tilde{r}_B = 0.06 \pm 0.02$ for cantilever B. However, as demonstrated by Allen *et al* [23], the tip of the bare cantilever A itself may have a non-negligible effect. We can geometrically estimate $\tilde{m}_A \operatorname{tip} \approx 0.04$ and $\tilde{r}_{A \operatorname{tip}} \approx 0.02$ for a 17 μ m tall pyramidal tip.

The measurement is performed with a home-made interferometric deflection sensor [15], inspired by the original design of Schonenberger [21], with a quadrature phase detection technique [22]: the interference between the reference laser beam reflecting on the chip of the cantilever and the sensing beam on the cantilever gives a direct measurement of the deflection with a very high accuracy (see figure 1). This technique offers a very low intrinsic noise (down to 10^{-14} m Hz^{-1/2}). It is intrinsically calibrated as it directly measures the deflection against the wavelength of the laser beam, in contrast to the standard optical lever technique, which actually measures an angular deflection. Lastly, the focused beam size resolution is tuned to as small as 10 μ m to ensure a good spatial resolution.

We apply the methodology of Paolino *et al* [15]: at every position x and y on a 5 μ m × 5 μ m grid, we measure the deflection z(x, y, t) produced by the sole thermal excitation of the cantilever and we evaluate the power spectrum density (PSD) $S_z(x, y, f)$ on a 20 s signal sampled at 2 MHz. For a quantitative characterization of the shape of the modes, the mean squared amplitude of each resonance $\langle A_n^2(x, y) \rangle$ is determined as a function of positions x and y by integrating the PSD in a convenient frequency interval $2\Delta f$ around each resonance frequency f_n :

$$\left\langle A_n^2(x,y) \right\rangle = \int_{f_n - \Delta f}^{f_n + \Delta f} S_z^2(x,y,f) \, \mathrm{d}f. \tag{26}$$

This quantity is computed directly from the experimental spectra, without any fitting process. We take care to subtract

the background noise contribution of the interferometer, and we also compensate for the finite integration range in frequency [15].

The complete set of results for cantilever A is reported in figure 6, where the rms amplitude $\sqrt{\langle A_n^2(x, y) \rangle}$ is represented with a color coded scale. The first three vibration modes can be clearly seen with their respective number of nodes. A weak component in torsion can be seen for the third mode with the bead, certainly because it has not been glued perfectly on the axis. However, in the following, we will neglect this effect and focus on the flexural modes along the *x* axis. Therefore, at each position *x*, the median $\sqrt{\langle A_n^2(x) \rangle}$ along the *y* axis of $\sqrt{\langle A_n^2(x, y) \rangle}$ is calculated. Due to the higher reflectivity of the gold-coated cantilever B, the background noise is lower, and we accurately measure the thermal noise up to the fifth resonant mode for this last sample.

3.2. Results and discussion

3.2.1. Resonant frequency ratios. In a first attempt to estimate the added mass parameters \tilde{m} and \tilde{r} , which must be known for proper normalization of the normal modes (value of a_n), let us focus on the ratio between the resonant frequency f_n of the successive modes. Indeed, equation (23) translates into:

$$\frac{\alpha_n(\tilde{m}, \tilde{r})}{\alpha_1(\tilde{m}, \tilde{r})} = \sqrt{\frac{f_n}{f_1}}.$$
(27)

The resonant frequencies of each mode are easily found by a simple harmonic oscillator fit of each resonant peak in the thermal noise spectrum. We report in table 2 the square root of the frequencies of the first three modes of cantilever A and the first five modes of cantilever B, normalized to the frequency of mode 1. We can then compare those measurements to the output of the analytical model (figure A.1), and try to estimate the values of \tilde{m} and \tilde{r} for our samples.

For cantilever A without the bead, the presence of the tip can be detected on the measured modes, leading to the estimation $\tilde{m}_{A \text{ tip}} = 0.057 \pm 0.010$ and $\tilde{r}_{A \text{ tip}} = 0.03 \pm 0.02$



Figure 6. Maps of thermal noise of cantilever A. The rms amplitude of the first three flexural modes, with (right) and without (left) a bead glued at the free cantilever end, are plotted using a color coded amplitude map (color bar at the bottom of the figure). The resonant frequency and full scale amplitude are given below each map. A weak component in torsion can be seen clearly for the third mode with the bead, hinting at a slightly off-axis gluing of the colloidal particle.

Table 2. Frequency ratio between modes and frequency shift upon gluing of the bead. The square root of these ratios should be equal to the ratio of the corresponding spatial eigenvalues $\alpha_n(\tilde{m}, \tilde{r})$ (with a correction factor L^w/L^{wo} for the frequency shift, which we suppose equal to 0.98 here). Using estimated values of \tilde{m} and \tilde{r} for each measurement set (bare and loaded cantilevers A and B), we get a good overall agreement for every mode, especially for sample A. The model reaches its limitation for the higher modes of cantilever B, whose triangular shaped end is not taken into account.

Mode number <i>n</i>	1	2	3	1	2	3	4	5	
Bare cantilever	A: $\tilde{m}_{A \text{ tip}} = 0.057, \tilde{r}_{A \text{ tip}} = 0.03$			B: $\tilde{m}_{B tip} = 0$, $\tilde{r}_{B tip} = 0$					
f_n^{wo} (Hz)	14956	95017	267 530	7195	42975	117 220	224 500	363 370	
	1.000 1.000 0.0	2.521 2.521 -0.0	4.229 4.236 -0.1	1.000 1.000 0.0	2.444 2.503 -2.4	4.036 4.189 -3.6	5.586 5.864 -4.7	7.107 7.539 -5.7	
Loaded cantilever	A: $\tilde{m}_{\rm A} = 0.35, \tilde{r}_{\rm A} = 0.03$			B: $\tilde{m}_{\rm B} = 1.18$, $\tilde{r}_{\rm B} = 0.06$					
$\overline{f_n^{\mathrm{w}}(\mathrm{Hz})}$	10974	83 050	246 380	3101	29319	79 023	152 400	270 980	
$\frac{\sqrt{f_n^w}/f_1^w}{\alpha_n(\tilde{m},\tilde{r})/\alpha_1(\tilde{m},\tilde{r})}$ Disagreement (%)	1.000 1.000 0.0	2.751 2.762 -0.4	4.738 4.762 -0.5	1.000 1.000 0.0	3.075 3.195 -3.8	5.048 5.190 -2.7	7.010 7.040 -0.4	9.348 9.347 0.0	
Frequency shift upon loading	Cantilever A			Cantilever B					
$ \frac{(L^{\rm w}/L^{\rm wo})\sqrt{f_n^{\rm w}/f_n^{\rm wo}}}{\alpha_n(\tilde{m},\tilde{r})/\alpha_n(\tilde{m}_{\rm tip},\tilde{r}_{\rm tip})} $ Disagreement (%)	$0.839 \\ 0.844 \\ -0.5$	0.916 0.924 -0.9	$0.940 \\ 0.948 \\ -0.8$	0.643 0.642 0.2	$0.809 \\ 0.820 \\ -1.2$	0.805 0.796 1.1	0.807 0.771 4.7	0.846 0.796 6.3	

(the standard deviations correspond to a 0.2% max distance between the analytical model and the measured frequency ratio on the three modes), close to the values expected from the geometrical analysis. Adding the bead changes those values to $\tilde{m}_{\rm A} = 0.336 \pm 0.008$ and $\tilde{r}_{\rm A} = 0.029 \pm 0.005$ (same criterion for error bars).

For bare cantilever B, the ratios of frequencies between modes cannot be explained by our model: the triangular shape of the end of the cantilever alters the results from those of a rectangular one. If, however, we look at the data for the loaded cantilever, we have a reasonable agreement for $\tilde{m}_{\rm B} =$ 1.18 ± 0.09 and $\tilde{r}_{\rm B} = 0.070 \pm 0.006$ (the standard deviations correspond to a 2% max distance between the analytical model and measured frequency ratio on the five modes), again in line with the geometrical analysis. 3.2.2. Resonant frequency shifts—a naive attempt. Another way to determine the \tilde{m} and \tilde{r} parameters is to analyze the resonant frequency shifts due to the loading, in an approach similar to the Cleveland method [24]. In a naive attempt, we suppose that the process of gluing a bead to the cantilever free end should have a limited effect on its stiffness and proper mass, thus we can relate the frequencies with (f_n^W) and without (f_n^{WO}) the mass to the spatial eigenvalues with $(\alpha_n(\tilde{m}, \tilde{r}))$ and without $(\alpha_n(0, 0))$ the mass through equation (23):

$$\frac{\alpha_n(\tilde{m}, \tilde{r})}{\alpha_n(0, 0)} = \sqrt{\frac{f_n^{\rm w}}{f_n^{\rm wo}}}.$$
(28)

The resonant frequencies of each mode are easily found by a simple harmonic oscillator fit of each resonant peak of the thermal noise spectrum. We can then compare those measurements to the output of the analytical model, and try to estimate the values of \tilde{m} and \tilde{r} for our samples.

As shown in figure 10, the first mode is almost independent of the value of \tilde{r} , and should thus reliably be used to measure \tilde{m} . This measurement is equivalent to the method of the added mass proposed by Cleveland [24]. We find, with this protocol, $\tilde{m} = 0.21 \pm 0.005$ for cantilever A, and $\tilde{m} = 1.065 \pm 0.010$ for cantilever B.

This estimation of \tilde{m} is hardly compatible with the expectation of the higher order modes for cantilever A, which would rather be $\tilde{m} = 0.10 \pm 0.01$ (mode 2) or $\tilde{m} = 0.055 \pm 0.010$ (mode 3), even considering the effect of \tilde{r} . As for cantilever B, the estimation could be compatible with higher order modes, but for both cantilevers the value of \tilde{m} is clearly underestimated with respect to the previous measurement through the frequency ratios between successive modes of the loaded cantilever.

3.2.3. Resonant frequency shifts—refined analysis. Two naive hypotheses are responsible for the shortcomings of the previous analysis of the frequency shifts upon loading: it first relies on the assumption that the effective length of the loaded cantilever is unchanged. It then assumes that the behavior of the cantilever without the load is that of an ideal bare rectangular cantilever.

Let us first consider the effect of a possible modification of the effective length L of the cantilever upon gluing a bead close to its free end. Indeed, this process may rigidify the end portion of the cantilever, thus shortening its effective length by the rigid part. Alternatively, the inertia of a large mass not fixed exactly at the free end, by bringing the nodes of the higher order modes closer to the position of the bead than to the free end, can also lead to a shortening of the effective length of the loaded cantilever. In order to evidence the dependence of the eigenvalue α_n on the cantilever length L, equation (3) can be expressed as:

$$\alpha_n^4 = \frac{3\mu L^4 \omega_n^2}{EI} \tag{29}$$

where *E* is Young's modulus of the cantilever, *I* its second moment of inertia, and μ its mass per unit length. As *E*, *I* and μ do not depend either on the cantilever's length or on the gluing of a bead, equation (28) should in fact be written as

$$\frac{\alpha_n^{\rm w}}{\alpha_n^{\rm wo}} = \frac{L^{\rm w}}{L^{\rm wo}} \sqrt{\frac{f_n^{\rm w}}{f_n^{\rm wo}}} \tag{30}$$

where the superscript ^w (respectively ^{wo}) designates a quantity with (respectively without) the load. Assuming α_n^{wo} is known, the parameters \tilde{m} and \tilde{r} for the loaded cantilever can be deduced by comparing the α_n^w values obtained from equation (30) to the tabulated values given in the appendix¹. Let us stress that if the resonant frequencies are measured with a very good accuracy, the effective length ratio is not known *a priori*. A small error in the effective length ratio leads only to the same relative error in the value of α_n^{w} . But since $\alpha_n(\tilde{m}, \tilde{r})$ varies only slowly with \tilde{m} (see the appendix), it can result in a much larger error on \tilde{m} and thus on the normalization of the normal modes (parameter a_n). In section 3.2.5, we estimate that the gluing of a bead reduces the effective length of cantilevers A and B by 10 μ m, resulting in a 2% shortening. We thus use this 2% correction in table 2 to compare the relative frequency shifts to the spatial eigenvalue ratios.

When analyzing the frequency ratios with and without a bead, it is also very important to use the right α_n^{wo} value for the cantilever without the bead. We have seen with the naive approach that ignoring this initial loading for cantilever A leads to inconsistent values between modes, underestimating the true loading. However, if $\alpha_n(\tilde{m}_{A \text{ tip}}, \tilde{r}_{A \text{ tip}})$ (with $\tilde{m}_{A \text{ tip}} = 0.057$ and $\tilde{r}_{A \text{ tip}} = 0.03$ as determined in section 3.2.1) is used as the unloaded reference value instead of $\alpha_n(0, 0)$, one obtains $\tilde{m}_A = 0.37 \pm 0.04$ and $\tilde{r}_A = 0.031 \pm 0.018$ with the first three modes, in much better agreement with the previous estimation.

The behavior of the tipless cantilever B also deviates from that of a bare rectangular cantilever because of its triangular end. However, since the frequency ratios between the modes of the unloaded cantilever B do not yield any consistent set of \tilde{m} and \tilde{r} values (see figure A.1), we choose to take $\tilde{m}_{B \text{ tip}} = \tilde{r}_{B \text{ tip}} = 0$ for the bare cantilever B. We thus compare the square root of the frequency ratio (corrected by the length ratio) with the $\alpha_n(\tilde{m}, \tilde{r})/\alpha_n(0, 0)$ ratio displayed on figure 10 for the first five modes of cantilever B. As shown in figure 10, the first mode is almost independent of the value of \tilde{r} , and can thus be used to measure \tilde{m} alone; we obtain $\tilde{m}_{\rm B} = 1.17 \pm 0.04$ for mode 1. The higher order modes are compatible with the estimation of $\tilde{m}_{\rm B} = 1.17$ and can be used to guess the value of $\tilde{r}_{\rm B}$. Using figure 10, we measure $\tilde{r}_{\rm B} = 0.068$ (mode 2), $\tilde{r}_{\rm B} = 0.057 \text{ (mode 3)}, \tilde{r}_{\rm B} = 0.046 \text{ (mode 4)} \text{ and } \tilde{r}_{\rm B} = 0.030$ (mode 5). The dispersion of results is quite large for $\tilde{r}_{\rm B}$, and points to the limitations of the model with respect to the actual cantilever shape. A simultaneous least squares minimization of the distance between the analytical model and measured frequency shifts on the five modes leads to $\tilde{m}_{\rm B} = 1.19 \pm 0.13$ and $\tilde{r}_{\rm B} = 0.051 \pm 0.006$ (standard deviation corresponding to a 3.5% max distance).

As a summary, we have two ways to estimate the normalized added mass and equivalent gyration radius from the measurement of the resonant frequencies of the loaded and unloaded cantilever: the frequency ratio between modes in one measurement, and the frequency shifts due to the addition of the bead. Provided the initial tip and effective length shortening are taken into account, both methods agree reasonably, though the dispersion on \tilde{r} is quite large. In the following we will retain the values:

- Unloaded cantilever A: $\tilde{m}_{A \text{ tip}} = 0.057$, $\tilde{r}_{A \text{ tip}} = 0.03$.
- Loaded cantilever A: $\tilde{m}_A = 0.35$, $\tilde{r}_A = 0.03$.
- Unloaded cantilever B: $\tilde{m}_{B \text{ tip}} = 0$, $\tilde{r}_{B \text{ tip}} = 0$.
- Loaded cantilever B: $\tilde{m}_{\rm B} = 1.18$, $\tilde{r}_{\rm B} = 0.06$.

¹ In order to retrieve the variation of the resonant frequencies with the fixation distance of a given bead to the free end of a cantilever, or to determine the added mass $m_{\rm b}$ and the gyration radius $r_{\rm g}$ from the \tilde{m} and \tilde{r} values, one must remember that the values of \tilde{m} and \tilde{r} determined from the resonant frequencies ratios or shifts also depend on the effective length of the loaded cantilever.



Figure 7. Amplitude of thermal noise for the first three flexural modes along cantilever A with and without a sphere (in red and blue respectively). The markers represent the data while the lines exhibit the fits: independent fits of each mode as a plain line, and simultaneous fit of all modes as a dashed line. The agreement is excellent for the bare and the loaded cantilever.

Eventually, using those estimations of \tilde{m} and \tilde{r} , we can compute the values of the eigenvalues $\alpha_n(\tilde{m}, \tilde{r})$ from the model, and compare them with the frequency shifts due to the bead. As one can see in table 2, the agreement is quite good, with an overall agreement better than 1% for cantilever A, and 3% for cantilever B (except for the highest order modes of cantilever B, where the limitations of the model appear more severe).

3.2.4. Spatial modes shapes. Figures 7 and 8 show the rms amplitude $\sqrt{\langle A_n^2(x) \rangle}$ of the first three resonant modes of cantilever A and the first five modes of cantilever B respectively, with (red circles) and without (blue crosses) the bead loading their ends. It is worth mentioning that the maximum of the last mode is only a few pm high, demonstrating the high resolution of our instrument. The behavior of the experimental data is clearly in line with the model illustrated in figure 3: the effect of the bead is almost negligible on the first longitudinal mode, and the nodes of higher order modes are shifted towards the free end of the cantilever.

In a first step to fit the data, we use independent fits for all modes. The fitting function is equivalent to equation (5). This generic function should be suitable for any cantilever clamped at its origin, whatever boundary conditions are applied at its



Figure 8. Amplitude of thermal noise for the first five flexural modes along cantilever B with and without a sphere (in red and blue respectively). The markers represent the data while the lines exhibit the fits: independent fits of each mode as a plain line, and simultaneous fit of all modes as a dashed line. The agreement is very good for the bare and the loaded cantilever.

other end. Note that x is now not normalized to the length L of the cantilever. We therefore use the following fitting function for $\langle A_n^2(x) \rangle$:

$$\psi^{2}\left(x,\bar{a},\bar{R},\frac{\bar{\alpha}}{\bar{L}},\bar{x_{0}}\right)$$

$$=\bar{a}^{2}\left(\cos\left(\bar{\alpha}\frac{x-\bar{x_{0}}}{\bar{L}}\right)-\cosh\left(\bar{\alpha}\frac{x-\bar{x_{0}}}{\bar{L}}\right)\right)$$

$$+\bar{R}\left[\sin\left(\bar{\alpha}\frac{x-\bar{x_{0}}}{\bar{L}}\right)-\sinh\left(\bar{\alpha}\frac{x-\bar{x_{0}}}{\bar{L}}\right)\right]\right)^{2}$$
(31)

Table 3. Eigenvalues normalized to the third mode for both cantilevers, bare and loaded with the bead: we compare the output of generic fits with equation (31) and the values expected from the model. The agreement is quite good for every mode but the first one, where the fit is not sufficiently constrained to converge to accurate estimations.

Mode number <i>n</i>	1	2	1	2	4	5
Bare cantilever	A: \tilde{m}_{A} $\tilde{r}_{A ti}$	$t_{tip} = 0.056,$ $t_{tip} = 0.03$	В	$\tilde{m}_{\rm B tip} = 0$, $\tilde{r}_{\rm Btip} = 0$	
Measurement: $\bar{\alpha}_n/\bar{\alpha}_3$ Theory: $\alpha_n(\tilde{m}_{tip}, \tilde{r}_{tip})/\alpha_3(\tilde{m}_{tip}, \tilde{r}_{tip})$	0.278 0.236	0.599 0.595	0.277 0.239	0.602 0.598	1.389 1.400	1.760 1.800
Disagreement (%)	-18	-0.7	-15	-0.7	0.8	2.2
Loaded cantilever	A: \tilde{n} \tilde{r}_{A}	$\dot{n}_{\rm A} = 0.35, \\ = 0.03$	B:	$\tilde{m}_{\rm B} = 1.18$	$\tilde{r}_{\rm B} = 0.06$	Ď
Measurement: $\bar{\alpha}_n/\bar{\alpha}_3$ Theory: $\alpha_n(\tilde{m}, \tilde{r})/\alpha_3(\tilde{m}, \tilde{r})$ Disagreement (%)	0.251 0.210 -19	0.580 0.580 0.0	$0.191 \\ 0.193 \\ -1.1$	0.561 0.616 8.8	1.389 1.356 -2.7	1.865 1.801 -3.5

with $\bar{a}, \bar{R}, \bar{\alpha}/\bar{L}$ and $\bar{x_0}$ being the four fitting parameters. As can be seen in figures 7 and 8, the result of this procedure is excellent, the model closely matching the experimental data.

The interesting output of the fit is the spatial eigenvalue $\bar{\alpha}_n/\bar{L}$ of each mode, however, since \bar{L} is not known precisely, only relative values of $\bar{\alpha}_n$ can be compared to the theory. We choose to normalize the values of $\bar{\alpha}_n$ to the third mode: the presence of two nodes and the 'high' amplitude constrain the fit to provide reliable values for $\bar{\alpha}_3$. The result of this procedure is shown in table 3. The agreement with the theoretical ratios is good for higher order modes, within a few per cent. Mode 1 stands apart, with a higher deviation: since no nodes are present, the fit is poorly constrained and the value of $\bar{\alpha}_1$ is not reliable.

3.2.5. Stiffness determination. If the generic independent fits are interesting to compare the eigenvalues and the shapes of the eigenmodes with the theory, it is not possible to compare the amplitude of the modes to the expectation from thermal noise excitation. Indeed, the normalization of the mode depends on the boundary conditions (kinetic energy of the bead), and cannot be guessed *a priori*. To go further, we therefore perform a *simultaneous fit of all modes*, imposing the values of α_n [15]. The fitting function for $\langle A_n^2(x) \rangle$ is now

$$\varphi_n^2(x,\bar{k},\bar{L},\bar{x_0}) = \frac{3}{\alpha_n^4(\tilde{m},\tilde{r})} \frac{k_{\rm B}T}{\bar{k}} \phi_n^2 \left(\frac{x-\bar{x_0}}{\bar{L}},\tilde{m},\tilde{r}\right) \quad (32)$$

where $\phi_n(x, \tilde{m}, \tilde{r})$ is the normal mode defined and normalized in section 2.2 (we write here explicitly the dependence in \tilde{m} and \tilde{r} to underline that those ϕ_n depend on the presence of the bead). The fitting parameters are \bar{k} , \bar{L} and $\bar{x_0}$, while the values of \tilde{m} and \tilde{r} are set to the estimation of section 3.2.3.

We shall thus perform a single simultaneous fit of all modes with those three free parameters on each cantilever, bare or loaded with a bead. We tried several weightings of the modes to compensate the decreasing amplitude of higher order modes driven by thermal noise: the function ϵ_i to

minimize during the fit is defined as

$$\epsilon_j(\bar{k}, \bar{L}, \bar{x_0}) = \sum_n \alpha_n^j \int_0^L dx |\langle A_n^2(x) \rangle - \varphi_n^2(x, \bar{k}, \bar{L}, \bar{x_0})|^2 \quad (33)$$

where *j* is a weighting parameter: j = 0 corresponds to natural weighting (mode 1 dominant) and increasing *j* weights to increasingly higher order modes ('flat' weighting for j = 8). We estimate the best fitting parameters \bar{k}, \bar{L} and $\bar{x_0}$ for j = 0-12, to test the robustness of the simultaneous fitting procedure. The dashed lines in figures 7 and 8 represent the result of this fitting process. Though not as perfect as independent fits, the results are in good agreement for all modes of each cantilever, with and without the bead.

For the bare cantilever A, the best fit values are $\bar{k} = (0.321 \pm 0.008)$ N m⁻¹ and $\bar{L} = (497 \pm 1) \mu$ m. The uncertainties correspond to the standard deviation in the full range of weighting parameter *j*, their low values demonstrating the robustness of the fit. For the loaded cantilever A, the best fit values are $\bar{k} = (0.339 \pm 0.010)$ N m⁻¹ and $\bar{L} = (485 \pm 7) \mu$ m. The stiffness of the cantilever experiences a small increase (6%) after the gluing of the bead. This is not surprising since the glue increases the rigidity of the end of the cantilever, shortening its effective length by about 10 μ m according to the fit. This 2% decrease in length translates into a 6% rise of the stiffness (*k* scales as $1/L^3$), in agreement with our estimation.

For the bare cantilever B, the best fit values are $\bar{k} = (0.151 \pm 0.002)$ N m⁻¹ and $\bar{L} = (463 \pm 4) \mu$ m. Again, the dispersion of estimated parameters is very low, hinting at the robustness of the model and fitting procedure. The value of \bar{L} is quite small with respect to the manufacturer specifications, however this effective length takes into account the triangular end of the cantilever. For the loaded cantilever B the best fit values are $\bar{k} = (0.169 \pm 0.010)$ N m⁻¹ and $\bar{L} = (448 \pm 5) \mu$ m. Again, this 12% increase of the static spring constant goes in the expected direction, and is consistent with a reduction of the effective length of the cantilever by 15 μ m (which should translate into a 10% increase in k).



Figure 9. Multiplicative correction factor to apply to the thermal noise measurement of the first resonant mode with an angular deflection measurement technique to compute the cantilever stiffness: the higher the added mass, the smaller the correction.

4. Conclusion

Our work demonstrates that even a glued bead as large as 10% of the length of a soft cantilever ($k \sim 0.1 \text{ N m}^{-1}$) modifies only slightly its first flexural mode and its static stiffness. A simultaneous fit of the thermal modes leads to a small decrease in the effective length and a small increase in the effective static stiffness of a cantilever upon gluing the bead. However, since these variations respect the $1/L^3$ scaling of k, the stiffness at the geometrical free end of the cantilever appears to be the same with and without a bead loading. Thus, the classic method to determine the spring constant by measuring the thermal spectrum of the first flexural mode at the free end of the cantilever can still be used even if the microlever is functionalized. In fact, we have shown that the thermal calibration based on the first mode only gets even closer to the static stiffness when the size of the bead increases, since the first mode gathers a higher fraction of the thermal energy at the free end. We have also shown that the mass model considering the load as a modification of the boundary conditions at the free end of a beam is a good approximation and fits well with all our results. On the way, we have introduced a proper normalization method of the resonant modes, an unavoidable step in computing the thermal noise amplitude of each mode. Once the stiffness of the colloidal probe is determined through the classic thermal noise calibration, one must make sure to take into account other corrections due to the tip geometry, as demonstrated by Edwards *et al* [25].

One of the main difficulties in testing the models is to estimate correctly the mass and gyration radius of the glued bead. We have seen that it can be important to consider the initial loading due to the AFM tip and the effective length shortening in order to get consistent results: an error of 30% on \tilde{m} could have been made by only considering the frequency shift due to the bead, if one refers to a tipless cantilever. The ratio between resonant frequencies of a cantilever offers an interesting way to estimate the load properties (without any prior knowledge about the cantilever, such as its unloaded properties), by comparison to tabulated values of $\alpha_n(\tilde{m}, \tilde{r})$. We



Figure 10. Spatial eigenvalues $\alpha_n(\tilde{m}, \tilde{r})$ numerically computed for modes 1–5 as a function of \tilde{m} and \tilde{r} (\tilde{r} increases from 0 to 0.1 from the top to bottom curve in each plot), normalized to their value at $\tilde{m} = 0$. The values of these ratios for cantilevers A and B, estimated by the frequency shift due to the addition of the bead, and supposing an effective length decrease of 2%, are plotted as a labeled horizontal line. Mode 1 is almost independent of the value of \tilde{r} , and can thus be used to estimate \tilde{m} . This curve is very close to the prediction given in the Cleveland method [24].

provide a set of such values in the appendix, for the first five modes, $0 \le \tilde{m} \le 2$ and $0 \le \tilde{r} \le 0.1$

As a final remark, let us study how our findings apply to the common AFM detection scheme. Indeed, our differential interferometer allows us to measure the actual deflection at any point of the cantilever, whereas most AFMs use an optical angular deflection measurement. The latter technique needs to be calibrated to infer from the 4-quadrant photodiode output the true deflection of the probe. This step is usually done by a rigid contact between the probe and a hard surface, the calibrated displacement of the sample providing a benchmark to measure the sensitivity (in nm V^{-1}). This sensitivity is thus valid for a static deformation only, and a mode-dependent correction factor must be applied to estimate the actual deflection of a resonant mode [1]. This multiplicative factor is 0.817 when one wants to use the thermal noise measurement of the first mode to calibrate the spring constant of a classic cantilever. In the current framework, we can easily compute how this correction factor depends on the normalized bead mass \tilde{m} by comparing the slope of the first eigenmode to that of a static deflection with the same deflection at its free end. We plot the result in figure 9 (computed with $\tilde{r} = 0$): the correction rapidly vanishes as \tilde{m} increases. Indeed, we have seen that the effect of the added mass is to decrease the eigenvalue α_1 (see figure 10), thus equation (2) tends to $z_1^{(4)} = 0$ when \tilde{m} increases. This last equation is that describing the static deflection, thus the first normal mode tends to the static deflection when \tilde{m} increases. The effect of the added mass is thus doubly beneficial for the angular measurement: both the sensitivity of the sensor tends to be more accurate (figure 9) and the first mode gathers most of the thermal noise at the cantilever free end (figure 4). More generally, an accurate coefficient can be extracted from our analysis for any \tilde{m} , and applied to the thermal noise calibration of AFM colloidal probes in any commercial device².

Acknowledgments

We are grateful to Clémence Devailly for valuable assistance during the experiments, and Sergio Ciliberto for our fruitful scientific interactions. This work has been funded by the ERC project Outeflucop.

Appendix. Spatial eigenvalues $\alpha_n(\tilde{m}, \tilde{r})$

In this appendix, we plot in figure 10 the spatial eigenvalues $\alpha_n(\tilde{m}, \tilde{r})$ numerically computed for modes 1–5 as a function of \tilde{m} and \tilde{r} , normalized to their value at $\tilde{m} = \tilde{r} = 0$. The curve for α_1 is very close to the prediction given in the Cleveland method [24], linking the frequency shift to the added mass. This figure can be used to estimate \tilde{m} and \tilde{r} from the value of the frequency shift of various modes, if the initial situation corresponds to a rectangular tipless cantilever and the potential effective length decrease after gluing the particle is known.

In figure A.1, we plot the same computed eigenvalues $\alpha_n(\tilde{m}, \tilde{r})$, but normalized to the value of the first mode for the same added mass $\alpha_1(\tilde{m}, \tilde{r})$. When one has no prior knowledge



Figure A.1. Spatial eigenvalues $\alpha_n(\tilde{m}, \tilde{r})$ numerically computed for modes 1–5 as a function of \tilde{m} and \tilde{r} (\tilde{r} increases from 0 to 0.1 from the top to bottom curve in each plot), normalized to the value of mode 1. The values of these ratios for cantilevers A and B, estimated by the frequency ratios between modes, are plotted as a labeled horizontal line for unloaded (index w) and loaded (index w) cantilevers. The model is not suited to the description of the bare cantilever B (B_{wo}), but allows the estimation of \tilde{m} and \tilde{r} for the other three situations.

of the unloaded resonant frequencies, the ratio of the resonant frequencies between modes can be used to estimate \tilde{m} and \tilde{r} .

References

- Butt H J, Cappella B and Kappl M 2005 Force measurements with the atomic force microscope: technique, interpretation and applications *Surf. Sci. Rep.* **59** 1–152
- [2] Fisher T E, Marszalek P E, Oberhauser A F, Vazquez M C and Fernandez J M 1999 The micromechanics of single molecules studied with atomic force microscopy *J. Physiol.* 520 5–14
- [3] Vazquez M C, Oberhauser A F, Fowler S B, Marszalek P E, Broedel S E, Clarke J and Fernandez J M 1999 Mechanical and chemical unfolding of a single protein: a comparison *Proc. Natl Acad. Sci. USA* **96** 3694–9

² Integrated commercial AFM software may include calibration coefficients corresponding to a bare cantilever (most probably tipless), which should be taken into account for precise calibration of the photodiode sensitivity and cantilever stiffness.

- [4] Frederix P, Bosshart D and Engel A 2009 Atomic force microscopy of biological membranes *Biophys. J.* 96 329–30
- [5] Radmacher M 2007 Methods in Cell Biology vol 83, ed Y L Wang and D E Discher (San Diego, CA: Elsevier Academic Press) p 347 (Studying the Mechanics of Cellular Processes by Atomic Force Microscopy)
- [6] Sbrana F, Sassoli C, Meacci E, Nosi D, Squecco R, Paternostro F, Tiribilli B, Zecchi-Orlandini S, Francini F and Formigli L 2008 Role for stress fiber contraction in surface tension development and stretch-activated channel regulation in C2C12 myoblasts *Am. J. Physiol. Cell Physiol.* **295** 160–72
- [7] Lavrik N V, Sepaniak M J and Datskos P G 2004 Cantilever transducers as a platform for chemical and biological sensors *Rev. Sci. Instrum.* 75 2229–53
- [8] Bhushan B 2008 Nanotribology, nanomechanics and nanomaterials characterization *Phil. Trans. R. Soc.* A 366 1351–81
- [9] Cook S M, Schaffer T E, Chynoweth K M, Wigton M, Simmonds R W and Lang K M 2006 Practical implementation of dynamic methods for measuring atomic force microscope cantilever spring constants *Nanotechnology* 17 2135
- [10] Levy R and Maaloum M 2002 Measuring the spring constant of atomic force microscope cantilevers: thermal fluctuations and other methods *Nanotechnology* 13 33
- [11] Hutter J L and Bechhoefer J 1993 Calibration of atomic-force microscope tips *Rev. Sci. Instrum.* 64 1868
- [12] Sader J E, Chon J W M and Mulvaney P 1999 Calibration of rectangular atomic force microscope cantilevers *Rev. Sci. Instrum.* **70** 3967–9
- Burnham N A, Chen X, Hodges C S, Matei G A, Thoreson E J, Roberts C J, Davies M C and Tendler S J B 2003 Comparison of calibration methods for atomic-force microscopy cantilevers *Nanotechnology* 14 1–6

- [14] Butt H J and Jaschke M 1995 Calculation of thermal noise in atomic force microscopy Nanotechnology 6 1
- [15] Paolino P, Tiribilli B and Bellon L 2009 Direct measurement of spatial modes of a microcantilever from thermal noise *J. Appl. Phys.* **106** 094313
- [16] Maali A and Bhushan B 2008 Nanorheology and boundary slip in confined liquids using atomic force microscopy J. Phys.: Condens. Matter 20 315201
- [17] Maali A, Pan Y, Bhushan B and Charlaix E 2012 Hydrodynamic drag-force measurement and slip length on microstructured surfaces *Phys. Rev.* E 85 066310
- [18] Laurent J, Sellier H, Mosset A, Huant S and Chevrier J 2012 Casimir force measurements in Au–Au and Au–Si cavities at low temperature *Phys. Rev.* B 85 035426
- [19] Oguamanam D C D 2003 Free vibrations of beams with finite mass rigid tip load and flexural-torsional coupling *Int. J. Mech. Sci.* 45 963
- [20] Bellon L 2008 Thermal noise of microcantilevers in viscous fluids J. Appl. Phys. 104 104906
- [21] Schonenberg C and Alvarado S F 1989 A differential interferometer for force microscopy *Rev. Sci. Instrum.* 60 3131
- [22] Bellon L, Ciliberto S, Boubaker H and Guyon L 2002 Differential interferometry with a complex contrast *Opt. Commun.* 207 49
- [23] Allen M S, Sumali H and Penegor P C 2009 DMCMN: experimental/analytical evaluation of the effect of tip mass on atomic force microscope cantilever calibration *J. Dyn. Syst. Meas. Control* 131 064501
- [24] Cleveland J P, Manne S, Bocek D and Hansma P K 1993 A nondestructive method for determining the spring constant of cantilevers for scanning force microscopy *Rev. Sci. Instrum.* 64 403
- [25] Edwards S A, Ducker W A and Sader J E 2008 Influence of atomic force microscope cantilever tilt and induced torque on force measurements J. Appl. Phys. 103 064513