

Verification of the Crooks fluctuation theorem and recovery of RNA folding free energies

D. Collin^{1*}, F. Ritort^{2*}, C. Jarzynski³, S. B. Smith⁴, I. Tinoco Jr⁵ & C. Bustamante^{4,6}

Atomic force microscopes and optical tweezers are widely used to probe the mechanical properties of individual molecules and molecular interactions, by exerting mechanical forces that induce transitions such as unfolding or dissociation. These transitions often occur under nonequilibrium conditions and are associated with hysteresis effects—features usually taken to preclude the extraction of equilibrium information from the experimental data. But fluctuation theorems^{1–5} allow us to relate the work along nonequilibrium trajectories to thermodynamic free-energy differences. They have been shown to be applicable to single-molecule force measurements⁶ and have already provided information on the folding free energy of a RNA hairpin^{7,8}. Here we show that the Crooks fluctuation theorem⁹ can be used to determine folding free energies for folding and unfolding processes occurring in weak as well as strong nonequilibrium regimes, thereby providing a test of its validity under such conditions. We use optical tweezers¹⁰ to measure repeatedly the mechanical work associated with the unfolding and refolding of a small RNA hairpin¹¹ and an RNA three-helix junction¹². The resultant work distributions are then analysed according to the theorem and allow us to determine the difference in folding free energy between an RNA molecule and a mutant differing only by one base pair, and the thermodynamic stabilizing effect of magnesium ions on the RNA structure.

The Crooks fluctuation theorem⁹ (CFT) predicts a symmetry relation in the work fluctuations associated with the forward and reverse changes a system undergoes as it is driven away from thermal equilibrium by the action of an external perturbation. This theorem applies to processes that are microscopically reversible, and its experimental evaluation in small systems is crucial to understand better the foundations of nonequilibrium physics¹³. A consequence of the CFT is Jarzynski's equality¹⁴, which relates the equilibrium free-energy difference ΔG between two equilibrium states to an exponential average (denoted by angle brackets) of the work done on the system, W , taken over an infinite number of repeated nonequilibrium experiments, $\exp(-\Delta G/k_B T) = \langle \exp(-W/k_B T) \rangle$. The equality has been developed⁶ into a formalism that allows us to use nonequilibrium single-molecule pulling experiments to reconstruct free-energy profiles or potentials of mean force¹⁵ along reaction coordinates. Experimental testing of Jarzynski's equality in single-molecule force experiments¹⁶ used the P5ab RNA hairpin^{7,8}, which can be folded and unfolded quasi-reversibly. But for processes that occur far from equilibrium, the applicability of Jarzynski's equality is hampered by large statistical uncertainties that arise from the sensitivity of the exponential average to rare events^{17,18} (low values of W). Moreover, although the equality $\langle W \rangle = \Delta G$ holds for processes occurring near equilibrium, spatial drift in the experimental

system usually makes it difficult in practice to extract unfolding free energies using small loading rates (below a few pN s⁻¹). Drift effects decrease noticeably for larger pulling speeds, making it possible to obtain more reliable experimental data (and also good statistics as a large number of pulls can be executed in a reasonable time), but at the expense of a more irreversible unfolding process. Here we show that significant improvements can be obtained by using the CFT, which provides a more robust and more rapidly converging method to extract equilibrium free energies from non-equilibrium processes.

The CFT allows us to quantify the amount of hysteresis observed in the values of the irreversible work done to unfold and refold a macromolecule. Let $P_U(W)$ denote the probability distribution of the values of the work performed on the molecule in an infinite number of pulling experiments along the unfolding (U) process, and define $P_R(W)$ analogously for the reverse (R) process. For the CFT to be applicable, the unfolding and refolding processes need to be related by time-reversal symmetry, that is, in our experiments, the optical trap used to manipulate the molecule must be moved at the same speeds during unfolding and refolding. Moreover, the molecular transition probed always has to start in an equilibrium state (folded in the unfolding process, and denatured or unfolded in the refolding process) and reach a well-defined final state. The CFT⁹ then predicts that:

$$\frac{P_U(W)}{P_R(-W)} = \exp\left(\frac{W - \Delta G}{k_B T}\right) \quad (1)$$

where ΔG is the free-energy change between the final and the initial states, and thus equal to the reversible work associated with this process. Note that the CFT does not require that the system studied reaches its final equilibrium state immediately after the unfolding and refolding processes have been completed; it is only the control parameter that needs to attain its final value, whereas the system may continue to equilibrate to a well-defined state that is consistent with the final value of the control parameter. The equilibration occurs without change of the control parameter, and therefore contributes no work. In principle, W is an integral over the external variation of a control parameter⁹, for example, the position of the optical trap or the time⁶. In our experiments, W is well approximated by the familiar force-versus-extension integral:

$$W = \sum_{i=1}^{N_s} F_i \Delta x_i \quad (2)$$

where x_i is the distance between the ends of the molecule and N_s is the number of intervals used in the sum (see ref. 6 for a thorough discussion of this issue). Relation (1) quantifies hysteretic effects in the pulling experiment: work values larger than ΔG occur most often

¹Merck & Co. Inc., Automated Biotechnology Department, North Wales, Pennsylvania 19454, USA. ²Departament de Física Fonamental, Facultat de Física, Universitat de Barcelona, Diagonal 647, 08028 Barcelona, Spain. ³T-13 Complex Systems, Los Alamos National Laboratory, Los Alamos, New Mexico 87545, USA. ⁴Howard Hughes Medical Institute, ⁵Department of Chemistry, ⁶Departments of Physics and Molecular & Cell Biology, University of California, Berkeley, California 94720, USA.

*These authors contributed equally to this work.

along the unfolding path while (absolute) values smaller than ΔG occur more often along the refolding path. As can be seen from equation (1), the CFT states that although $P_U(W)$, $P_R(-W)$ depend on the pulling protocol, their ratio depends only on the value of ΔG . Thus the value of ΔG can be determined once the two distributions are known. In particular, the two distributions cross at $W = \Delta G$:

$$P_U(W) = P_R(-W) \Rightarrow W = \Delta G \quad (3)$$

regardless of the pulling speed. Although the simple identity (3) already gives an estimate of ΔG , it is not necessarily very precise because it uses only the local behaviour of the distribution around $W = \Delta G$. Using the whole work distribution increases the precision of the free-energy estimate¹⁹. In particular, as we show below, when the overlapping region of work values between the unfolding and refolding work distributions is too narrow (as may happen for large values of the average dissipated work, defined as $\langle W_{\text{dis}} \rangle = \langle W \rangle - \Delta G$), the use of Bennett's acceptance ratio method²⁰ makes it possible to extract accurate estimates of ΔG using the CFT (see the Supplementary Information).

We first experimentally test the validity of the CFT for a molecular transition occurring near equilibrium. For this, we use a short interfering (si)RNA hairpin that targets the messenger RNA of the CD4 receptor of the human immunodeficiency virus (HIV)¹¹ and that unfolds irreversibly but not too far from equilibrium at accessible experimental pulling speeds (dissipated work values less than $6k_B T$). Under these conditions, the unfolding and refolding work distributions overlap over a sufficiently large range of work values to justify the use of the direct method to experimentally test equation (1). The work done on the molecules during either pulling or relaxation is given by the areas below the corresponding force-extension curves (Fig. 1).

Unfolding and refolding work distributions at three different pulling speeds are shown in Fig. 2. Irreversibility increases with the pulling speed and unfolding-refolding work distributions become progressively more separated. Note, however, that the unfolding and the refolding distributions cross at a value of the work $\Delta G = 110.3 \pm 0.5 k_B T$ that does not depend on the pulling speed, as predicted by equation (3). Moreover, the work distributions also

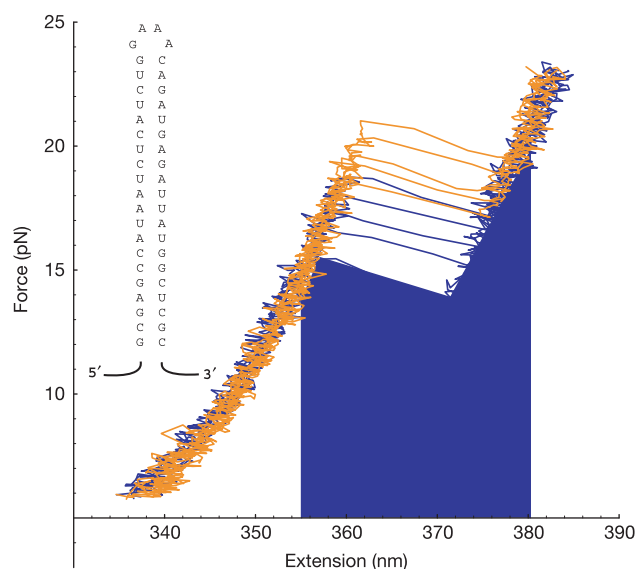


Figure 1 | Force-extension curves. The stochasticity of the unfolding and refolding process is characterized by a distribution of unfolding or refolding work trajectories. Five unfolding (orange) and refolding (blue) force-extension curves for the RNA hairpin are shown (loading rate of 7.5 pN s^{-1}). The blue area under the curve represents the work returned to the machine as the molecule switches from the unfolded to the folded state. The RNA sequence is shown as an inset.

satisfy the CFT, that is, equation (1) (see the Supplementary Information). We also notice that work distributions are compatible with, and can be fitted to, gaussian distributions (data not shown). After subtracting the contribution arising from the entropy loss due to the stretching of the molecular handles attached on both sides of the hairpin ($\Delta G^{\text{handles}} = 23.8 k_B T$) and of the extended single-stranded (ss)RNA ($\Delta G^{\text{ssRNA}} = 23.7 \pm 1 k_B T$) from the total work, $\Delta G^{\text{exp}} = 110.3 \pm 0.5 k_B T$, we obtain for the free energy of unfolding at zero force $\Delta G_0^{\text{exp}} = 62.8 \pm 1.5 k_B T = 37.2 \pm 1 \text{ kcal mol}^{-1}$ (at 25°C , in 100 mM Tris-HCl, pH 8.1, 1 mM EDTA), in excellent agreement with the result obtained using the Visual OMP from DNA software²¹ $\Delta G_0^{\text{fold}} = 38 \text{ kcal mol}^{-1}$ (at 25°C , in 100 mM NaCl).

To extend the experimental test of the validity of the CFT to the very-far-from-equilibrium regime where the work distributions are no longer gaussian, we apply the CFT to determine: (1) the difference in folding free energy between an RNA molecule and a mutant that differs only by one base-pair, and (2) the thermodynamic stabilizing effect of Mg^{2+} ions on the RNA structure. The RNA we consider is a three-helix junction of the 16S ribosomal RNA of *Escherichia coli*¹² that binds the S15 protein. The secondary structure of this RNA is a common feature in RNA structures^{22–24} that plays, in this case, a crucial role in the folding of the central domain of the 30S ribosomal subunit. For comparison, and to verify the accuracy of the method, we have pulled the wild type and a C-G to G-C mutant (C754G to G587C) of the three-helix junction.

Figure 3 depicts the unfolding and refolding work distributions for the wild-type and mutant molecules (work values were binned into about 10–20 equally spaced intervals). For both molecules, the distributions display a very narrow overlapping region. In contrast with the hairpin distribution, the average dissipated work for the unfolding pathway is now much larger—in the range $20\text{--}40 k_B T$ —and the unfolding work distribution shows a large tail and strong deviations from gaussian behaviour. Thus, these molecules are ideal to test the validity of equation (1) in the far-from-equilibrium regime. As shown in the inset of Fig. 3, the plot of the log ratio of the unfolding to the refolding probabilities versus total work done on the molecule can be fitted to a straight line with a slope of 1.06, thus establishing the validity of the CFT (see equation (1)) under far-from-equilibrium conditions. Our measurements reveal the presence of long tails in the work distribution $P_U(W)$ along the unfolding path

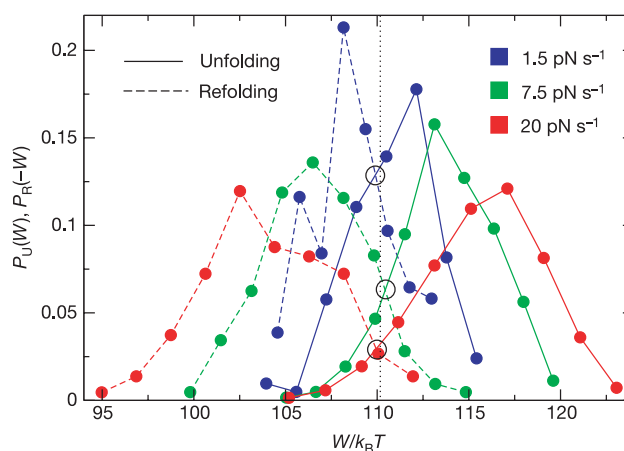


Figure 2 | Test of the CFT using an RNA hairpin. Work distributions for RNA unfolding (continuous lines) and refolding (dashed lines). We plot negative work, $P_R(-W)$, for refolding. Statistics: 130 pulls and three molecules ($r = 1.5 \text{ pN s}^{-1}$), 380 pulls and four molecules ($r = 7.5 \text{ pN s}^{-1}$), 700 pulls and three molecules ($r = 20.0 \text{ pN s}^{-1}$), for a total of ten separate experiments. Good reproducibility was obtained among molecules (see Supplementary Fig. S2). Work values were binned into about ten equally spaced intervals. Unfolding and refolding distributions at different speeds show a common crossing around $\Delta G = 110.3 k_B T$.

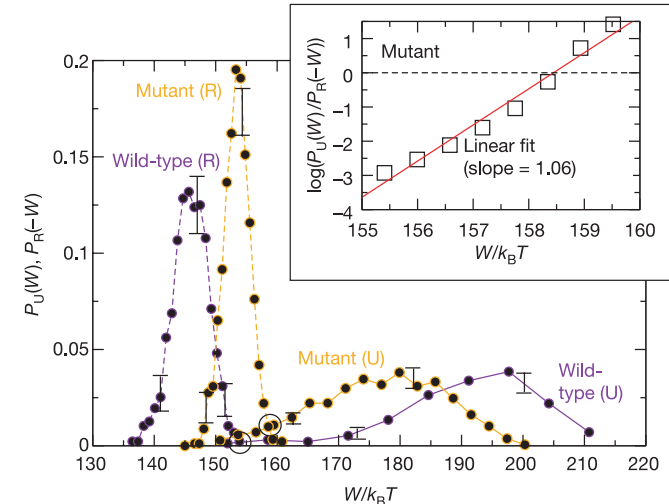


Figure 3 | Free-energy recovery and test of the CFT for non-gaussian work distributions. Experiments were carried out on the wild-type and mutant S15 three-helix junction without Mg^{2+} . Unfolding (continuous lines) and refolding (dashed lines) work distributions. Statistics: 900 pulls and two molecules (wild type, purple); 1,200 pulls and five molecules (mutant type, orange). Crossings between distributions are indicated by black circles. Work histograms were found to be reproducible among different molecules (error bars indicating the range of variability). Inset, test of the CFT for the mutant. Data have been linearly interpolated between contiguous bins of the unfolding and refolding work distributions.

and narrow work distributions $P_R(W)$ along the refolding path. These distributions complement each other, one being large where the other is small, thereby providing thermodynamically important information about the free-energy landscape.

Bennett's acceptance ratio method gives $\Delta G^{\text{exp}} = 154.1 \pm 0.4 k_B T$ and $\Delta G^{\text{exp}} = 157.9 \pm 0.2 k_B T$ for unfolding the wild-type and mutant types, respectively, giving a difference between the two forms $\Delta \Delta G_0^{\text{exp}} = \Delta \Delta G^{\text{exp}} = 3.8 \pm 0.6 k_B T$. After subtracting the (identical for both molecules) handle and RNA entropy loss contributions ($97 \pm 1 k_B T$) we get $\Delta G_0^{\text{exp}} = 57 \pm 1.5 k_B T$ (wild type) and $\Delta G_0^{\text{exp}} = 60.8 \pm 1.5 k_B T$ (mutant), the error increasing owing to the uncertainty in the contributions coming from the stretching of ssRNA. Free-energy prediction programs such as Mfold²⁵ and Visual OMP²¹ give a $\Delta \Delta G_0^{\text{fold}} = 2 \pm 2 k_B T$ between the forms (at 25 °C and 100 mM NaCl). Thus, when combined with acceptance ratio methods, the CFT furnishes a method precise enough to determine the difference in the folding free energies of RNA molecules differing only by one base pair in 34 base pairs.

Finally we apply equation (1) to obtain the free energy of

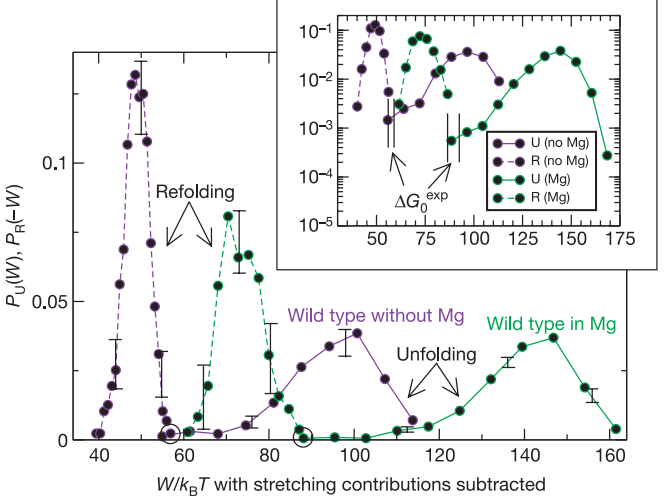


Figure 4 | Use of CFT to extract the stabilizing contribution of Mg^{2+} to the free energy of the S15 three-helix junction (wild type). Unfolding (continuous lines) and refolding (dashed lines) work distributions. Green curves, 450 pulls and two molecules in Mg^{2+} ; purple curves, 900 pulls and two molecules without Mg^{2+} . Crossings between distributions are indicated by black circles. Work histograms are reproducible between the molecules (error bars indicating the range of variability). Inset, the same histograms in logarithmic scale (axes labels as for the main panel) showing (vertical black bars) the regions of work values where unfolding and refolding distributions are expected to cross each other by Bennett's acceptance ratio method (Supplementary Information).

stabilization by Mg^{2+} of the S15 three-helix junction. These values are often difficult to access using bulk methods because melting temperatures of tertiary folded RNAs are frequently higher than the boiling point of water, and Mg^{2+} catalyses the hydrolysis of RNA at increased temperatures²⁶. Figure 4 depicts the work histograms in the presence and absence of Mg^{2+} (at constant ionic strength); stretching contributions differ in the presence and absence of magnesium ions ($116.8 k_B T$ and $97 k_B T$, respectively). These values have been subtracted from the work data to properly compare the unfolding free energies of both molecules. The strong increase of irreversibility due to Mg^{2+} can be seen in the large value of the average dissipated work (about $50 k_B T$ along the unfolding reaction and $16 k_B T$ along the refolding path). Applying Bennett's acceptance ratio method for the molecule in the presence of magnesium yields $\Delta G^{\text{exp}} = 205.5 \pm 1.5 k_B T$ and (after subtracting the stretching contributions) gives $\Delta G_0^{\text{exp}} = 88.7 \pm 2.5 k_B T$ for the unfolding reaction of the wild-type junction in 4 mM $MgCl_2$. The difference in free energies of unfolding in the presence and absence of Mg^{2+} , $\Delta \Delta G_0^{\text{exp}} = -31.7 \pm 2 k_B T$, gives the free energy of stabilization associated with the binding of Mg^{2+}

Table 1 | Summary of results obtained for all molecules

Molecule	W_m^U	W_m^R	σ_U	σ_R	$W_{\text{cum}}^{(2)}$	W_J^U	W_J^R	W_J^{est}	ΔG^{exp}	ΔG_0^{exp}	W_{dis}^U	W_{dis}^R	R_U	R_R
Hairpin (1.5 pN s ⁻¹)*	110.9	108.7	2.35	2.21	109.7 (0.2)	107.4 (0.7)	110.9 (0.2)	109.1 (0.5)	110.0 (0.2)	62.5 (1.2)	0.9	1.3	3.1	1.9
Hairpin (7.5 pN s ⁻¹)	113.8	106.6	2.63	2.84	110.3 (0.2)	109.7 (0.7)	110.9 (0.5)	110.3 (0.5)	110.3 (0.5)	62.8 (1.5)	3.5	3.7	0.98	1.10
Hairpin (20 pN s ⁻¹)	115.7	104.1	3.2	3.5	110.1 (0.2)	110.2 (0.7)	108.6 (0.2)	109.4 (0.4)	110.2 (0.6)	62.9 (1.6)	5.4	6.2	0.94	0.98
S15 (wild, no Mg)	191.3	145.9	11.3	2.9	158.7 (0.8)	155.2 (1.4)	149.3 (0.2)	152.2 (0.7)	154.1 (0.4)	57.0 (1.5)	36.3	9.1	1.75	0.46
S15 (mutant, no Mg)	176.5	153.4	10.6	2.1	156.0 (0.4)	152.4 (5.0)	155.7 (0.2)	154.1 (0.3)	157.9 (0.2)	60.8 (1.5)	18.6	4.5	3.02	0.49
S15 (wild, Mg)	256.4	190.3	12.2	5.0	213.0 (1.3)	207.0 (4.0)	199.8 (0.6)	203.6 (2.0)	205.5 (1.5)	88.7 (2.5)	50.9	15.2	1.46	0.82

W_m^U and W_m^R , σ_U and σ_R , W_J^U and W_J^R are the average total work, standard deviations and predictions obtained by using Jarzynski's equality along the unfolding (U) and refolding (R) paths. $W_{\text{cum}}^{(2)}$ is the estimate obtained by Hummer³⁰, $W_{\text{cum}}^{(2)} = (W_m^U + W_m^R)/2 - (\sigma_U^2 - \sigma_R^2)/12 k_B T$, which gives the leading correction to the linear response prediction, W_J^{est} is the average of the estimates obtained by using Jarzynski's equality along the unfolding and refolding paths $W_J^{\text{est}} = (W_J^U + W_J^R)/2$. ΔG^{exp} is our best estimate obtained by using the acceptance ratio method; ΔG_0^{exp} is the final estimate for the unfolding free energy at zero force after subtracting the handles contribution, $W_{\text{dis}}^R = |W_J^R - \Delta G^{\text{exp}}|$ is the average dissipated work (for the analysis of the hairpin data we took $\Delta G^{\text{exp}} = 110.3$ for all pulling rates) and $R_{U,R} = \frac{\sigma_{U,R}^2}{2k_B T W_{\text{dis}}^R}$ is a parameter that is equal to 1 for gaussian work distributions⁸. Statistical errors are given for Jarzynski's equality and the crossing estimates. These were obtained using the bootstrap method. All work values (except ΔG_0^{exp}) include the handle and RNA stretching contributions and are given in units of $k_B T$ at $T = 298$ K. In parentheses we indicate the errors in units of $k_B T$.

*Data for the RNA hairpin at 1.5 pN s⁻¹ are also included for completion; however, at such low loading rates drift effects are very large and data are very noisy as revealed by the values of R_U and R_R , which differ too much from 1.

ions to the S15 three-helix junction through both specific and non-specific (shielding) interactions.

These results illustrate that when used in conjunction with an appropriate fluctuation theorem, nonequilibrium single-molecule force measurements can provide equilibrium information such as folding free energies, even if the process studied occurs under far-from-equilibrium conditions. The approach works using soft optical traps but is probably limited to processes that dissipate less than $100 k_B T$. Whether it can be extended to studies that use much stiffer atomic-force-microscope cantilevers to pull proteins²⁷ is at present being examined in our laboratory. Finally, the initial and final states of molecular interactions such as ligand binding or macromolecular assembly usually do not correspond to measurable molecular extensions; in such cases, the approach as described cannot at present be applied.

METHODS

Sample preparation. The RNA molecules were prepared as previously described by ref. 16. Each DNA sequence corresponding to the three different RNAs was cloned separately into pBR322 vector between *EcoRI* and *HindIII* sites. A polymerase chain reaction (PCR) was used to amplify a DNA sequence containing an upstream T7 promoter, the RNA sequence of interest and flanking DNA sequences corresponding to the 'handles'. The handles correspond to a sequence of pBR322 (NCBI ID 'J01749') from nucleotides 3838 to 1 and from 29 to 629, respectively. The three RNA sequences were transcribed *in vitro* using T7 RNA polymerase²⁸. Two DNA handles were synthesized by PCR. The DNA handle upstream of the RNA was biotinylated at the 3'-end, whereas a digoxigenin moiety was attached to the 5'-end of the other handle. The RNA and two DNA handles were annealed by heating samples to 85 °C, followed by a slow cooling down to room temperature. The RNA hairpin was pulled in 100 mM Tris-HCl, pH 8.1, 1 mM EDTA buffer. The S15 three-helix junction and the mutant have been pulled in 62 mM KCl, 10 mM HEPES pH 7.8 buffer. In 4 mM MgCl₂ the KCl concentration was adjusted to 50 mM to work at the same ionic strength as in the absence of Mg²⁺.

Work measurements. Work probability distributions were obtained from many force-extension curves for a given molecule and aligned to a worm-like chain curve that best fitted the force-extension data at forces below the range of forces where the molecule unfolds. This procedure minimizes the effect of machine drift on the measured work values. For the worm-like chain fits we used $P \approx 10$ nm and $P \approx 1$ nm for the persistence lengths of the DNA/RNA hybrid handles and ssRNA respectively. Work values were integrated along the range of extension: [355 nm, 380 nm] for the hairpin; [326 nm, 392 nm] for the S15 three-helix junction without magnesium (wild and mutant); and [337 nm, 398 nm] for the S15 three-helix junction with magnesium. The free-energy contributions from stretching the handles and the ssRNA were then obtained by numerical integration of the worm-like chain reference curves using the values for the persistence and contour lengths of the polymers. To estimate the free energy of unfolding at zero, ΔG_0^{exp} , we subtract the free-energy contribution of the hybrid handles and ssRNA from the total reversible work across the transition, ΔG^{exp} , by using the expression^{7,29}: $\Delta G_0^{\text{exp}} = \Delta G^{\text{exp}} - \Delta G^{\text{handles}} - \Delta G^{\text{ssRNA}}$, where $\Delta G^{\text{handles}}$, ΔG^{ssRNA} are the entropy loss contributions due to the stretching of the molecular handles attached on both sides of the hairpin and of the extended ssRNA, respectively.

Received 11 April; accepted 2 July 2005.

- Evans, D. J., Cohen, E. G. & Morriss, G. P. Probability of second law violations in shearing steady states. *Phys. Rev. Lett.* **71**, 2401–2404 (1993).
- Gallavotti, G. & Cohen, E. G. D. Dynamical ensembles in nonequilibrium statistical mechanics. *Phys. Rev. Lett.* **74**, 2694–2697 (1995).
- Ciliberto, S. & Laroche, C. An experimental test of the Gallavotti-Cohen fluctuation theorem. *J. Phys. IV* **8** (Proc. 6), 215–220 (1998).
- Evans, D. J. & Searles, D. J. The fluctuation theorem. *Adv. Phys.* **51**, 1529–1585 (2002).
- Wang, G. M., Sevick, E. M., Mittag, E., Searles, D. J. & Evans, D. J. Experimental demonstration of violations of the second law of thermodynamics for small systems and short timescales. *Phys. Rev. Lett.* **89**, 050601 (2002).
- Hummer, G. & Szabo, A. Free-energy reconstruction from nonequilibrium

- single molecule experiments. *Proc. Natl Acad. Sci. USA* **98**, 3658–3661 (2001).
- Liphardt, J., Dumont, S., Smith, S. B., Tinoco, I. Jr & Bustamante, C. Equilibrium information from nonequilibrium measurements in an experimental test of the Jarzynski equality. *Science* **296**, 1832–1835 (2002).
- Ritort, F., Bustamante, C. & Tinoco, I. Jr A two-state kinetic model for the unfolding of single molecules by mechanical force. *Proc. Natl Acad. Sci. USA* **99**, 13544–13548 (2002).
- Crooks, G. E. Entropy production fluctuation theorem and the nonequilibrium work relation for free-energy differences. *Phys. Rev. E* **60**, 2721–2726 (1999).
- Smith, S. B., Cui, Y. & Bustamante, C. An optical-trap force transducer that operates by direct measurement of light momentum. *Methods Enzymol.* **361**, 134–162 (2003).
- McManus, M. T., Petersen, C. P., Haines, B. B., Chen, J. & Sharp, A. P. Gene silencing using micro-RNA designed hairpins. *RNA* **8**, 842–850 (2002).
- Serganov, A. et al. Role of conserved nucleotides in building the 16S rRNA binding site for ribosomal protein S15. *J. Mol. Biol.* **305**, 785–803 (2002).
- Ritort, F. Work fluctuations, transient violations of the second law and free-energy recovery methods. *Semin. Poincaré* **2**, 193–226 (2003).
- Jarzynski, C. Nonequilibrium equality for free energy differences. *Phys. Rev. Lett.* **78**, 2690–2693 (1997).
- Park, S. & Schulten, K. Calculating potentials of mean force from steered molecular dynamics simulations. *J. Chem. Phys.* **120**, 5946–5961 (2004).
- Liphardt, J., Onoa, B., Smith, S. B., Tinoco, I. Jr & Bustamante, C. Reversible unfolding of single RNA molecules by mechanical force. *Science* **292**, 733–737 (2001).
- Zuckerman, D. M. & Woolf, T. B. Theory of systematic computational error in free energy differences. *Phys. Rev. Lett.* **89**, 180602 (2002).
- Gore, J., Ritort, F. & Bustamante, C. Bias and error in estimates of equilibrium free-energy differences from nonequilibrium measurements. *Proc. Natl Acad. Sci. USA* **100**, 12564–12569 (2003).
- Shirts, R., Bair, E., Hooker, G. & Pande, V. S. Equilibrium free energies from nonequilibrium measurements using maximum likelihood methods. *Phys. Rev. Lett.* **91**, 140601 (2003).
- Bennett, C. H. Efficient estimates of free energy differences from Monte Carlo data. *J. Comp. Phys.* **22**, 245–268 (1976).
- SantaLucia, J. Jr & Hicks, D. The thermodynamics of DNA structural motifs. *Annu. Rev. Biophys. Biomol. Struct.* **33**, 415–440 (2004).
- Robertus, D. J. et al. Structure of yeast phenylalanine tRNA at 3 Å resolution. *Nature* **250**, 546–551 (1974).
- Long, D. M. & Uhlenbeck, O. C. Self-cleaving catalytic RNA. *FASEB J.* **7**, 25–30 (1993).
- Cate, J. H. & Doudna, J. A. Metal binding sites in the major groove of a large ribozyme domain. *Structure* **4**, 1221–1229 (1996).
- Zuker, M. Mfold web server for nucleic acid folding and hybridization predictions. *Nucleic Acids Res.* **31**, 3406–3415 (2003).
- Turner, D. H. in *Nucleic Acids: Structures, Properties and Functions* (eds Bloomfield, D. A., Crothers, D. M. & Tinoco, I. Jr) Ch. 7 (Univ. Press, New York, 2000).
- Carrion-Vazquez, M., et al. Protein nanomechanics studied by AFM single-molecule force spectroscopy. In *Emerging Techniques in Biophysics* (eds Arrondo, J. L. R. & Alonso, A.) (Biophys. Monogr. Ser., Springer, Heidelberg, in the press).
- Milligan, J. F., Groebe, D. R., Witherell, G. W. & Uhlenbeck, O. C. Oligoribonucleotide synthesis using T7 RNA polymerase and synthetic DNA templates. *Nucleic Acids Res.* **15**, 8783–8798 (1987).
- Manosas, M. & Ritort, F. Thermodynamic and kinetic aspects of RNA pulling experiments. *Biophys. J.* **88**, 3224–3242 (2005).
- Hummer, G. Fast-growth thermodynamics integration: Error and efficiency analysis. *J. Chem. Phys.* **114**, 7330–7337 (2001).

Supplementary Information is linked to the online version of the paper at www.nature.com/nature.

Acknowledgements We thank G. Hummer and A. Szabo for many discussions and G. E. Crooks, D. Chandler and J. Liphardt for a careful reading of the manuscript. F.R. was supported by the Spanish Research council and the Catalan Government (Distinció de la Generalitat). C.J. was supported by an NIH grant and the US Department of Energy. I.T. was supported by an NIH grant. C.B. was supported by the Howard Hughes Medical Institute and the David and Lucile Packard Foundation.

Author Information Reprints and permissions information is available at npg.nature.com/reprintsandpermissions. The authors declare no competing financial interests. Correspondence and requests for materials should be addressed to C.B. (carlos@alice.berkeley.edu) or F.R. (ritort@ffn.ub.es).