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Cellogram: On the Fly Traction Force Microscopy

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Complete List of Authors:	Lendenmann, Tobias; Eidgenössische Technische Hochschule Zurich; New York University Courant Institute of Mathematical Sciences Schneider, Teseo; New York University Courant Institute of Mathematical Sciences Dumas, Jérémie ; New York University Courant Institute of Mathematical Sciences; nTopology Tarini, Marco; Università degli Studi di Milano Giampietro, Costanza; Empa Materials Science and Technology Bajpai, Apratim ; New York University, Mechanical and Aerospace Engineering Chen, Weiqiang; New York University, Department of Mechanical and Aerospace Engineering Gerber, Julia; Eidgenössische Technische Hochschule Zurich Poulikakos, Dimos; Eidgenössische Technische Hochschule Zurich, Ferrari, Aldo; Eidgenössische Technische Hochschule Zurich Departement Maschinenbau und Verfahrenstechnik, Panzica, Daniele; New York University Courant Institute of Mathematical Sciences

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Cellogram: On the Fly Traction Force Microscopy

Tobias Lendenmann^{1,2,¢}, Teseo Schneider^{2,¢}, Jérémie Dumas^{2,3}, Marco Tarini⁴, Costanza

Giampietro⁷, Apratim Bajpai⁵, Weiqiang Chen⁵, Julia Gerber¹, Dimos Poulikakos¹, Aldo

Ferrari^{1,6,7*}, Daniele Panozzo^{2*}

1 ETH Zurich, Laboratory of Thermodynamics in Emerging Technologies, 8092 Zurich,

Switzerland

2 New York University, Courant Institute of Mathematical Sciences, New York 10003, USA

3 nTopology, New York 10013, USA

4 Università degli Studi di Milano, Department of Computer Science, 20133 Milano, Italy

5 New York University, Department of Mechanical and Aerospace Engineering, New York
11201, USA

6 ETH Zurich, Institute for Mechanical Systems, 8092 Zürich, Switzerland

7 EMPA, Swiss Federal Laboratories for Materials Science and Technology, 8600 Dübendorf,
Switzerland

¢ - Equal Contribution

* - Corresponding authors, panozzo@nyu.edu and aferrari@ethz.ch

Abstract

Traction Force Microscopy (TFM) derives maps of cell-generated forces, typically in the nN range, transmitted to the extracellular environment upon actuation of complex biological processes. In traditional approaches, force rendering requires a terminal, time-consuming step of cell deadhesion to obtain a reference image. A conceptually opposite approach is provided by reference-free methods, opening to the on-the-fly generation of force maps from an ongoing experiment. This requires an image processing algorithm keeping the pace of the biological phenomena under investigation. Here, we introduce an integrated software pipeline rendering force maps from single reference-free TFM images seconds to minutes after their acquisition. The algorithm tackles image processing, reference image estimation, and finite element analysis as a single problem, yielding a robust and fully automatic solution. The method's capabilities are demonstrated in two applications. First, the mechanical annihilation of cancer cells is monitored as a function of rising environmental temperature, setting a population threshold at 45°C. Second, the fast temporal correlation of forces produced across individual cells is used to map physically connected adhesion points, yielding typical lengths that vary as function of the cell cycle phase.

Introduction

Forces transmitted by mammalian cells to the extracellular environment contribute fundamental signals and actuation mechanisms to complex biological processes, including migration, organogenesis and tissue repair^{1,2}. Among cellular forces, actomyosin-generated cell contractility in the nN range, can be estimated by means of optical methods which capture the deformation imposed by cells to the substrate³. These experimental approaches have evolved from simple qualitative observations⁴ to the current family of traction force microscopy⁵ protocols yielding force maps with high spatial resolution⁶.

Despite these advancements, the accurate measurement of substrate deformations from optical images remains challenging. In particular, classical TFM techniques^{6,7} use stochastic patterns of randomly dispersed fluorescent beads in a continuous, compliant substrate⁸. Their intrinsic limitation is in the requirement of a reference, load-free image of the same field of view, to reveal the displacement of optical landmarks³. This is obtained upon de-adhesion of cells, implying that the force map can only be computed after the experiment is terminated, and thus hours after the original image is collected. For this reason, reference-based TFM methods are incompatible with on-the-fly computation of tractions during an ongoing experiment. In addition, they are not conducive to immunostaining⁹. A global pre-scanning¹⁰ of each substrate before cell seeding, although theoretically possible, practically requires a lengthy setup and yields to limited resolution.

In reference-free methods, randomly dispersed fluorophores are replaced by precisely arranged, regular patterns of fiducial markers or force sensors¹¹⁻¹³. Specifically, beds of silicone pillars or needles provide a well-established and scalable approach, which has seen its best application in the study of forces applied by individual adhesion points¹³⁻¹⁸. Upon force generation, flexible pillars are bent from their resting position and the ensuing

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3 deformation can be observed using fluorescence or transmission microscopy¹⁹. The resulting
4 quantification provides a time-resolved force vector in correspondence of each pillar¹³.
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6 Tractions maps with sub-adhesion resolution can similarly be obtained in reference-free
7 modality, using the, by comparison, confocal TFM (cTFM), an approach based on the
8 decoration of a compliant, continuum elastomeric substrate with a regular distribution of
9 electrohydrodynamic printed quantum dot (QD) discs with typical sizes in the nm range^{9, 11}.
10
11 Based on the high precision patterning of either pillars or QD nanodiscs, their load-free
12 configuration can be numerically computed^{11, 12}. Reference-free approaches are therefore
13 conducive to the online determination of force maps from a single image, in turn opening
14 the way to the iterative education of the ongoing experiment. This breakthrough
15 development is, however, hampered by the poor automation of currently available *in silico*
16 approaches, which are practically non-compatible with the online processing of large data
17 sets acquired at high frequency.
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20 Existing algorithmic solutions rely on the detection of large, undeformed regions in the
21 image of interest. Polio et al.¹² uses a radon transform to align a regular grid to the regular
22 regions²⁰ and then relies on PIV^{8, 21, 22} to estimate the displacement field. cTFM¹¹ uses a
23 Delaunay triangulation to bootstrap the detection (see Video 3 for the original method¹¹ and
24 Video 1 for ours), and an exhaustive combinatorial search to reconstruct the displacements
25 in the deformed regions. Weng et al.²³ relies on user interaction to identify regular regions
26 by manually aligning one row of undeformed pillars (see Video 4 for the original method²³
27 and Video 2 for the same image processed with ours). In all cases, these methods cannot be
28 used on problematic, yet very common, images with (i) large deformations, since the
29 synthetic reference image is too different from the acquired one, (ii) lack of a sufficiently
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3 large undeformed regions, which is common in monolayer²⁴, and (iii) images containing the
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5 substrate boundary.
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8 Here, we introduce Cellogram: the first analysis pipeline for reference-free TFM images
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10 which can robustly and automatically process the aforementioned problematic images,
11 enabling batch processing of large image sequences. Our algorithmic solution includes
12 automated image processing, reference state determination, and finite element analysis. An
13 estimate of the displacements is computed in seconds, while the accurate force
14 reconstruction is completed offline in minutes. The combination of these three steps is a
15 black-box analysis pipeline with unprecedented efficiency, compatible with the online
16 generation of force maps in long-term and/or high time resolution live cell experiments.
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18 These features are showcased by two challenging examples. First, using a cTFM approach,
19 we explore the temperature dependence of cellular force generation in cancer cells and
20 define a threshold value for thermally induced mechanical annihilation. Second, using micro-
21 pillar arrays, we detect the time correlation between forces exerted by cells on individual
22 adhesions, which provides a functional estimation of the typical length of basal stress fibers.
23 Altogether, these experimental settings yield an online information flow, which provides
24 access to additional layers of information and is far beyond the reach of the existing state of
25 the art.
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The Analysis Algorithm

Cellogram automatically converts images featuring optical landmarks (QD nanodiscs or pillars;^{11, 13}) into traction forces. Specifically, the algorithm detects the fiducial markers in the image, estimates their reference positions, and, depending on the reference-free methodology used, either estimates forces directly from displacements (pillars) or runs a finite element analysis to calculate the actuating tractions (QDs). Figure 1 a, shows a schematic view of the process, while Figure 1 b demonstrates it on an experimental image. Note that Figure 1 (c-h) provide statistics for both reference-free methodologies (first 2 columns pillars, last two columns QDs), but given the difference between the processes the timings are not directly comparable. For most experimental images (more than 95%), the processing is fully automatic. Minimal user interaction is required in the remaining cases in which high local deformations, substrate defects, or low image quality create ambiguities.

When compared to existing solvers^{11, 23, 25-27}, Cellogram proves superior in terms of calculation time for all sub-processes (Figure 1 c-g) and is overall at least one or several orders of magnitude faster (Figure 1 h). Note that the timing of the detection step is similar between cTFM and Cellogram (Figure 1 c) since the same algorithm was used. Considering the application to cTFM, the new computational approach is compatible with imaging at a time resolution of 5 s for the online estimation of displacements (for approximately 3,000 landmarks). This enables the generation of full traction maps fully automatically in less than 2 min. This time resolution compares with an average of more than 20 min for the previous method, which requires significant user input¹¹. For the application to the analysis of forces on arrays of deformable pillars, the fully automatic processing time is reduced to less than a second. This compares with 5 min of manual interaction for previous methods²³.

In addition to the fast and automatic generation of force maps, which can run in parallel with the ongoing image acquisition at high temporal and spatial resolution, Cellogram extends the analysis to all available images, including those featuring large substrate deformation. This includes data, which proved unsolvable using previous approaches. In these settings, reference-free TFM becomes a routine technique (Figure 1 i, j), which can be applied to study transient processes, without investing large amounts of time or requiring trained users. Together, the increase in speed and robustness opens up reference-free platforms to high throughput TFM and high temporal resolution online TFM. In what follows, examples are presented in which the new capabilities are fully exploited.

Temperature dependence of traction forces generated by cancer cells

Localized temperature increase provides an efficient and non-invasive mechanism to ablate unresectable tumors in soft tissues²⁸. A safe application of thermal ablation must ensure the complete eradication of the metastatic seeds, while minimizing the collateral damage to the surrounding healthy tissues. In this context, it is relevant to evaluate the sensitivity of cancer cells to increasing temperatures, up to the definition of a threshold value at which adhesion and force generation are compromised. A state which, together with genetic and metabolic damage²⁹, irreversibly leads to cell death³⁰. The response of cancer cells to hyperthermia is therefore a clinically relevant biological phenomenon which requires a statistically significant number of observations, with high temporal and spatial resolution, to distil a reliable trend of temperature dependency from a cell population.

Some information is available regarding the integrity of the cell cytoskeleton and integrin mediated adhesion at low temperatures (i.e. below 37°C³¹). However, a time-resolved evaluation of the mechanical cell response to supraphysiological thermal conditions remains

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3 unexplored. We approached this problem using cTFM to visualize substrate deformations
4 induced by HeLa cells in a temperature-controlled environment. This cell line was selected as
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6 its biochemical and adhesion properties are well described³².
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10 To verify that the selected reference-free method can capture dynamic variations of acto-
11 myosin contractility, HeLa cells were treated with Blebbistatin, a general myosin inhibitor
12 (Figure 2 a-c;³³). In particular, cells were imaged for 140 min upon addition of 50 μ m of the
13 drug to the culture medium³³. Tensions exerted by individual cells on the compliant
14 substrate decreased rapidly in the first 60 min, to then reach a stable plateau at ~40% of the
15 initial strain energy (Figure 2 c).
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18 Next, the mechanical adaptation of HeLa cells to thermal energy was investigated.
19 Environmental temperature was gradually increased from 37°C to 45°C with a $\Delta T/t$ of 1°C/h.
20 Multiple individual cells were imaged in parallel with a time resolution of 30 min. Cells
21 initially (i.e. at 37°C) conveyed different levels of mechanical strain energy U to the
22 substrate³⁴. Yet, following a sufficient number of individual cells over time (~700 images), a
23 general trend could be obtained (Figure 2 d-g). Cell-generated substrate deformation
24 remained unvaried until the temperature of 42°C was reached (Figure 2 d, e). Interestingly,
25 all cells increased significantly (>25%) their contractility during the time period in which
26 temperature increased from 42 to 44°C. In this phase, cell surfaces started to shrink (Figure 2
27 d, e). Above 44°C, the mechanical activity of all monitored cells rapidly decreased (Figure 2 f,
28 g) yielding an evident cell rounding. These results indicate that a threshold for the
29 mechanical annihilation of HeLa cells is reached at 45°C, a value that is in good agreement
30 with clinical reports²⁸.
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33 Finally, to visualize the dynamics of focal adhesion disassembly upon the mechanical
34 response to increasing temperature, the analysis was extended to cells expressing a
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3 fluorescent adhesion marker. For this, the thermal dependence of traction forces was
4 evaluated in rat embryo fibroblasts (REF-52) stably transfected with a YFP-paxillin reporter
5 (Supplementary Figure 7), a well-established model for the study of adhesion dynamics³⁵.
6
7 The overall response of these cells showed a monotone decrease of displacement and strain
8 energy, with complete mechanical annihilation at 45°C. This phase coincided with the loss of
9 visible focal adhesion (Supplementary Figure 7) and disassembly of the actin cytoskeleton
10 (Supplementary Figure 8).
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21 **Spatial and temporal correlation of traction forces during the cell cycle**

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23 The contractile machinery of cells transmits tractions to the underlying substrate in a
24 dynamic process that actuates spreading, shape change and migration³⁶. Acto-myosin
25 generated contractility is coupled to the extracellular environment by the physical
26 interaction between ventral actin stress fibers and integrin-based contacts³⁷. At the basal
27 side of adherent cells, bundled actin filaments define linear connections between oppositely
28 growing focal adhesions, therefore establishing tension across the cell body. While the
29 typical architecture of these structural elements has been described, the interaction with the
30 substrate and the resulting cell adaptation remain poorly understood³⁷.
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33 Force sensor arrays comprising elastomeric posts with diameter and period of few microns
34 offer a structured substrate to adhering cells, allowing the establishment of focal adhesions
35 at the upper surface of individual pillars¹³. The resulting one-to-one interaction renders a
36 digital, reference-free representation of the cell's mechanical activity opening to the
37 investigation of force transmission patterns during complex processes and their modulation
38 in different phases of the cell cycle^{9,38}. In this direction, a spatial and temporal analysis of
39 cell-generated tractions is possible, which however requires an agile and efficient approach
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3 to define the cross-correlation among several mechanical elements, at high frequency and
4 over long periods of observation.
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9 We approached this application using an array of deformable pillars (4 μm pitch, 1.8 μm
10 diameter, 7.1 μm height and 2.5 MPa Young's modulus) on which HeLa and MDCK cells
11 expressing a Fucci cell-cycle phase sensor were seeded (Figure 3,^{9,39}). Individual cells were
12 imaged for 160 min with a time resolution of $\Delta t = 4$ min, compatible with the typical
13 dynamics of focal adhesion maturation³⁷. The global analysis of pillar bending yielded a
14 deformation vector for all individual elements at each time of measure. The pair-wise
15 Pearson correlation coefficient of these field (on average 38 and 53 pillars per cell, for HeLa
16 and MDCK; respectively) was then calculated (Supplementary Figure 9). Pillar couples on
17 which force transmission increased or decreased at the same time received a
18 positive correlation score (+1). Couples that on the contrary displayed an antithetic behavior
19 received a negative correlation score (-1). Pairs of pillars which behaved independently or
20 showed no bending had zero correlation (Supplementary Figure 9 a, b). This temporal
21 correlation analysis was extended to all possible combinations of pillars interacting with
22 individual cells, yielding a total of ~51 thousand analyzed pairs.
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25 Negative temporal correlation between paired pillar movements (i.e. smaller than -
26 0.75) was seldom detected (0.01%). This result indicates that cellular forces are not likely to
27 generate sliding or oscillation of rigid bodies. On the other hand, significant positive
28 correlation (i.e. larger than 0.75) was evident for specific couples of pillars (17.2%), which
29 increased or decreased deformation at the same time along the entire observation period
30 (Figure 3 and Supplementary Figure 9). Correlated elements moved together, compatibly
31 with the increase or decrease of contractile actuation. Based on this global analysis, a plot of
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3 temporal correlation as function of the distance between the considered elements was
4 obtained (Figure 3 f, g and l, m for HeLa cells and MDCK; respectively). In general, the
5 correlation between pillar couples decreased exponentially at increasing interpillar distance
6 (Figure 3 f-g, l-n).
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10 The temporal correlation analysis was extended to cancerous and normal epithelial cells
11 along the cell cycle to investigate how the dynamic patterns of force transmission vary as
12 function of the cell cycle phase. In particular, HeLa cells in the G1 phase, the most
13 mechanically active^{9,38}, yielded a correlation distance up to 40 μm with average values of
14 18 μm . The transition to the ensuing S/G2 phase reduced the average correlation length to
15 12 μm (Figure 3 a and f-g). Corresponding values were obtained when the actual length of
16 actin stress fibers was measured in fixed specimens (average and maximal length of 15 ± 0.5
17 μm and $44.5 \mu\text{m}$; respectively. Figure 3 b-e). Consistently, focal adhesions were established
18 in correspondence of individual pillars and positively correlated pillar pairs aligned along
19 visible stress fibres (Figure 3 h-k).
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22 The same investigation was performed on MDCK cells stably expressing the Fucci sensor,
23 rendering however a different picture (Figure 3 l-n). Subconfluent cells (single cells or small
24 islands of 2-4 cells) showed shorter actin fibers (average and maximal length of $6.5 \pm 0.5 \mu\text{m}$
25 and $20.5 \mu\text{m}$; respectively. Supplementary Figure 10) mostly restricted to the cell periphery.
26 Cells displayed a dynamic pattern of pillar bending whereby individual pillars were engaged
27 and deformed for short periods of time ($10 \pm 4 \text{ min}$ and $14 \pm 4 \text{ min}$ for MDCK cells in G1 or
28 S/G2; respectively. Figure 3 n) as compared to the corresponding values measured in HeLa
29 cells ($35 \pm 4 \text{ min}$ and $42 \pm 4 \text{ min}$; for HeLa cells in G1 or S/G2; respectively. Figure 3 n)
30 supporting more active shape changes. The temporal analysis of tractions exerted on pillars
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3 by MDCK cells revealed shorter correlation distances (Figure 3 m, n and Supplementary
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5 Figure 7) with no significant differences between the G1 and S/G2 phase.
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10 Altogether, the on-the-fly representation of tractions exerted by cells on pillar arrays was
11 compatible with the temporal and spatial analysis of force correlation. Forces exerted by
12 HeLa cells displayed stable correlations over long distances, particularly extended in the G1
13 phase of the cell cycle. On the contrary, MDCK cells generated a more dynamic pattern of
14 tractions resulting in limited force correlation along the entire cell cycle. This pattern
15 supported higher motility and active shape changes consistent with a mesenchymal
16 phenotype typical of subconfluent epithelial cells⁴⁰.
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Discussion

The Cellogram algorithm applies to reference-free TFM methods. Quickly after the acquisition of a fluorescent image, it provides a full representation of traction forces computed from the displacement of fiducial markers (Figure 1). Therefore, the mechanical activity of cells becomes accessible at the pace of the biological phenomenon under study opening to high resolution and high-throughput TFM, reducing, and in most cases eliminating completely, the extensive manual labour required by previous algorithms for reference-free TFM. In addition to describing the algorithm and providing a public, reference implementation, we showcase these new possibilities in two applications based on complementary reference-free TFM technologies.

cTFM Approach¹¹: A temperature increase above 37°C affects membrane fluidity and permeability, leading to cell volume and surface reduction and eventually to cell death⁴¹. Cellogram was used to render the mechanical response of HeLa and REF-52 cells to increasing environmental temperature (Figure 2 and Supplementary Figures 7 and 8). The resulting population analysis revealed a threshold value at which the cytoskeleton and focal adhesions to the substrate were thermally destabilized (at 45°C) and cells mechanically annihilated. In cancer cells, this phase was preceded by a transient increase of substrate deformation, which may be caused by a rapid cell membrane retraction while cells are still holding on their adhesion to the substrate.

Micro-Pillar Arrays²³: Mechanically active stress fibers have a typical length between 10 and 100 μm, depending on the cell type and activity³⁷. This specific architecture was never linked to the cell cycle phase. The application of Cellogram enabled the processing of large datasets reporting the dynamic deformation of elastomeric pillars interacting with HeLa and MDCK cells expressing a fluorescent reporter of the cell cycle phase (Figure 3). The

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3 resolved spatial and temporal cross-correlation analysis revealed a typical fingerprint of
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5 force generation which coincided with the extension of stress fibers. Specific variations along
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7 the cell cycle were revealed in cancer cells, whereby the G1 phase featured high correlation
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9 over longer distances (up to 40 μ m) consistent with high mechanical activity⁹.
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Materials and Methods

Cell Culture

Cells were cultured in an incubator at 37°C and in 5% CO₂ humidified atmosphere. For the temperature series on the QD nanodisc array substrates HeLa cells (American Type Culture Collection; ATCC) and rat embryonic fibroblasts stably expressing YFP-Paxillin (REF-52;³⁵) were used. They were cultured in high-glucose Dulbecco's modified Eagle's medium (DMEM) supplemented with 2 mM L-Glutamine, 100 U/ml penicillin, 100 µg/ml streptomycin, and 10% fetal bovine serum. For cell cycle synchronization, HeLa cells were pre-incubated overnight with the G2/M blocker RO-3306⁴² at a final concentration of 10 µm. The experiments were started at a temperature of 37°C degrees. The rate of temperature increase was set to one Celsius per hour. The temperature of the cell culture medium was measured with TSic temperature sensors (IST AG, Switzerland) and cells were imaged every 30 min.

For the cell cycle experiments we used HeLa³⁹ and MDCK⁴³ cells stably transfected with the Fucci2 biomarker and seeded them on micro-contact printed micropillars. The MDCK-Fucci stable cell line was a gift from A. Cavalcanti Adam. The cells were seeded onto the pillar array 12 to 18 h prior to the experiment. Images were taken every 4 min for a total of 160 min.

Live-cell imaging

Cells were allowed to spread on the target substrates overnight before imaging. Temperature, CO₂ and humidity were controlled during imaging using an incubation chamber. QD nanodisc array imaging was performed with a Nikon TI N DIC with a 60x oil

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3 immersion objective with 1.35 NA. Micropillar array imaging was done with a Zeiss Axio
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5 Observer. Z1 and 40X (EC Plan-NEOFLUAR 40x/0.75) objective with 0.55 NA.
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9 *Substrate preparation*

10 Arrays of QD nanodiscs were generated as reported in¹¹. Briefly, the QDs were deposited
11 on the substrate by electrohydrodynamic nanodrip-printing⁴⁴. The colloidal ink containing
12 the QDs is ejected from a micro-sized gold-coated nozzle in an electric field onto the
13 elastomeric substrate. By modulating the electric field and position of the substrate the QDs
14 are deposited in a controlled orderly fashion. The electro-hydrodynamic nanodrip-printing
15 technology can be commercially obtained through an ETH Zurich spin-off company
16 (<http://www.srona.ch>). Before performing the experiments, the temperature dependence
17 of the substrate mechanical properties was evaluated, resulting perfectly stable within the
18 analyzed temperature range⁴⁵. The surfaces were homogenously coated with fibronectin as
19 previously reported¹¹.
20
21

22 The PDMS micropillar array was manufactured by a two-step molding process⁴⁶⁻⁴⁹. First, a
23 Si, master mold was created using photo lithography. The master mold was used to create a
24 negative mold using a 10:1 Polydimethylsiloxane (PDMS) solution, (w/w, base: curing agent
25 Sylgard 184, Dow, Corning). This negative mold was functionalized with oxygen plasma and
26 Trichloro (1H,1H,2H,2H-perfluorooctyl) silane and placed it on top of a cover glass with a
27 drop of 10:1 PDMS and cured for 40 hours in 110 °C. The PDMS negative mold was then
28 peeled off to recover the PDMS micropillar arrays.
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31 To visualize and functionalize the PDMS micropillar, micro-contact printing was used to
32 modify the top of PDMS micropillar with fluorescent ECM proteins to promote cell
33 attachment. 7 mm x 7 mm 30:1 PDMS stamp were treated with 30 µL of mixed solution
34 drops comprising of fibronectin (50 µg/ml; Sigma-Aldrich) and Alexa-Fluor 647-conjugated
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3 fibrinogen (25 µg/ml; Life Technologies) for 1 h at room temperature. Treated PDMS stamps
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5 were rinsed with distilled (DI) water and blown dry with nitrogen gas. Meanwhile, PDMS
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7 micropillar arrays were pre-treated with ultraviolet (UV) ozone for 7 min in a UVO cleaner
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9 (Jelight). Functionalized PDMS stamp was placed in contact with the top of the PDMS
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11 micropillar arrays. Through this process, the functionalization proteins were transferred
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13 from the PDMS stamps to the tops of PDMS micropillar. PDMS stamp was removed and the
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15 PDMS micropillar array was washed by pure ethanol, 70% ethanol and DI water. 0.2%
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17 Pluronic (Sigma-Aldrich) solution was used to block the un-stamped regions of the PDMS
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19 micropillar array. After 60 min of immersion, DI water and PBS (Life Technologies) was used
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21 to wash away the excess Pluronic solution.
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25 *Immunofluorescence*

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29 Cells were fixed with 4% paraformaldehyde in phosphate-buffered saline (PBS) for 10 min
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31 at room temperature (RT) and permeabilized with 0.5% Triton in PBS for 10 min. After
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33 blocking with 5% bovine serum albumin (BSA) in PBS for 1 h, samples were incubated at 4°C
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35 overnight with primary antibodies. The following commercial primary antibody was used:
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37 rabbit anti-phospho-paxillin Tyr118, Cell Signaling (#2541S), dilution: 1:100⁵⁰. Secondary
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39 antibody was Alexa Fluor 647 chicken anti-rabbit, Life Technologies (#A-21443), used at
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41 1:200 for 1 h at RT. Actin cytoskeleton has been stained with Alexa Fluor 555 phalloidin, Life
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43 Technologies (#A34055), while nuclei were stained with Hoechst, Sigma (#H6024).
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47 *Temporal and spatial correlation*

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50 Temporal correlation calculations were performed using MATLAB's 'corrcoef' function
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52 (Figure 3 f, g, l, m and Supplementary Figure 7). Here, the full temporal displacement profiles
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54 for pillars were correlated against each other to find the highest Pearson correlation
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3 coefficients. Pillars showing displacement profiles that deflect simultaneously with similar
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5 magnitudes have higher correlation coefficients.
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8 For the spatial correlation (Figure 3 j, k) at a fixed time point, the correlation coefficient or
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10 similarity coefficient was defined as $\frac{u \cdot v}{\|u\| \cdot \|v\|} \cdot \min\left\{\frac{\|u\|}{\|v\|}, \frac{\|v\|}{\|u\|}\right\}$, where u and v are the
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12 displacement vectors for two pillars, respectively. This yields a value of 1 for displacements
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14 in the same direction and magnitude, -1 for opposite direction but same magnitude, and 0
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16 for perpendicular vectors. Hence, two vectors with spatial correlation of -1 are being pulled
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18 together with the same force.
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23 *Statistics*
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25 Statistical comparison of population means was performed using a non-parametric Mann-
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27 Whitney U test ($\alpha = 0.05$). All quantitative measurements reported are expressed as average
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29 values \pm the standard error of the mean. The analysis was always based on three or more
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31 independent experiments.
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Author contributions

T.L., D. Panizzo, D. Poulikakos, and A.F. designed the study; T.L., A.F. C.G. and J.G produced QD substrates and performed cell experiments and their analysis; A.B., W.C., produced pillar substrates and performed the according experiments. T.L., D. Panizzo, T.S., J.D., M.T. developed the algorithms and software. All authors contributed to the manuscript.

Supporting Information

- **Supplementary:** contains the full description of Cellogram's algorithm.
- **Software documentation:** explains how to use Cellogram's software.
- **Software:** contains the packaged Cellogram's binaries.
- **S_Video_<1-4>:** Screen recording to compare Cellogram with state-of-the-art traction force tools.

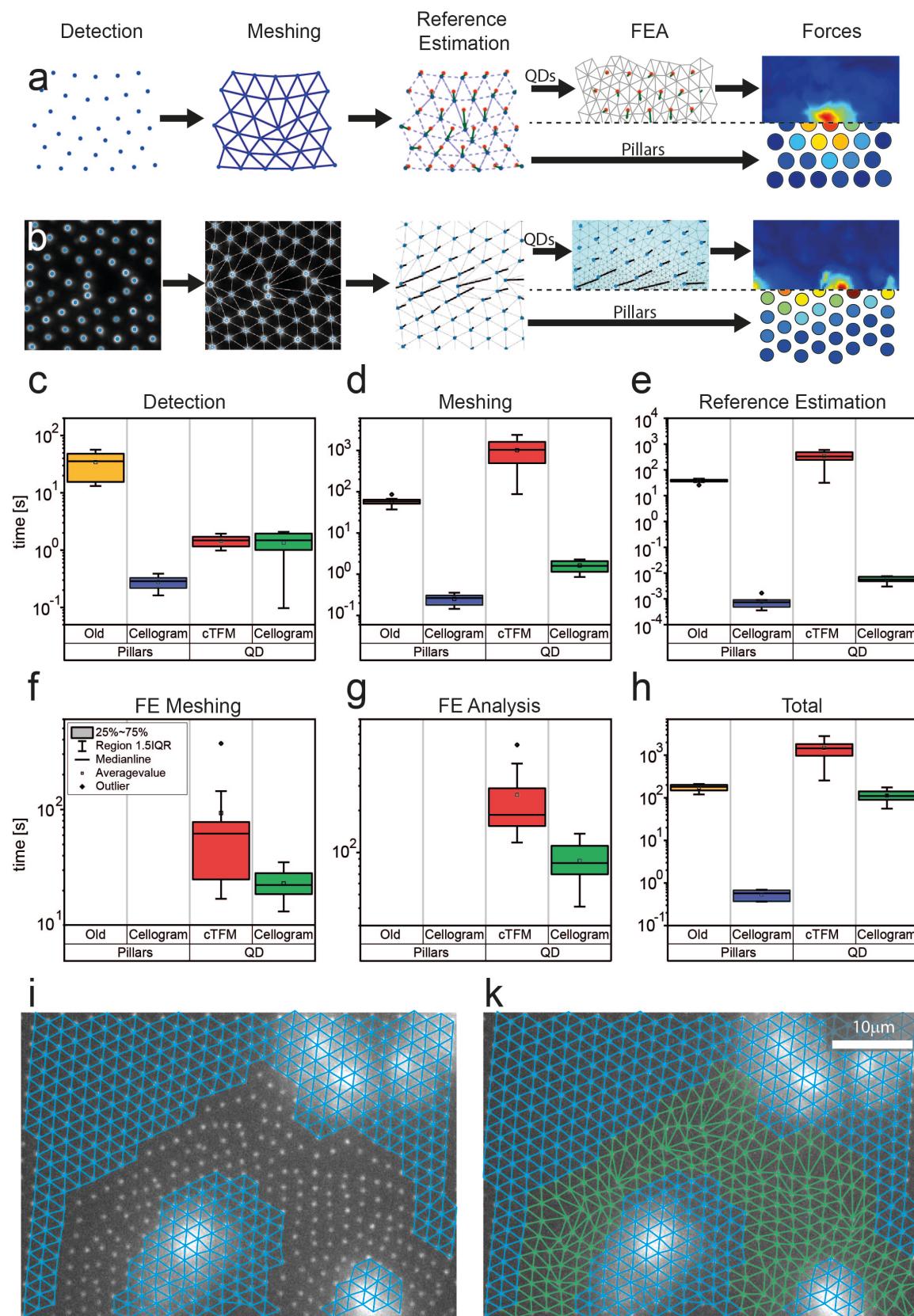


Figure 1. Processing pipeline improvements (a) Schematic of the Cellogram processing pipeline. (b) Software screenshots of the Cellogram pipeline corresponding to the steps in (a). (c-g) comparison of computation times for individual pipeline stages in Cellogram and Old methods. (h) Total computation time comparison. (i-k) FE Meshing and Analysis results.

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3 previously used software on 10 images each. (h) Computation time comparison for the total
4 analysis time per image. (i) Unsuccessful meshing attempt in cTFM (j) Completed meshing in
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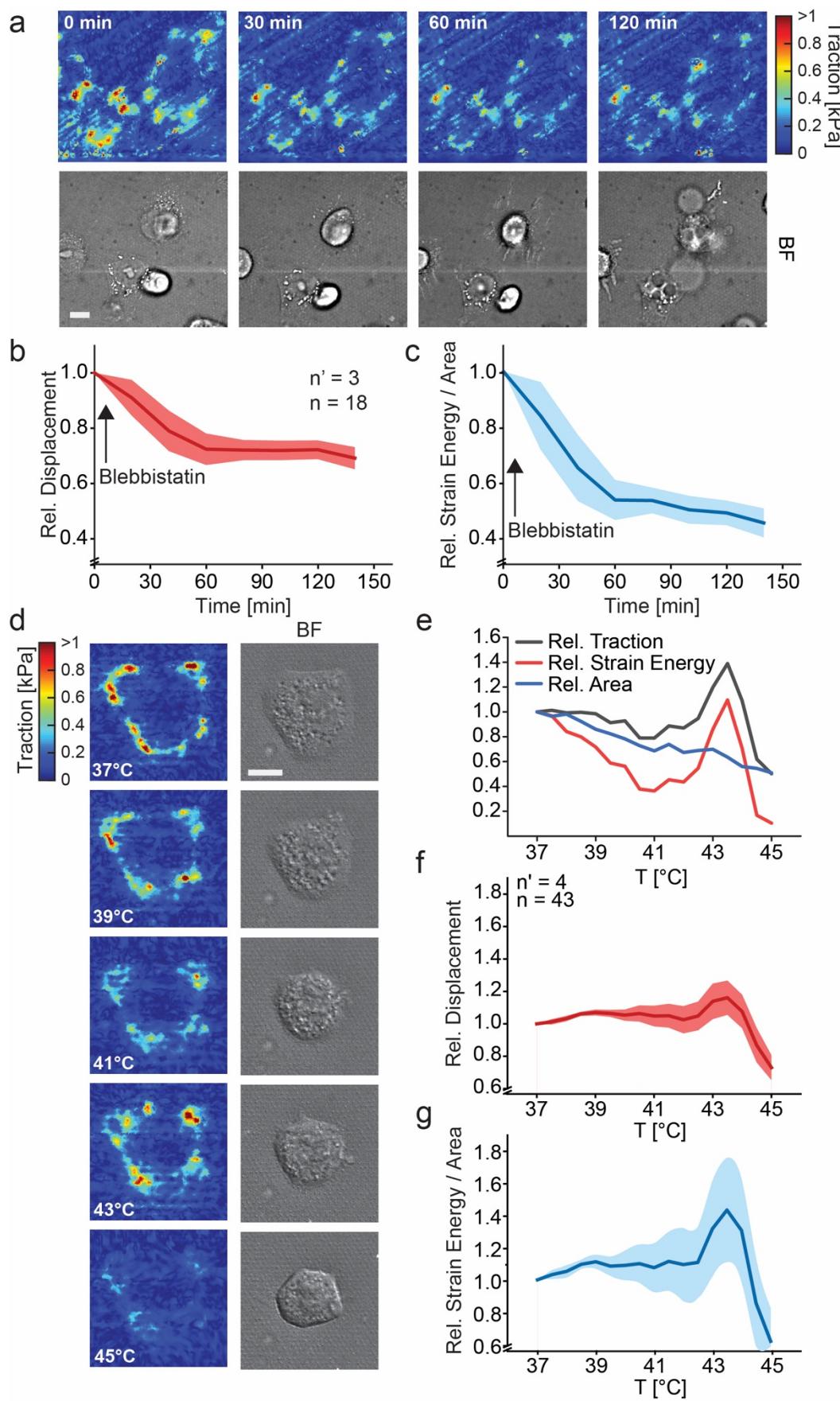
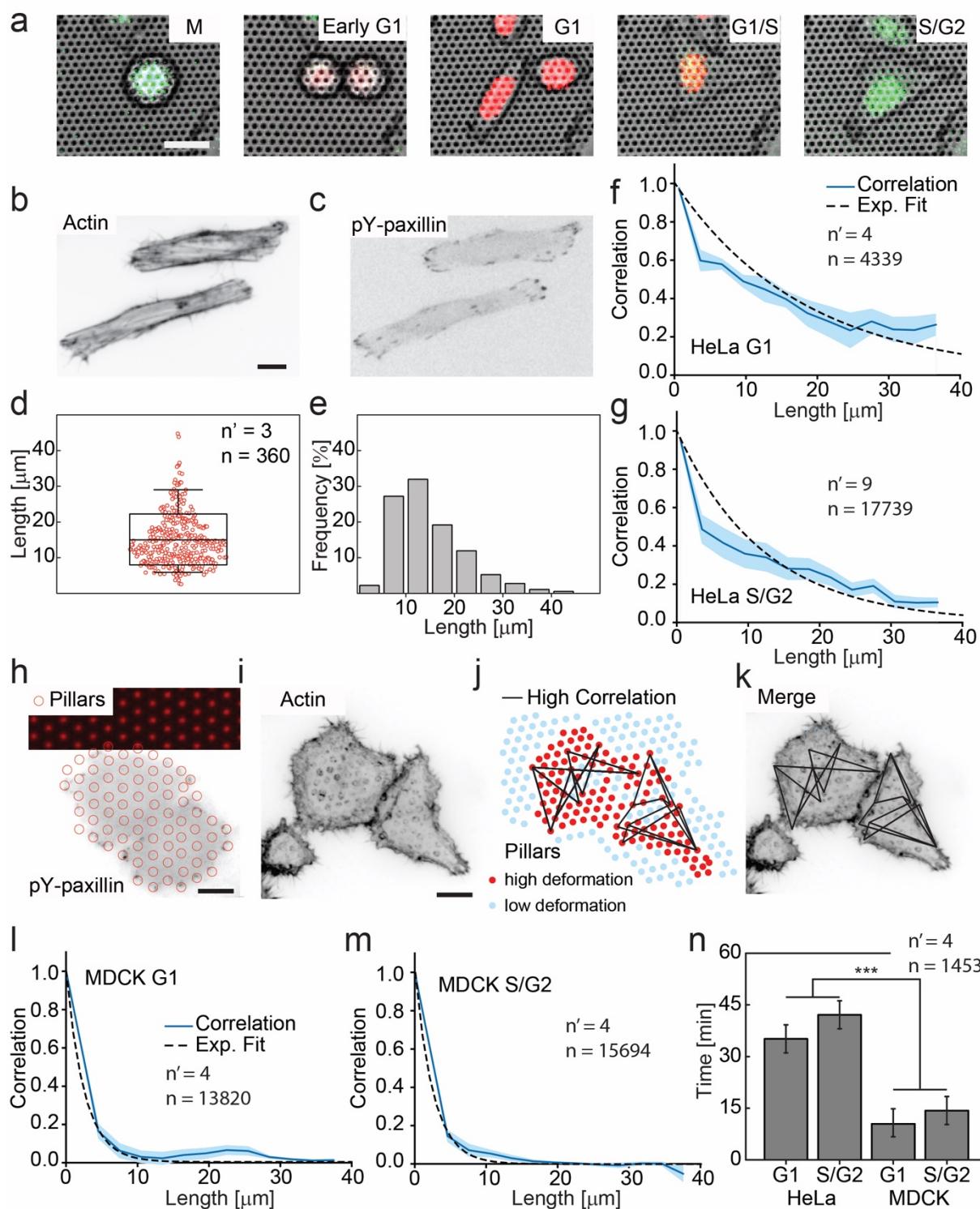


Figure 2. Mechanical and thermal annihilation of cancer cells. (a) Temporal variation of tractions exerted by HeLa cells upon treatment with Blebbistatin (50 μ M). Representative

example of cellular traction maps (upper panels) and corresponding brightfield images (BF, lower panels). Scale bar is 10 μm . (b-c) Population analysis reporting the displacement (b) and strain energy/area (c) relative to the values measured before treatment (i.e. Time = 0 min). A red or blue line defines the average value. The shaded area corresponds to the standard error of the mean. (d-g) Mechanical response of HeLa cells to increasing environmental temperature. (d) Representative example of cellular traction maps (left panels) and corresponding brightfield images (BF, right panels). Scale bar is 10 μm . (e) Corresponding evolution of traction, strain energy and area relative to the values measured at 37°C. Population analysis reporting the displacement (f) and strain energy/area (g) relative to the values measured at 37°C. A red or blue line defines the average value. The shaded area corresponds to the standard error of the mean. For all panels, n = number of individual measures, n' = number of independent experiments.



nuclear colouring in HeLa cells expressing a Fucci reporter. Scale bar is 20 μm . (b, c) Representative inverted fluorescent images reporting the distribution of filamentous actin (Actin; b) and phosphorylated paxillin (pY-paxillin; c) in HeLa cells. Scale bar is 20 μm . (d) Boxplot reporting the calibration of actin stress fiber length in fixed samples. The bars

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3 extend from the 25th to the 75th percentile. A line in the box represents the mean value
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5 while the box height corresponds to its standard deviation. Individual measures are reported
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7 as open red circles. (e) Corresponding frequency distribution (in percentage) of measured
8 values. (f, g) Correlation of traction forces as a function of interpillar distance for HeLa Fucci
9 cells in the G1 (f) or S/G2 (g) phase of the cell cycle. A blue line defines the average value.
10 The shaded blue area corresponds to the standard error of the mean. A dashed black line
11 displays the exponential fit for the experimental data from which a typical length
12 (correlation = 0.5) is extrapolated. (h) Colocalization of focal adhesions (individuated by the
13 inverted fluorescent signal of pY-paxillin in fixed HeLa cells) and pillar tops (open red circles).
14 Scale bar is 10 μ m. (i-j) Colocalization of actin filaments (individuated by the inverted
15 fluorescent signal of filamentous actin in fixed HeLa cells; i) and highest positive interpillar
16 correlations (j). (k) Merged representation of panels i and j. (l-m) Corresponding analysis of
17 traction force correlation during different phases of the cell cycle in MDCK cells. Correlation
18 of traction forces as a function of interpillar distance for MDCK cells in the G1 (l) or S/G2 (m)
19 phase. A blue line defines the average value. The shaded blue area corresponds to
20 the standard error of the mean. A dashed black line displays the exponential fit for the
21 experimental data from which a typical length (correlation = 0.5) is extrapolated. (n) Average
22 time of engagement for pillars interacting with HeLa or MDCK cells in the G1 or S/G2 phase
23 of the cell cycle. For all panels, n = number of individual measures, n' = number of
24 independent experiments.
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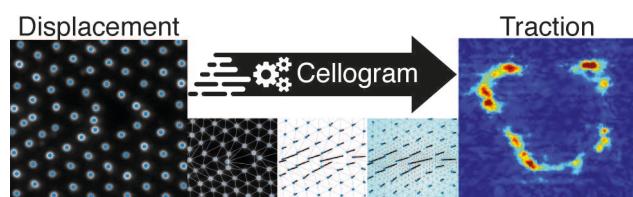
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