

Universality and Stability Phase Diagram of Two-Dimensional Brittle Fracture

Yuri Lubomirsky,¹ Chih-Hung Chen,² Alain Karma,^{2,*} and Eran Bouchbinder^{1,†}

¹*Chemical and Biological Physics Department, Weizmann Institute of Science, Rehovot 7610001, Israel*

²*Department of Physics and Center for Interdisciplinary Research on Complex Systems, Northeastern University, Boston, Massachusetts 02115, USA*

 (Received 19 July 2018; published 25 September 2018)

The two-dimensional oscillatory crack instability, experimentally observed in a class of brittle materials under strongly dynamic conditions, has been recently reproduced by a nonlinear phase-field fracture theory. Here, we highlight the universal character of this instability by showing that it is present in materials exhibiting widely different near crack tip elastic nonlinearity, and by demonstrating that the oscillations wavelength follows a universal master curve in terms of dissipation-related and nonlinear elastic intrinsic length scales. Moreover, we show that upon increasing the driving force for fracture, a high-velocity tip-splitting instability emerges, as experimentally demonstrated. The analysis culminates in a comprehensive stability phase diagram of two-dimensional brittle fracture, whose salient properties and topology are independent of the form of near tip nonlinearity.

DOI: [10.1103/PhysRevLett.121.134301](https://doi.org/10.1103/PhysRevLett.121.134301)

Cracks mediate materials failure and hence understanding their spatiotemporal dynamics is of prime fundamental and practical importance [1–5]. It is experimentally well established that cracks in brittle materials undergo various symmetry-breaking instabilities [5–13]. Most notably, straight cracks in three-dimensional (3D) samples of isotropic materials universally undergo a microbranching instability, where short-lived microcracks branch out sideways from the main crack, at low to medium propagation velocities [5–11]. Recent experiments performed on a class of neo-Hookean (NH)—a nonlinear extension of linear, Hookean elasticity—brittle materials revealed that upon reducing the system’s thickness, approaching the two-dimensional (2D) limit, the microbranching instability is severely suppressed; as a result, straight cracks accelerate to unprecedentedly high velocities, approaching the relevant sonic velocity, until they undergo an oscillatory instability [4,12,13].

Our understanding of these dynamic fracture instabilities is far from complete. In particular, the classical theory of brittle cracks—linear elastic fracture mechanics (LEFM) [1,2]—intrinsically falls short of explaining these instabilities [4,14,15]. Recently, integrating theoretical ideas about the existence and importance of elastic nonlinearity near cracks tips [4,13,16–19] into a phase-field fracture theory, the 2D oscillatory instability in brittle NH materials has been quantitatively reproduced in large-scale simulations for the first time [14], cf. Fig. 1(a). The theory has shown, in agreement with experiments, that the wavelength of oscillations scales linearly with an intrinsic nonlinear elastic length scale, which does not exist in LEFM, hence explicitly demonstrating the failure of the classical theory of brittle cracks.

This significant progress raised several pressing questions of fundamental importance. First, is the oscillatory instability universal, i.e., observed in various brittle materials independently of the nature and form of near tip elastic nonlinearity? Second, what are the minimal physical conditions for the existence of the oscillatory instability? Third, are there additional, previously undiscovered instabilities in 2D dynamic fracture? Finally, can one derive a comprehensive stability phase diagram of two-dimensional brittle fracture and if so, is it universal? In this Letter, we extensively address these important questions using theoretical considerations and large-scale simulations in the framework of the recently developed nonlinear phase-field theory of fracture [14].

This theory belongs to a broader class of diffuse-interface approaches to fracture [21–26] that avoid the difficulty of tracking the evolution of sharp crack surfaces, and at the same time allow a self-consistent selection of the crack’s velocity and path, which are far from being understood in general. These properties emerge from the dynamics of an auxiliary phase field ϕ and its coupling to other fields, which provide a mathematical machinery that renders the fracture problem self-contained. In particular, it gives rise to a dissipation-related length scale ξ and a fracture energy $\Gamma(v)$ (the energy dissipated per unit crack surface, where v is the crack propagation velocity). The smooth transition in space between the pristine ($\phi = 1$) and the fully broken ($\phi = 0$) states of the material is initiated when the elastic strain energy functional $e_{\text{strain}}(\mathbf{H})$ exceeds a threshold e_c , where $\mathbf{H} = \nabla \mathbf{u}$ is the displacement gradient tensor and $\mathbf{u}(x, y, t)$ is the displacement vector field.

Unlike previous phase-field approaches [22,23,25], the formulation in [14] maintains the wave speeds constant

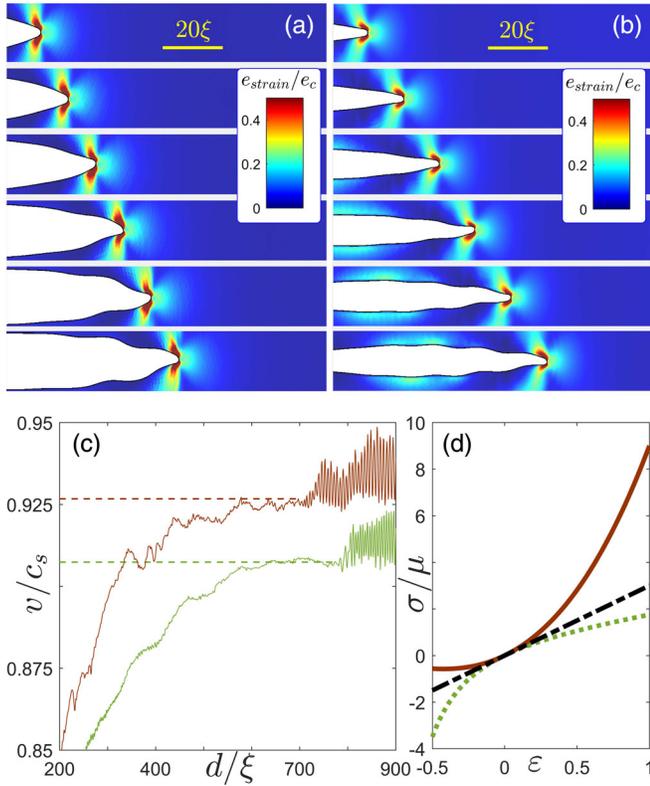


FIG. 1. (a) An enlargement of the 2D oscillatory instability in a phase-field simulation of brittle neo-Hookean (NH) materials under tensile (mode I) loading [14]. The color code corresponds to the normalized strain energy density and ξ is the dissipation length. (b) The same, but for brittle Saint-Venant-Kirchhoff (SVK) materials, see text and [20] for details. (c) Examples of the normalized crack speed v/c_s (c_s is the shear wave speed) vs normalized crack propagation distance d/ξ for SVK (top curve) and NH materials (bottom curve). The onset of oscillations is marked by the horizontal dashed lines. (d) The uniaxial stress σ (normalized by the shear modulus μ) vs strain ϵ for SVK (solid line) and NH materials (dashed line), along with their linear approximation (dash-dotted line).

inside the dissipation zone, thereby allowing cracks to accelerate to unprecedentedly high velocities without undergoing tip-splitting instabilities at marginally dynamic velocities [27,28]. Moreover, $e_{\text{strain}}(\mathbf{H})$ is taken to be nonlinear, introducing nonlinearity on a length ℓ in the near vicinity of the crack tip, while the linear elastic (quadratic) approximation to $e_{\text{strain}}(\mathbf{H})$ remains very good on larger scales. Hence, this theory features an intrinsic dissipation length ξ and intrinsic length ℓ associated with near tip elastic nonlinearity, which can be independently varied.

The strain energy functional $e_{\text{strain}}^{\text{NH}}(\mathbf{H}) = \frac{1}{2}\mu\{\text{tr}(\mathbf{F}^T\mathbf{F}) + [\det(\mathbf{F})]^{-2} - 3\}$ of 2D incompressible NH materials [29], where $\mathbf{F} = \mathbf{I} + \mathbf{H}$ (\mathbf{I} is the identity tensor) and μ is the shear modulus, has been shown [14] to quantitatively predict the experimentally observed oscillatory instability in 2D brittle polymeric elastomers under tensile (mode I) loading

conditions [4,12,13]. Most notably, the oscillations [see example in Fig. 1(a)] emerge at a critical velocity v_c [see the lower horizontal dashed line in Fig. 1(c)] and their wavelength follows $\lambda = \alpha\xi + \beta\Gamma_0/\mu$ [shown by the squares in Fig. 2(a)], both in quantitative agreement with experiments [13,14]. Here, α and β are dimensionless numbers, ξ is the dissipation length defined above, $\Gamma_0 \equiv \Gamma(v \rightarrow 0)$, and Γ_0/μ is proportional to the intrinsic nonlinear elastic length ℓ [4,13,17,19]. $e_{\text{strain}}^{\text{NH}}$ is of entropic origin, which is well understood in terms of the statistical thermodynamics of cross-linked polymer chains [30].

To assess the generality of these results, our first goal is to consider near tip elastic nonlinearity which is sufficiently general, yet of qualitatively different physical origin and emerging properties compared to $e_{\text{strain}}^{\text{NH}}$. To that aim, we invoke the minimal elastic nonlinearity associated with the rotational invariance of isotropic materials. That is, we use the rotationally invariant (Green-Lagrange) metric strain tensor $\mathbf{E} = \frac{1}{2}(\mathbf{F}^T\mathbf{F} - \mathbf{I}) = \boldsymbol{\epsilon} + \frac{1}{2}(\mathbf{H}^T\mathbf{H})$, instead of its widespread linear approximation $\boldsymbol{\epsilon} = \frac{1}{2}(\mathbf{H} + \mathbf{H}^T)$ [31]. When combined with constitutive linearity, i.e., with a quadratic energy functional in which the linearized strain measure $\boldsymbol{\epsilon}$ is replaced by its nonlinear rotationally invariant counterpart \mathbf{E} , we obtain

$$e_{\text{strain}}^{\text{SVK}}(\mathbf{H}) = \frac{1}{2}\tilde{\lambda}\text{tr}^2(\mathbf{E}) + \mu\text{tr}(\mathbf{E}^2). \quad (1)$$

This energy functional, corresponding to Saint-Venant-Kirchhoff (SVK) materials [32], is constitutively identical to linearized Hookean elasticity (where $\tilde{\lambda}$ is the first Lamé constant [31]), but features geometric nonlinearity embedded inside \mathbf{E} .

While materials typically feature constitutive nonlinearity in addition to geometric nonlinearity, they should *at least* feature the latter, and hence Eq. (1) constitutes the minimal possible elastic nonlinearity. $e_{\text{strain}}^{\text{SVK}}$ is not only of a qualitatively different physical origin compared to $e_{\text{strain}}^{\text{NH}}$, but it also exhibits significantly different properties. In Fig. 1(d), we plot the uniaxial tension response corresponding to the two functionals, in addition to their linear elastic approximation (the elastic constants are chosen such that the latter is identical for both functionals). We observe that $e_{\text{strain}}^{\text{SVK}}$ features a stronger nonlinearity than $e_{\text{strain}}^{\text{NH}}$ and is of a strain-stiffening nature, while the latter is of a strain-softening nature. Do materials described by $e_{\text{strain}}^{\text{SVK}}$ experience the same oscillatory instability as those described by $e_{\text{strain}}^{\text{NH}}$?

In Fig. 1(b) we present the results of a large-scale numerical simulation of the nonlinear phase-field fracture theory discussed above, using $e_{\text{strain}}^{\text{SVK}}$ (see simulation details in [20]). An oscillatory instability that is strikingly similar to the oscillatory instability shown in Fig. 1(a) for brittle NH materials is observed. The onset of instability, see example in Fig. 1(c) (top curve), takes place at an ultrahigh critical velocity v_c , whose normalized value is similar to the

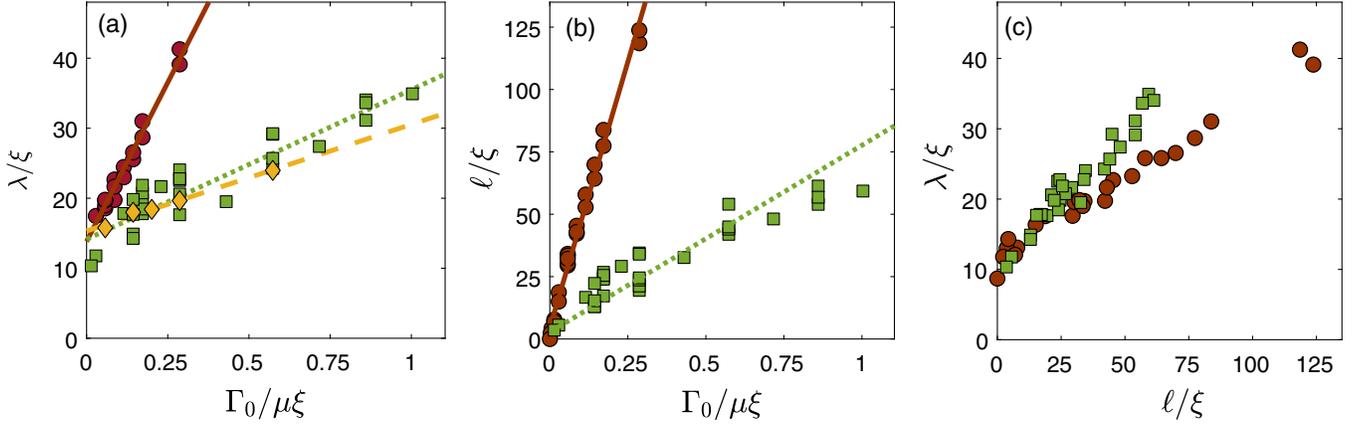


FIG. 2. (a) The normalized oscillations wavelength λ/ξ vs $\Gamma_0/\mu\xi$ for SVK (circles), NH (squares), and truncated NH (diamonds) materials. (b) The normalized nonlinear length ℓ/ξ (ℓ is defined in the text) vs $\Gamma_0/\mu\xi$ for SVK (circles) and NH (squares) materials. (c) λ/ξ vs ℓ/ξ for SVK (circles) and NH (squares) materials. The lines are guides to the eye.

one of brittle NH materials (bottom curve). The fact that the oscillatory instability exists for widely different forms of near crack tip elastic nonlinearity provides a strong indication in favor of its universal nature. In Fig. 2(a), we present the oscillations wavelength of brittle SVK materials demonstrating that it scales linearly with Γ_0/μ , $\lambda = \alpha\xi + \beta\Gamma_0/\mu$, exactly as it does for brittle NH materials, yet again supporting the universality of the oscillatory instability. Remarkably, while β is significantly larger for brittle SVK materials compared to brittle NH materials, $\alpha = 13 \pm 1$ is essentially identical for the two classes of materials.

To address the apparent independence of α on the form of near tip elastic nonlinearity and its physical implications, we consider yet another energy functional. In selecting the latter, we aim at addressing also a different question: Is the wavelength significantly affected by strong elastic nonlinearity deep inside the near tip nonlinear elastic zone or mainly by weak elastic nonlinearity, as predicted by the weakly nonlinear theory of fracture [4,15,17]? To address these questions, we consider the incompressible NH functional truncated to leading order nonlinearity

$$e_{\text{strain}}^{\text{TNH}}(\mathbf{H})/\mu \simeq \text{tr}(\boldsymbol{\epsilon}^2) + \text{tr}(\boldsymbol{\epsilon})^2 - \frac{1}{2}\text{tr}(\mathbf{H})^3 - \frac{3}{2}\text{tr}(\mathbf{H})\text{tr}(\mathbf{H}^2) - \frac{1}{8}\text{tr}(\mathbf{H})^4 + \frac{9}{4}\text{tr}(\mathbf{H})^2\text{tr}(\mathbf{H}^2) + \frac{3}{8}\text{tr}(\mathbf{H}^2)^2 + \mathcal{O}(\mathbf{H}^5). \quad (2)$$

Unlike the weakly nonlinear theory of fracture, which is an analytic perturbative theory that takes into account only cubic nonlinearity in the energy, here we are looking for global numerical solutions and hence should include also quartic nonlinearity to ensure the well-posedness of the global fracture problem [20].

Phase-field theory calculations with $e_{\text{strain}}^{\text{TNH}}$ of Eq. (2) demonstrated the existence of an oscillatory instability (not shown), whose wavelength is essentially indistinguishable from that of $e_{\text{strain}}^{\text{NH}}$, as shown in Fig. 2(a) (diamonds).

This result indicates that weak elastic nonlinearity controls the oscillatory instability and further highlights the independence of α on the form of nonlinearity. The latter raises an intriguing possibility; if elastic nonlinearity vanishes, $\ell \propto \Gamma_0/\mu \rightarrow 0$, the results of Fig. 2(a) imply that $\lambda \approx 13\xi$. That is, the oscillatory instability may exist in the absence of elastic nonlinearity, where its wavelength is determined by the intrinsic dissipation length ξ .

This possibility is quite intriguing because, if true, it means that near tip elastic nonlinearity is not necessary for the existence of the oscillatory instability, which can alternatively inherit its characteristic length from the dissipation zone. The extensive calculations that gave rise to the wavelength λ in Fig. 2(a), where the ratio Γ_0/μ has been reduced as much as possible within numerical limitations, also showed that the *amplitude* of the oscillations diminishes as $\ell \propto \Gamma_0/\mu$ is reduced (see Fig. S3 in [20]). This observation might support the possibility that the instability exists with a finite wavelength and a vanishingly small amplitude as elastic nonlinearity vanishes. Hence, the observation in [14] in which no oscillatory instability is observed when $\ell = 0$, i.e., using the linear elastic (quadratic) approximation of e_{strain} to begin with, may simply be explained in terms of a vanishing amplitude, not as indicating the absence of instability in this limit. On the other hand, based on presently available evidence, we cannot exclude the possibility that the instability disappears at a finite (yet very small) value of $\ell \propto \Gamma_0/\mu$, as further discussed below.

While α appears to be independent of the form of near tip elastic nonlinearity, the slope β in the linear $\lambda-\Gamma_0/\mu$ relation does depend on it. To understand this dependence, recall that the nonlinear length ℓ is proportional to Γ_0/μ , but is not identical to it (e.g., Γ_0/μ exists also in the absence of elastic nonlinearity, while ℓ does not) [4]. Consequently, material dependence (in addition to v dependence) is expected to be encapsulated in the proportionality factor.

To test this possibility, we should directly calculate ℓ , instead of using its dimensional estimate. To calculate ℓ in our large-scale simulations, we need to estimate the actual length at which near tip nonlinearity becomes significant. Any energy functional e_{strain} can be uniquely decomposed into its linear $e_{\text{strain}}^{\text{le}}$ and nonlinear $e_{\text{strain}}^{\text{nl}} \equiv e_{\text{strain}} - e_{\text{strain}}^{\text{le}}$ parts. The region near the tip at which $\|\partial_{\mathbf{H}} e_{\text{strain}}^{\text{nl}}\|/\|\partial_{\mathbf{H}} e_{\text{strain}}^{\text{le}}\|$ becomes non-negligible can be used to define ℓ ($\|\cdot\|$ is the magnitude of a tensor, i.e., the square root of the sum of the squares of its elements). In particular, we define $\ell \equiv \sqrt{\mathcal{A}}$, where \mathcal{A} is the area of the region in which $\|\partial_{\mathbf{H}} e_{\text{strain}}^{\text{nl}}\|/\|\partial_{\mathbf{H}} e_{\text{strain}}^{\text{le}}\| \geq 0.5$ (see Fig. S1 in [20]), consistently with earlier definitions [4]. While the existence of a threshold (here 0.5) involves some degree of arbitrariness in the definition of ℓ , the results themselves do not strongly depend on the exact threshold (see Fig. S2 in [20]).

In Fig. 2(b) we plot ℓ (as just defined) vs Γ_0/μ for both NH and SVK brittle materials. For both types of materials we observe $\ell \propto \Gamma_0/\mu$, as predicted theoretically. Moreover, the prefactor in this relation is significantly larger for SVK materials compared to NH materials, which is consistent with the already discussed stronger nonlinearity of the former. In fact, the variability in the prefactor appears to be similar in magnitude to the variability in the slope of the linear λ - Γ_0/μ relation shown in Fig. 2(a). To test whether the material dependence of the prefactor indeed accounts for the material dependence of the wavelength shown in Fig. 2(a), we plot λ vs ℓ in Fig. 2(c). We observe that the two curves approximately collapse one on top of the other, suggesting that in fact the oscillations' wavelength follows a universal master curve $\lambda = \alpha\tilde{\xi} + \beta\ell$, where α and $\beta = 0.3 \pm 0.07$ are nearly material independent. These results yet again strongly support the universal character of the 2D oscillatory instability.

When the driving force for fracture W , i.e., the stored elastic energy per unit area ahead of the crack, is sufficiently large to allow cracks to accelerate to the critical velocity v_c , the oscillatory instability emerges. Are there additional, previously undiscovered instabilities in 2D dynamic fracture triggered when cracks are driven even more strongly? When W is further increased, there exists a range of driving forces for which steady-state oscillatory cracks exist. As W is increased even more, a tip-splitting instability emerges. It can emerge either from oscillatory crack states (resulting in asymmetric tip splitting, as shown in Fig. 3 and in the Supplemental Material, Movie S1 [20]) or directly from straight crack states (resulting in symmetric tip splitting, as shown in Movie S2 in the Supplemental Material [20]). This ultra-high-velocity tip-splitting instability in 2D dynamic fracture appears to be different from previously studied 2D instabilities occurring at lower, marginally dynamic, velocities [27,28,33–42] and from the microbranching instability in 3D experiments [5]. In Fig. 3(a) we present tip splitting that emerges after

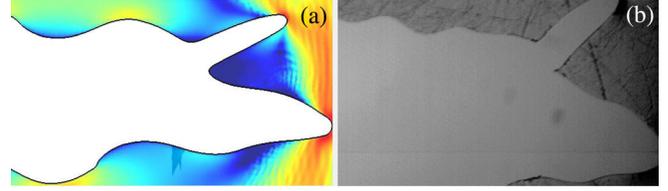


FIG. 3. (a) Oscillations followed by tip splitting in a phase-field simulation of NH materials under a large driving force (see Movie S1 in the Supplemental Material [20]). (b) The corresponding experimental observation (experimental details can be found in [10]).

oscillations (see Supplemental Material, Movie S1 [20]). Remarkably, such oscillations followed by tip splitting have been recently observed under strong driving force conditions in previously unpublished experiments on brittle NH materials, see Fig. 3(b).

We are now in a position to construct a comprehensive stability phase diagram of 2D dynamic fracture. The analysis presented above indicates that the relevant dimensionless parameters for such a stability phase diagram are the ratio between the intrinsic nonlinear scale $\ell \propto \Gamma_0/\mu$ and the intrinsic dissipation scale ξ , and the ratio between the driving force for fracture W and the fracture energy scale Γ_0 . The stability phase diagram in the W/Γ_0 - $\Gamma_0/\mu\xi$ plane is shown in Fig. 4 for both brittle SVK [panel (a)] and NH [panel (b)] materials. For both classes of materials, for a fixed and finite $\Gamma_0/\mu\xi$, the phase diagram exhibits the sequence of transitions with increasing W/Γ_0 described in the previous paragraph; for $W/\Gamma_0 < 1$ no crack propagation is possible (not shown), steady-state straight cracks exist upon increasing W/Γ_0 beyond unity over some range of W/Γ_0 (diamonds), then steady-state oscillatory cracks exist over some range for yet larger values of W/Γ_0 (circles), then tip splitting emerges (squares), either after

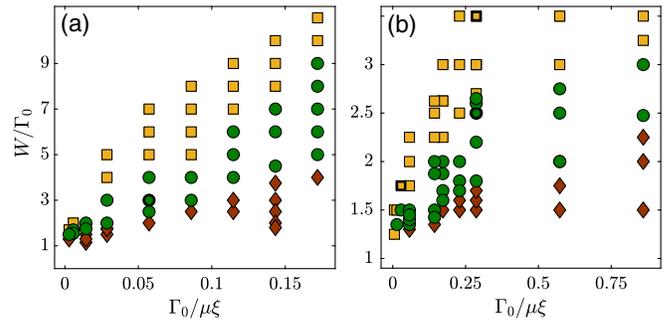


FIG. 4. Stability phase diagram of 2D dynamic fracture in the W/Γ_0 - $\Gamma_0/\mu\xi$ plane for SVK (a) and NH materials (b). Shown are straight cracks (diamonds), oscillatory cracks (circles), and oscillatory or straight cracks undergoing tip splitting (squares). See Supplemental Material movies corresponding to the thick-lined squares in panel (b) [20]. The upper thick-lined square also corresponds to Fig. 3(a). The thick-lined circles in panels (a) and (b) correspond to Figs. 1(b) and 1(a), respectively.

oscillations (asymmetric tip splitting, see Movie S1 in the Supplemental Material [20]) or before (symmetric tip splitting, see Movie S2 in the Supplemental Material [20]).

Interestingly, the W/Γ_0 range over which oscillatory cracks exist decreases with decreasing $\Gamma_0/\mu\xi$. In the limit $\Gamma_0/\mu\xi \rightarrow 0$, it appears to vanish altogether such that straight cracks make a direct transition to tip-split cracks. While present numerical limitations do not allow us to conclude with certainty if the oscillatory instability exists or not in the absence of near tip elastic nonlinearity, they at least show that, if present, this instability would only exist over a vanishingly small range of loading, with a vanishingly small oscillations amplitude. Our results further strongly indicate that minute near tip nonlinearity, which is likely to exist in most materials, is sufficient to dramatically affect the instability whose wavelength is essentially determined by the dissipation length ξ . Finally, and most remarkably, the salient properties and topology of the stability phase diagram appear to be universal; i.e., they are the same for two widely different forms of near tip nonlinearity.

In summary, we highlighted in this Letter the universal nature of the 2D oscillatory instability whose wavelength follows a universal master curve in terms of dissipation and nonlinear elastic intrinsic lengths, demonstrated the existence of a high-velocity tip-splitting instability, and constructed a universal stability phase diagram of 2D dynamic fracture. Future research should address the existence of the oscillatory instability in the absence of near tip elastic nonlinearity and the origin of the high-velocity tip-splitting instability.

This research was supported in part by the U.S.-Israel Binational Science Foundation (BSF) Grant No. 2012061, which provided partial support for C.-H. C. E. B. acknowledges support from the William Z. and Eda Bess Novick Young Scientist Fund and the Harold Perlman Family. A. K. acknowledges the support of Grant No. DE-FG02-07ER46400 from the U.S. Department of Energy, Office of Basic Energy Sciences. We are grateful to J. Fineberg for providing the experimental result shown in Fig. 3(b).

Y. Lubomirsky and C.-H. Chen contributed equally to this work.

*a.karma@northeastern.edu

†eran.bouchbinder@weizmann.ac.il

- [1] L. B. Freund, *Dynamic Fracture Mechanics* (Cambridge University Press, New York, 1998).
- [2] K. B. Broberg, *Cracks and Fracture* (Academic Press, San Diego, 1999).
- [3] E. Bouchbinder, J. Fineberg, and M. Marder, *Annu. Rev. Condens. Matter Phys.* **1**, 371 (2010).
- [4] E. Bouchbinder, T. Goldman, and J. Fineberg, *Rep. Prog. Phys.* **77**, 046501 (2014).
- [5] J. Fineberg and M. Marder, *Phys. Rep.* **313**, 1 (1999).
- [6] K. Ravi-Chandar and W. G. Knauss, *Int. J. Fract.* **26**, 141 (1984).
- [7] J. Fineberg, S. P. Gross, M. Marder, and H. L. Swinney, *Phys. Rev. Lett.* **67**, 457 (1991).
- [8] E. Sharon and J. Fineberg, *Phys. Rev. B* **54**, 7128 (1996).
- [9] A. Livne, G. Cohen, and J. Fineberg, *Phys. Rev. Lett.* **94**, 224301 (2005).
- [10] T. Goldman Boué, G. Cohen, and J. Fineberg, *Phys. Rev. Lett.* **114**, 054301 (2015).
- [11] I. Kolvin, G. Cohen, and J. Fineberg, *Nat. Mater.* **17**, 140 (2018).
- [12] A. Livne, O. Ben-David, and J. Fineberg, *Phys. Rev. Lett.* **98**, 124301 (2007).
- [13] T. Goldman, R. Harpaz, E. Bouchbinder, and J. Fineberg, *Phys. Rev. Lett.* **108**, 104303 (2012).
- [14] C.-H. Chen, E. Bouchbinder, and A. Karma, *Nat. Phys.* **13**, 1186 (2017).
- [15] E. Bouchbinder, *Phys. Rev. Lett.* **103**, 164301 (2009).
- [16] A. Livne, E. Bouchbinder, and J. Fineberg, *Phys. Rev. Lett.* **101**, 264301 (2008).
- [17] E. Bouchbinder, A. Livne, and J. Fineberg, *Phys. Rev. Lett.* **101**, 264302 (2008).
- [18] E. Bouchbinder, A. Livne, and J. Fineberg, *J. Mech. Phys. Solids* **57**, 1568 (2009).
- [19] A. Livne, E. Bouchbinder, I. Svetlizky, and J. Fineberg, *Science* **327**, 1359 (2010).
- [20] See Supplemental Material at <http://link.aps.org/supplemental/10.1103/PhysRevLett.121.134301> for additional information.
- [21] B. Bourdin, G. A. Francfort, and J.-J. Marigo, *J. Mech. Phys. Solids* **48**, 797 (2000).
- [22] A. Karma, D. A. Kessler, and H. Levine, *Phys. Rev. Lett.* **87**, 045501 (2001).
- [23] V. Hakim and A. Karma, *J. Mech. Phys. Solids* **57**, 342 (2009).
- [24] A. J. Pons and A. Karma, *Nature (London)* **464**, 85 (2010).
- [25] R. Spatschek, E. Brener, and A. Karma, *Philos. Mag.* **91**, 75 (2011).
- [26] B. Bourdin, J.-J. Marigo, C. Maurini, and P. Sicsic, *Phys. Rev. Lett.* **112**, 014301 (2014).
- [27] A. Karma and A. E. Lobkovsky, *Phys. Rev. Lett.* **92**, 245510 (2004).
- [28] H. Henry, *Europhys. Lett.* **83**, 16004 (2008).
- [29] J. K. Knowles and E. Sternberg, *J. Elast.* **13**, 257 (1983).
- [30] P. J. Flory and J. Rehner Jr., *J. Chem. Phys.* **11**, 512 (1943).
- [31] L. D. Landau and E. M. Lifshitz, *Theory of Elasticity*, 3rd ed. (Pergamon Press, London, 1986).
- [32] G. A. Holzapfel, *Nonlinear Solid Mechanics* (Wiley, Chichester, 2000).
- [33] F. F. Abraham, D. Brodbeck, R. A. Rafey, and W. E. Rudge, *Phys. Rev. Lett.* **73**, 272 (1994).
- [34] X. P. Xu and A. Needleman, *J. Mech. Phys. Solids* **42**, 1397 (1994).
- [35] M. Marder and S. Gross, *J. Mech. Phys. Solids* **43**, 1 (1995).
- [36] M. Adda-Bedia and M. Ben Amar, *Phys. Rev. Lett.* **76**, 1497 (1996).
- [37] M. Adda-Bedia, R. Arias, M. Ben Amar, and F. Lund, *Phys. Rev. Lett.* **82**, 2314 (1999).

- [38] M. J. Buehler, F. F. Abraham, and H. Gao, *Nature (London)* **426**, 141 (2003).
- [39] M. Adda-Bedia, *Phys. Rev. Lett.* **93**, 185502 (2004).
- [40] E. Bouchbinder, J. Mathiesen, and I. Procaccia, *Phys. Rev. E* **71**, 056118 (2005).
- [41] R. Spatschek, M. Hartmann, E. Brener, H. Muller-Krumbhaar, and K. Kassner, *Phys. Rev. Lett.* **96**, 015502 (2006).
- [42] D. Pilipenko, R. Spatschek, E. A. Brener, and H. Muller-Krumbhaar, *Phys. Rev. Lett.* **98**, 015503 (2007).