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Dissipative dynamic fracture of disordered systems

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Breakdown of two-dimensional disordered systems is studied with a time-dependent network model. The dependence of fracture process on the local relaxation of the force field is included within the framework of Maxwellian viscoelasticity. The dynamics and characteristics of crack formation and propagation are shown to depend on disorder and relative time scales of dissipation and loading. Brittle behavior is encountered in the adiabatic limit of slow straining. At finite strain rates, the development of damage shows ductile behavior with increasing dissipation. Nucleation of cracks in various dynamical situations is discussed.

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The development of failure in a medium depends on defects and disorder, initially present. In such far-from-perfect systems, brittle failure has received a lot of attention due to intrinsic technological interest [1]. The strength of a material depends also on its ability to withstand damage, which is determined by the rate small scale defects grow and get connected. The final fracture occurs when a loaded specimen has a crack that has grown over a critical size. Many real materials, such as composites and polymers, display time-dependent and dissipative behavior under stress. Such materials are called viscoelastic, and they show creep and relaxation of applied stress [2] and stress-strain behavior that depends directly on the loading rate. The microscopic reasons for viscoelastic and reversible elastic responses differ, and in polymer melts, for example, the cause is believed to be the slow readjustment of the microscopic constituents to the applied stress. It is similar to the dynamics of dislocations, which underlies plasticity in the stress-strain characteristics of metals.

In this Rapid Communication we analyze the interplay between disorder and dissipation in dynamic fracture with a simple time-dependent mesoscale model. The beginning and development of failure have usually been studied with discrete lattice models [1,3], in which the strain is applied adiabatically. In this case the simulations are conducted in a way that corresponds to a slow rate of straining; the local stress field is allowed to readjust itself to the changes in the environment over a scale of the correlation length. Hence such quasistatic models exhibit no real dynamics, but a series of equilibrium states.

Our model is constructed to be initially a square lattice, each site being assigned a unit mass, whose dynamical behavior will be monitored. As in usual quasistatic lattice models for fracture, the sites are connected by bonds. The elastic interaction strength of a bond \( ij \) between nearest neighbor sites \( i \) and \( j \) is described by the Born Hamiltonian [4]

\[
H_{ij} = \frac{\alpha}{2}([\hat{u}_i - \hat{u}_j] \cdot \hat{d}_j)^2 + \frac{\beta}{2}([\hat{u}_i - \hat{u}_j] \cdot \hat{d}_j)^2, \tag{1}
\]

in a system of size \( L \times L \). The displacement vector of site \( k \) is denoted by \( \hat{u}_k \), and \( \hat{d}_j \) stands for the unit vector, either parallel or perpendicular to the vector connecting the sites in the undeformed lattice. Periodic boundary conditions are applied perpendicular to the external loading, as depicted in Fig. 1. The Born potential, defining a noncentral force system, is the simplest one in describing angular stiffness, but it is computationally less demanding compared with various bond-bending Hamiltonians [5]. It also allows the study of fairly large system sizes, up to \( 10^5 \) particles. In our model, disorder is introduced as quenched, i.e., by removing a fraction of bonds at random with probability \( (1-p) \).

Apart from the energetics described above, the choice for the dynamics of the system plays a crucial role in modeling fracture. This is because simultaneously a time scale is introduced. The dynamics of site-spring models has been analyzed earlier in some special cases [6], and it is closely related to discrete cellular automata and Hamiltonian systems [7]. The classical molecular dynamics simulations of fracture are also closely analogous [8,9]. Recently, some time-dependent lattice models of fracture have been devised [10–12] in which the time scale arises from stochastic rules, or is, in other words, externally dictated. In our model, a dissipation mechanism is introduced to the microscopic forces in order to mimic the local adjustment of the material structure to the stress. This is done through the classical Maxwellian viscoelasticity [13], which allows the description of relaxation and dissipation of elastic energy as a dynamical decay of the local forces. The constitutive equation for the forces acting at each bond is taken to be

\[
\begin{align*}
\text{FIG. 1.} & \text{ Description of the model. Neighboring mass sites are coupled with dissipative Born springs, parametrized with } \alpha \text{ and } \beta. \text{ The dynamics of the local bond forces are given by Eq. (2). A bond exists with probability } p. \\
\end{align*}
\]
steady-state stress, which is achieved with sufficient straining. In the limit of a vanishing strain rate and dissipation, the ensuing force-elongation characteristics should be equal to those of a corresponding adiabatic model [14,15].

Next we investigate the dependence of the fracture process on the strain rate and internal dissipation via the $\tau$ parameter. General features of fracture are shown in the inset of Fig. 2, in which a 100x100 system is strained up to the point of failure. The initial bond dilution was chosen to be 30%, i.e., $p = 0.7$, in the simulation. The calculation is stopped once the system becomes mechanically disconnected, which is checked after each bond breakage. As a fracture criterion we use the elongation of a bond: it is broken if its length exceeds a critical value, here taken as 1% of the original bond length. Note that the beginning of fracture in an “ideal” force-elongation curve is roughly given by the chosen fracture length (cf. Fig. 2). After the onset of fracture, the decrease in the force-elongation curve shows fluctuations. This is caused partly by the relaxation of the internal elastic energy contained in the lattice, and partly by disorder.

Compared to the linear dependence between local stress and strain in adiabatic models, our choice for the constitutive relation implies that the local stress depends not only on the present strain, but on the whole preceding strain history. The fracture condition in this work can be compared with that used by Sornette and Vanneste (SV), whose lattice model is based on the rupture of fuse elements, when the local temperature exceeds a critical value [12]. In that case the temperature field controlled by the individual fuse currents is analogous to the plastic strain field. However, in our model the local strain is directly responsible for the onset of fracture. In both cases, the instantaneously high stresses, or currents in the SV model, do not necessarily break the bond, but the “memory” from previous deformations—included in the bond length, or fuse temperature in the SV model—may cause the rupture at low local stress levels. In interpreting our simulations, it should also be remembered that the redistribution of internal stresses occurs with a finite speed. This is in contrast to the quasistatic models.

For a moderate to large strain rate and in a slightly diluted system, two different regimes can be seen in the fracture process. The increase in damage during the elongation can be used to demonstrate this. Figure 3 shows, for a system with $p = 0.9$, how the dissipation constant $\tau$ affects the time development of the number of bonds cut during straining, $N_c$. For a small enough $\tau$, $\tau = 1$ at $d e/d t = 2.5 \times 10^{-4}$, the proportion of ruptured bonds grows rapidly in the beginning of the fracture history. But then the accumulation of damage saturates and the system is able to withstand additional straining. This can be attributed to local viscoelastic dissipation, which arrests the growth of the crack. Note that during loading, the distribution of strain in the network shows a clear bias towards the strained edge, if the strain rate is high enough. Thus, in spite of disorder, the bonds near the strained edge have a higher probability to reach the fracture condition first [16]. For a larger value of $\tau = 10$, one observes that the time dependence of $N_c$ is different. After the initiation of fracture, isolated bond breakages build up slowly, until the final rupture phase, during which most of the bonds are broken in a correlated manner [17]. In Fig. 3 the latter behavior is seen as a steeper increase in $N_c$. This can be
interpreted as the start of the crack propagation phase. In general, at finite strain rates and a large $\tau$, there can exist competing microcracks, which have nucleated from several different original defects—not only from the most critical one [18], as is common in models of brittle fracture with fixed disorder. The microcracks exhibit correlated growth and can eventually coalesce. This illustrates how fracture depends on the dynamical evolution of the individual mass sites.

Judging from the width of the fracture regime in strain or strain to final fracture (in Fig. 3), the system shows increased ductility for small $\tau$ or non-negligible dissipation. In the limit of vanishing strain rate, the differences between the two fracture regimes described above become obscure, and the fracture dynamics become eventually identical despite a varying degree of dissipation, as shown in the inset of Fig. 3. The phenomenon of the fracture behavior being independent of the dissipation in the case without almost any inertia for very slow strain rates can be attributed to the fact that the local dissipation does not change much the ordering of the internal forces, which in turn dictates the local dynamics. The almost identical morphologies of the corresponding cracks shows that the adiabatic behavior of quasistatic lattice models [14,19] is encountered in this limit. We emphasize the difference between the fracture process and the load history with varying dissipation and strain rate. The former shows signs of generality stemming from the decisive role of the quenched disorder, while the latter macroscopic behavior stems from the Maxwellian viscoelastic dynamics of individual bonds.

Figure 4 shows how the individual fracture events occur in the direction perpendicular to the external strain. For the propagation speed of microcracks we have found a rough dependence $v_{\text{crack}} \approx \beta$. In our model, there seems to be no evidence of crack bifurcation in the form exhibited by some recent models of crack dynamics [20], which may be caused by the differences in symmetry and loading arrangements of the system. On the other hand, it is known from studies of the quasistatic Born model in the adiabatic limit that branching and crack arrest can be controlled with the $\alpha/\beta$ ratio [14]. This has its roots in the interactions between rows of springs parallel to the applied strain. For small values of $\alpha/\beta$, the fracture path becomes quite rough. However, we have chosen a moderate value in the sense that in the adiabatic case the expected cracks are rather straight in character [14]. The increased ductility for $\tau=1$ compared with $\tau=10$, shown in Fig. 3, is obvious also in Figs. 4(a) and 4(c). However, decreasing $\beta$ in Figs. 4(b) and 4(d) has a similar effect. Now it is easy for bonds to rotate to load-bearing directions, which increases the ability of the system to withstand further straining. It is also seen that the number of simultaneously propagating microcracks grows, which is attributable to the diminished interactions between the rows of bonds. Thus, concerning ductility and elastic stiffness of the systems, the $\alpha/\beta$ ratio has a similar effect on mechanical properties as dissipation: increased ductility corresponds to a decrease in the capability of the system to store elastic energy. If the disorder is increased, i.e., $p$ decreases, the picture emerging from our model follows that seen in adiabatic simulations: the wandering of the final crack becomes more erratic, as the breakage of distant, individual bonds is sufficient to bring about the final fracture. The ductility and elastic modulus decrease with increasing disorder as happens in adiabatic models. The detailed study of crack statistics in the presence of branching and arrest effects is left for future study.

The main conclusion of our dissipative dynamical model is the appearance of different regimes in the fracture characteristics, when increasing the strain rate. These regimes differ in many aspects, as can be seen in the dynamics of crack growth. The most striking features are the dynamics of the final crack and the characteristics of fracture ductility. Far from the adiabatic limit, dissipation determines the ductility...
of the system. Although a strongly dissipative system suffers
damage at lower external strains than a weakly dissipative
one, it is, however, able to accommodate more external
strain, when breaking down. The differences compared to the
adiabatic models arise solely from the nonlinear dissipative
response of the system. This stems from the microscopic
dynamics of individual sites together with the chosen frac-
ture criterion, and is reflected in the macroscopic behavior of
the system. In the opposite limit of slow dynamics, the frac-
ture characteristics of our model approach those of standard
adiabatic models. Though beyond the scope of this study, an
important question arises: how do stress waves generated by
individual bond breakages affect the fracture itself? It can be
expected that, in addition to disorder, the boundary condi-
tions of the system are crucial. Finally, in judging the results,
the fact should be considered that disorder in our model is
assumed independent of the viscoelastic relaxation pro-
cesses. This can be expected to be true in the case of physical
systems with natural disorder at a mesoscopic length scale
dissipation on a molecular scale, as in such a viscoelastic
material as paper. On the other hand, in polymeric materials
the situation may be different.

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