

1 **The initiation of frictional motion - the nucleation**
2 **dynamics of frictional ruptures**

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5 **Key Points:**

- 6 • Frictional ruptures initiate via a characteristic nucleation process that triggers dy-
7 namic rupture essentially
8 • Nucleation replaces the concept of a ‘static friction coefficient’
9 • The nucleation process possesses unique characteristic general properties
10 • Nucleation details depend on local topography

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Abstract

Frictional interfaces lose stability via earthquake-like ruptures, which are close analogues of shear cracks that are well-described by fracture mechanics. Interface ruptures, however, need to be first formed - or nucleated. Rupture nucleation therefore determines the onset of friction, replacing the concept of a characteristic ‘static friction coefficient’. Utilizing rupture arrest at an imposed barrier, we experimentally determine nucleation locations, times and stresses at the origin of each subsequent rupture event. This enables us study the nucleation process via real-time measurements of real contact area and local strain. Nucleation events initiate as 2D patches that expand at nearly constant velocities, v_{nuc} , that are orders of magnitude lower than the dynamic rupture velocities described by conventional fracture mechanics. We find that: (1) Nucleation has location-dependent stress thresholds, (2) v_{nuc} is roughly proportional to the local stress level, (3) the nucleation process continues until the patch size reaches $L_{tran} \sim L_G$, the Griffith length for the onset of dynamic fracture (4) scaling time by $\tau = L_{tran}/v_{nuc}$, nucleation patches exhibit self-similar dynamics (5) dynamic ruptures’ cohesive zones are not fully established until significantly beyond L_{tran} . Many details of nucleation are governed by the local contact area topography, which is roughly invariant under successive rupture events in mature interfaces. Topography-dependent details of the nucleation process include: precise nucleation site location, patch geometry, critical stress thresholds and the proportionality constant of v_{nuc} with stress. We believe that these results shed considerable light on both how frictional motion is triggered and earthquake initiation.

Plain Language Summary

Recent experiments have demonstrated that rapid rupture fronts, akin to earthquakes, mediate the transition to frictional motion. Moreover, once these dynamic rupture fronts (“laboratory earthquakes”) are created, their singular form, dynamics and arrest are well-described by fracture mechanics. Ruptures, however, need to be created within initially rough frictional interfaces, before they are able to propagate. This is the reason that “static friction coefficients” are not well-defined; frictional ruptures can nucleate for a wide range of applied forces. A critical open question is, therefore, how the nucleation of rupture fronts actually takes place. We experimentally demonstrate that rupture front nucleation is prefaced by slow nucleation fronts. These nucleation fronts, which are self-similar, are not described by our current understanding of fracture mechanics. The nucleation fronts emerge from initially rough frictional interfaces at well-defined stress thresholds, evolve at characteristic velocity and time scales governed by stress levels, and propagate within a frictional interface to form the initial rupture from which fracture mechanics take over. These results are of fundamental importance to questions ranging from earthquake nucleation and prediction to processes governing material failure.

Introduction

The initiation of ‘stick-slip’ frictional motion is caused by propagating rupture fronts that detach the ensemble of discrete contacts that form a frictional interface. When occurring along a natural fault that is sandwiched between tectonic plates, this rupture process and the associated release of energy describes an earthquake (Byerlee & Brace, 1968; Scholz, 2019). These fronts have been recorded and experimentally studied over the past two decades in both brittle plastics (Rubinstein et al., 2004; Xia et al., 2004; Nielsen et al., 2010; Schubnel et al., 2011) and in rock (Wu & McLaskey, 2019; Xu et al., 2018; Passelègue et al., 2013). These rupture fronts behave precisely like shear cracks; the framework of fracture mechanics (LEFM or linear elastic fracture mechanics) fully describes rupture front characteristics, such as their propagation dynamics (Svetlizky, Kammer, et al., 2017; Passelègue et al., 2020; Kammer & McLaskey, 2019), arrest conditions (Bayart

et al., 2016, 2018), and the stress fields surrounding them (Svetlizky & Fineberg, 2014; Xu et al., 2019; Kammer & McLaskey, 2019; Mello et al., 2016).

Experiments have, moreover, demonstrated (Ben-David, Cohen, & Fineberg, 2010; Svetlizky, Kammer, et al., 2017) that a frictional system at constant nominal conditions (e.g. normal stress) will be stable to rupture for a large range of imposed initial shear stresses. Were an interface wholly homogeneous, applied stresses could be large enough to approach the maximal shear stress of a given material before the interface would ‘fracture’. According to LEFM, in order for an interface to be unstable to fracture and propagate, it first requires an initial rupture or ‘seed’; if no initial ‘seed’ exists within an interface, then the extreme (putatively singular) stress amplification that forms at a rupture tip could not take place, and ruptures would not be excited. Furthermore, this initial seed needs to surpass a critical length, called the Griffith length, L_G . LEFM predicts that a system under stress will, therefore, remain stable until an initial rupture of sufficient length is either imposed or somehow develops whose length exceeds L_G . L_G is a well-defined function of both the applied stress and system geometry; roughly speaking, the larger the imposed stress, the smaller L_G . The experimental observations of fault stability over a very wide range of initial shear stresses simply imply that a range of initial ‘seed ruptures’ exist; for a given ‘seed’ size, imposed stresses need to be sufficiently large to reduce L_G to this size. Rupture nucleation, or the formation and growth of this initial rupture within the random rough surface that characterizes a frictional interface, is, therefore, the process that *determines* when and how rupture onset will initiate in initially stressed plates.

In effect, the rupture nucleation process replaces the concept of a ‘static friction’ coefficient. This simple criterion for stick-slip initiation, the idea that a material characteristic ‘static friction coefficient’ determines whether frictional motion will occur has long been known (Rabinowicz, 1951) to be invalid. Experiments have shown that the same system can nucleate at very different initial stress levels (Ben-David, Cohen, & Fineberg, 2010; Ben-David & Fineberg, 2011; Passelègue et al., 2020). Rupture propagation, structure, and dynamics, which are all described by LEFM, are largely independent of any characteristic frictional resistance of the interface, so long as this resistance does not rapidly vary with slip velocity (Barras et al., 2020; Palmer C. & Rice R., 1973). We will demonstrate that the *local* conditions at the exact location and time of nucleation, and not a *global* ‘friction law’, will determine the stress at which a rupture nucleates. Once nucleation takes place, the stress distribution along the interface will then govern a rupture’s eventual characteristics (Bayart et al., 2016, 2018; Ben-David, Cohen, & Fineberg, 2010; Svetlizky, Kammer, et al., 2017).

The nucleation problem is an important and open question in both the friction and fracture fields, and is crucial for the understanding of earthquakes. Below L_G , it is unclear how a small defect will grow to reach this critical length. The nucleation stage in laboratory studies of friction has been described in many studies (Ohnaka & Shen, 1999; Latour et al., 2013; Dresen et al., 2020; Popov et al., 2010; Lapusta & Rice, 2003; Uenishi & Rice, 2003; Gori et al., 2021; Hulbert et al., 2019; Leeman et al., 2018; Marone, 2019) to be slow and ‘aseismic’. This is in contrast to the propagation stage, in which ruptures rapidly accelerate to high velocities and therefore emanate ‘seismic’ radiation. The nucleation stage is, therefore, often assumed to be qualitatively different from the dynamic rupture process. The empirical distinction between this slow process and slow rupture fronts that are described by the framework of fracture mechanics (Svetlizky, Kammer, et al., 2017; Rubinstein et al., 2007) (i.e. with crack lengths that are barely **above** the Griffith length) is rather elusive (Leeman et al., 2018). While rupture around L_G is one possible scenario for slow ruptures, it is certainly not the only possible mechanism. This important issue is still entirely open.

Several studies (Ripperger et al., 2007; Uenishi & Rice, 2003; Schär et al., 2021) have attributed the nucleation process to heterogeneity of the applied stress field. In this

114 view, stress heterogeneity may initiate local slip events at locations where an assumed
 115 peak strength value is surpassed. When a small slipping patch is artificially created be-
 116 yond a critical length, an instability is reached and the initial patch expands spontaneously,
 117 leading to a rupture process and macroscopic sliding. This can happen with a single patch
 118 (Uenishi & Rice, 2003), or through a coalescence of several small patches (Schär et al.,
 119 2021). Other studies have proposed an avalanche-like process at the microscopic asper-
 120 ity level (de Geus et al., 2019), or the nucleation and propagation of excited slip pulses
 121 (Brener et al., 2018) as possible mechanisms leading to the nucleation of rapid ruptures.

122 Despite the importance of this process, only a relatively few detailed experimen-
 123 tal descriptions of the nucleation process exist (Ohnaka & Shen, 1999; Latour et al., 2013;
 124 Guérin-Marthe et al., 2019; Fukuyama et al., 2018; McLaskey, 2019; Gvirtzman & Fineberg,
 125 2021; Cebry & McLaskey, 2021; Gori et al., 2021). The reason for this lies in the very
 126 nature of the nucleation process - it is unpredictable both in space and in time, and ne-
 127 cessitates closely following the dynamics of slip at relatively small scales. Therefore, con-
 128 ducting a controlled experiment that will be able to describe the process in detail is ex-
 129 tremely challenging. For this reason, our knowledge of nucleation in the laboratory has
 130 often stemmed from examining spontaneous events that ‘happen’ to nucleate in a con-
 131 venient location along the interface under study. While some features of this process can
 132 be understood by these means, the initial conditions at the nucleation location are gener-
 133 ally difficult to precisely define. For example, some studies noted the significance of
 134 the rate of the nucleation process (Xu et al., 2018; Guérin-Marthe et al., 2019), but the
 135 resulting timescale could not be measured directly, since no clear starting point could
 136 be experimentally detected. As a result, in these studies the full ‘nucleation time’ could
 137 not be defined.

138 Recently, we developed a novel experimental approach that utilizes a ‘remote trig-
 139 gering’ method, which can overcome these difficulties (Gvirtzman & Fineberg, 2021). This
 140 approach enables us to conduct sequences of experiments where both the nucleation point
 141 is pre-determined and the local stress at the rupture nucleation point is measured. Pre-
 142 cise control of the nucleation location enables us to gain a detailed description of nucle-
 143 ation processes by imaging the nucleation process in real time with sufficient spatial and
 144 temporal resolution. This work demonstrated that rupture nucleation is a 2D process
 145 that is not described within the current framework of fracture mechanics. The 2D spa-
 146 tial character of the nucleation zone has also been observed in recent experiments (Fukuyama
 147 et al., 2018; McLaskey, 2019) in rock. This work highlighted the importance of trans-
 148 verse rupture dynamics within the ‘width’ of the interface during the nucleation process.
 149 To address this issue, we utilized our optical measurement of the interface to image the
 150 *full* 2D evolution of the nucleation process. Moreover, in (Gvirtzman & Fineberg, 2021)
 151 the nucleation time was shown to depend on the local stress level, up to a threshold be-
 152 low which nucleation does not occur.

153 In this paper, we will describe the detailed characteristics of the nucleation pro-
 154 cess. In particular, we will show that all nucleation events share basic features, such as
 155 self-similar dynamics. The details of the nucleation process can significantly vary with
 156 the nucleation location, and we will demonstrate that this dependence is related to the
 157 local ‘topography’ at the nucleation site. Moreover, we will show the details of rupture
 158 evolution as the transition from nucleation to dynamic rupture comes about.

159 **Experimental Apparatus**

160 *Sample construction:* As in previous studies (Svetlizky & Fineberg, 2014; Ben-David,
 161 Cohen, & Fineberg, 2010; Bayart et al., 2016; Berman et al., 2020; Gvirtzman & Fineberg,
 162 2021), we use 2 PMMA blocks to construct a frictional interface (Fig. 1). The x, y, z di-
 163 mensions of the top (bottom) block are 200,100,5.5 (290,28,30)mm, respectively. The lon-
 164 gitudinal and shear wave velocities of the blocks were ultrasonically measured (Shlomai

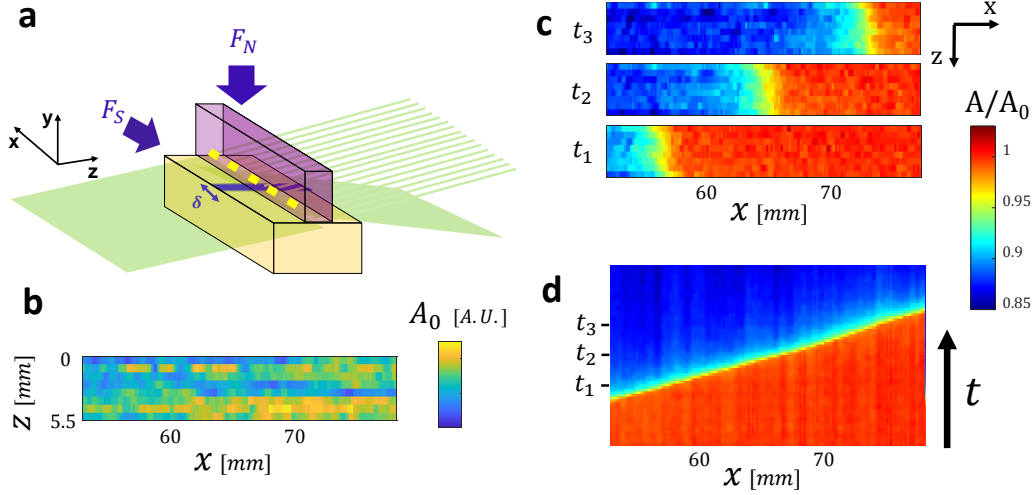


Figure 1. Experimental Setup for Nucleation Experiments. (a) A frictional interface is formed by pressing 2 PMMA blocks together with a normal force, F_N . Stick-slip behavior ensues by applying a shear load, F_S . A thick blue line marks an imposed barrier of high fracture energy at a localized position within the interface. Yellow squares denote strain gauges that are placed slightly above ($y \sim 3.5\text{mm}$) the frictional interface. The interface is illuminated by a sheet of light that is incident at an angle well beyond that of total internal reflection at a PMMA-air interface. Light transmitted through the upper block is then roughly proportional to the real area of contact, $A(x, z, t)$ at each spatial location. (b) Measurements of (the *un-normalized*) $A(x, z, t)$ of a section of the 2D interface. Colors represent the intensity of light that is transferred through each pixel. As the light intensity corresponds to the total area of the contacts within each pixel, these un-normalized measurements provide us with a map of the ‘topography’ of the real contact area, $A(x, z, t)$. (c) 3 snapshots of the *normalized* contact area, $A(x, z, t)/A_0(x, z)$ of the section in (b) at different times; each snapshot is normalized by its value, $A_0(x, z)$ long before rupture took place. The images are obtained at intervals $t_{(i+1)} - t_i = 20\mu\text{s}$. Sequential images throughout an experiment enable tracking of the propagating rupture front. (d) 1 dimensional representation of the interfaces are obtained by averaging over the width of the interface in the z axis. Each 2D snapshot, therefore, corresponds to a single row in the spatio-temporal picture. A rupture front is the boundary between the broken and unbroken contacts. The temporal locations of the snapshots in (c) are indicated.

165 et al., 2020) under plane strain conditions to obtain values of 2680 and 1361m/s, respec-
 166 tively, with an error of 10 m/s. These values yield a Rayleigh velocity of 1255 ± 10 m/s.
 167 The density of PMMA was measured to be $1170 \pm 10 \text{ kg m}^{-3}$, and the Poisson ratio is
 168 0.3300 ± 0.0007 . PMMA (Read et al., n.d.) is viscoelastic, with a dynamic Young’s mod-
 169 ulus of 5.75 ± 0.15 GPa and a static value of 3.62 ± 0.3 GPa.

170 *Loading system:* The upper block was clamped at its upper edge and was pressed to the
 171 bottom block with a normal force F_N in the y direction with a mean normal stress of
 172 4.5 MPa (Fig 1a). F_N was held constant throughout all experiments. The bottom block
 173 was mounted on a low-friction sliding stage, which was quasistatically loaded in the $-x$
 174 direction by applying a shear force, F_S , to produce stick-slip sequences, in which spon-
 175 taneous rupture events nucleated near the $x=0$ edge and propagated in the x direction.
 176 Applied forces, F_N and F_S , were measured by application of these forces in series with
 177 load cells having accuracies better than 1Nt.

178 *Real contact area measurements:* We used a high-power blue LED (CBT-120) to illumi-
 179 nate the entire interface at an incident angle (70°) that was well beyond the total in-
 180 ternal reflection angle (41.8°) between PMMA and air. In this way, light only passes through
 181 contact points, and therefore the intensity of transmitted light is roughly proportional
 182 to the real contact area at each point (Rubinstein et al., 2004). To visualize the entire
 183 interface, we utilized a fast camera (Phantom V710) whose 1280×8 frames were mapped
 184 to pixels of size $165 \times 688 \mu\text{m}$ in the x and z directions, respectively. The real contact
 185 area, $A(x, z, t)$, was, by this method, continuously measured at a rate of ~ 580000 fps,
 186 enabling us to track the propagation of fast rupture fronts over the entire interface by
 187 visualizing the changes of $A(x, z, t)$. Rupture visualization was accomplished by normal-
 188 izing each spatial point by its value, $A_0(x, z)$, long before the event started; $A(x, z, t)/A_0(x, z)$
 189 (Fig. 1c). The non-normalized (raw) measurement, $A_0(x, z, t)$, is a map of the local dis-
 190 tribution of contacts, or the local ‘topography’ (Fig. 1b). *1D* dynamics, $A(x, t)/A_0(x)$
 191 was followed by averaging over the interface width, in the z axis (Fig. 1d). Normaliza-
 192 tion of $A(x, z, t)$ is necessary since the changes in A surrounding the rupture nucleation
 193 process are quite small (in general below $\sim 2\%$), so, without such normalization, are ef-
 194 fectively masked by the underlying topography.

195 *Construction of a local barrier to rupture propagation:* A barrier to rupture propagation
 196 is introduced at specific locations along the interface by locally increasing the fracture
 197 energy. This is achieved by painting the interface with a permanent marker (Staedtler;
 198 size M). The (blue) marker used was transparent to the incident (blue) light used to mea-
 199 sure $A(x, z, t)$. The marker increased (Gvirtzman & Fineberg, 2021) the local fracture
 200 energy by about a factor of 5. This was sufficient to arrest rapid spontaneously prop-
 201 agating ruptures that encountered it, as, locally, the energy flux to the rupture front was
 202 insufficient to overcome the increased fracture energy (Bayart et al., 2016). In the ex-
 203 periments presented here, we created barriers of widths 1-4 mm. After each experimen-
 204 tal sequence, the barrier was removed by cleaning with isopropanol, and, generally, drawn
 205 at a new location. While the increase in the fracture energy due to the marker can be
 206 measured, the precise mechanism for this increase is still unknown. In general, markers
 207 are composed of minute dye particles that are chemically linked to the PMMA. They are
 208 also immersed in a solvent that dries after application. Under the huge pressures sur-
 209 rounding any surface contacts, it is difficult to say whether the increase in fracture en-
 210 ergy is due to slight adhesion, due to the linker molecules, a granular (gouge-like) effect
 211 due to friction of the particles or a softening of the PMMA in a sub-micron layer below
 212 the interface resulting from the solvent.

213 *Induced stress by rupture arrest:* Ruptures, which initiated spontaneously at the sam-
 214 ple edge ($x = 0$), immediately arrested, upon their arrival at a barrier. From their ve-
 215 locity immediately preceding arrest, v_{arr} , one can extract the static stress intensity fac-
 216 tor, $K_S = K(v = 0)$, of the arrested crack as follows. We used the ruptures’ equation
 217 of motion immediately prior to arrest, as described by LEFM (Freund, 1998; Svetlizky,

218 Kammer, et al., 2017):

$$\Gamma = G_s \cdot g(v_{arr}) = K_S^2 \cdot (1 - \nu^2) / E \cdot g(v_{arr}) \quad (1)$$

219 where $g(v)$ is a known dynamic function (Freund, 1998), E and ν are, respectively, the
 220 dynamic Young's modulus and Poisson ratio, and G_s is the static energy release rate.
 221 It is important to note that $g(v)$ is a monotonically decreasing function of v such that
 222 $g(0) = 1$ and $1 \geq g(v) > 0$. The value of the fracture energy outside the barrier, Γ ,
 223 was measured to be 1 J/m^2 , for our experimental conditions (Svetlizky & Fineberg, 2014;
 224 Gvirtzman & Fineberg, 2021). Eq. 1 enabled us to determine the static stress intensity
 225 factor, K_S . The static stress intensity factor, K_S , was determined by using Eq. 1, while
 226 inputting both the measured values of Γ and v_{arr} to provide us with the value of $g(v_{arr})$.
 227 The resulting stress field ahead of the arrested crack is therefore:

$$\Delta\sigma_{xy}(v_{arr}, x) = K_S \cdot (2\pi x)^{-1/2} = \sqrt{E\Gamma/(1 - \nu^2)} \cdot g(v_{arr})^{-1/2} \cdot (2\pi x)^{-1/2}. \quad (2)$$

228 In Eq. 2, the predicted stress field is $\Delta\sigma_{xy} = \sigma_{xy} - \sigma_{xy}^{res}$, where σ_{xy}^{res} is the residual shear
 229 stress value that remains after the passage of a rupture (Bayart et al., 2016). We define
 230 the induced stress at the nucleation point as $\sigma_{ind} = \Delta\sigma_{xy}(r^*)$, where r^* is the distance
 231 between the arrest and nucleation locations, x_{nuc} .

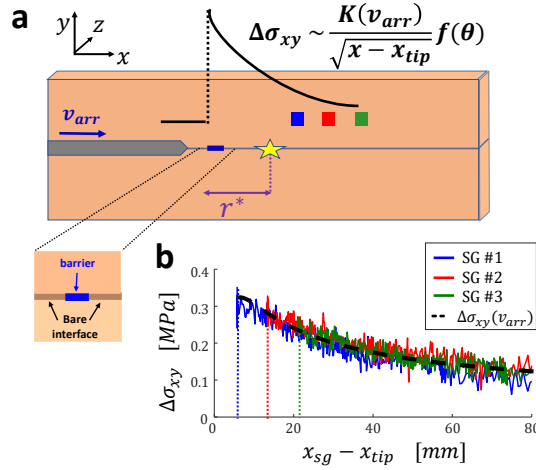


Figure 2. Validation of induced stress calculation. (a) Schematic description of the induced stress field, σ_{ind} , resulting from ruptures arrested at a barrier. The singular stress field induced by an arrested rupture, $\sigma_{ind} = \Delta\sigma_{xy}(r^*)$, was calculated using Eq. 2 using its velocity, v_{arr} , upon arrival at the barrier. This enables calculation of the stress at the nucleation point (star) located a distance r^* from the point of arrest. (b) Comparison of the calculated stress changes $\Delta\sigma_{xy}(v)$ (dashed line) induced by a moving and arrested rupture to stresses obtained via strain measurements at 3 locations, x_{sg} , along and above the interface relative to the instantaneous rupture tip locations, x_{tip} (Svetlizky & Fineberg, 2014; Freund, 1998). Vertical lines: distances between SG locations, x_{sg} , and rupture arrest point. Note that the measurements appearing beyond the vertical lines are stresses measured prior to the rupture arrest at the barrier.

232 While stresses could not be directly measured on the interface (the strain gages could
 233 only be mounted 3mm away from the interface), the values of the induced stresses pre-

234 dictated by Eq. 2 were verified by direct measurements of the stress changes, during rup-
 235 ture propagation and upon rupture arrest, at strain gages located, slightly above the in-
 236 terface, at locations $x > x_{nuc}$. Comparison of the measured σ_{xy} at the first three strain
 237 gages located beyond the barrier with the computed induced stress from the rupture tip
 238 is presented in Figs. 2a,b. The agreement between measured and computed values is ex-
 239 cellent, and justifies our use of the computed values to obtain σ_{ind} .

240 Nucleation Experiments

241 A typical sequence of stick-slip events is shown in Fig. 3b in which a barrier was
 242 imposed at the point $x = 100\text{mm}$. In each event, spontaneous rupture fronts nucleated
 243 near the sample edge at $x = 0$ and propagated in the x direction until reaching the bar-
 244 rier created by the application of our marker. In the experiments considered, rupture fronts
 245 instantaneously arrested upon arrival at the barrier. This behavior is predicted by LEFM
 246 (Freund, 1998; Bayart et al., 2016) when the fracture energy of the barrier is higher than
 247 the value of the energy release rate of the incoming rupture, $G(v)$, for all values of $0 \leq$
 248 $v < v_{arr}$. In particular, at arrest (see Eq. 1) the static energy release rate, $G_s = \Gamma/g(v_{arr})$,
 249 is insufficient to overcome the fracture energy of the barrier, $\Gamma_{barrier}$.

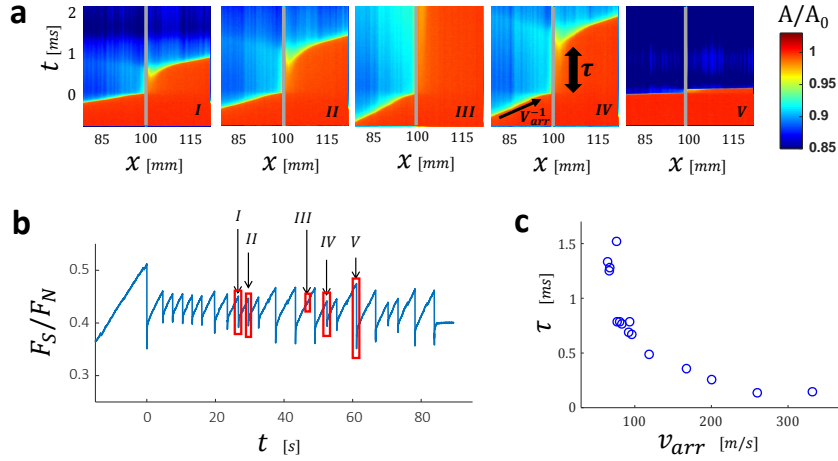


Figure 3. Nucleation Experiments. (a) Rupture arrest occurred when a rupture encountered a high fracture energy ‘barrier’, here imposed at location $x = 100\text{mm}$. Shown are the propagation and arrest of 5 spontaneous ruptures chosen from within a single stick-slip sequence. For events I, II, IV, and V, a nucleation event spontaneously initiated at the far side of the barrier (gray rectangle). Indicated are the velocities of the spontaneous ruptures prior to arrest, v_{arr} , and the respective timescales, τ , of the resulting nucleation process. Note that in event III, no secondary nucleation was observed. (b) The (typical) stick-slip sequence in which the events in (a) occurred. Each stress drop corresponds to a single rupture event. Noted are the events presented in (a). (c) The relation between the nucleation timescales, τ , and the velocities of the arrested crack, v_{arr} , of the full stick-slip sequence. Faster arrested cracks result in shorter nucleation processes.

250 Examples of rupture arrest and consequent nucleation events are presented in Fig.
 251 3a for different values of v_{arr} . Each abrupt rupture arrest produces shear waves of suf-
 252 ficiently high amplitude to create a slight amount of damage as they propagated beyond
 253 the barriers and along the interface. This damage rendered them visible in the contact
 254 area measurements, as they generated $\sim 1\%$ reductions of $A(x, z, t)$ with their passage

255 (Gvirtsman & Fineberg, 2021). When this ‘damage’ is sufficiently large, these shear waves
 256 trigger the nucleation of a new rupture front near the far edge of the barrier. We believe
 257 that this damage results from the detachment of the weakest contacts in the region of
 258 high shear stress, near the barrier’s far edge, created by the stress singularity of the ar-
 259 rested crack. A full discussion of this initial damage is found in (Gvirtsman & Fineberg,
 260 2021). The nucleation onset is defined by the time elapsed after this shear wave passed
 261 x_{nuc} , where $x = x_{nuc}$ is the nucleation location (see Fig. 2). *This onset time is used*
 262 *to determine the duration, τ , of the nucleation phase.*

263 We define the timescale, τ , as the elapsed time between the nucleation onset and
 264 the onset of a *dynamic* rupture triggered by the nucleation process (see Fig. 3a). τ is
 265 indicative of the duration time of the nucleation process and, as shown in Fig. 3c, is a
 266 continuous, monotonically decreasing function of v_{arr} .

267 If nucleation occurs, it will be driven by the induced stress that is produced by the
 268 arrested rupture, $\Delta\sigma_{xy}(v_{arr}, x)$, that is described by Eq. 2. Knowledge of $\Delta\sigma_{xy}(v_{arr}, x)$
 269 enables us to quantify the resulting stress increase at the nucleation site, $\sigma_{ind} \equiv \Delta\sigma_{xy}(v_{arr}, x_{nuc})$.
 270 In this way we are able to study the nucleation process with a known ‘nucleation stress’
 271 and clear ‘nucleation onset time’.

272 In Fig. 4a we present the dependence of τ with σ_{ind} for a typical experiment. The
 273 approximate proportionality for this event, $\tau^{-1} \propto \sigma_{ind}$, is typical (Gvirtsman & Fineberg,
 274 2021). For each sequence of ruptures having the same nucleation point, a well defined
 275 $\sigma_{ind} = \sigma_{thresh}$ threshold exists. Below σ_{thresh} , rupture nucleation will not take place
 276 beyond the barrier (e.g. event III in Fig. 3a). The value of σ_{thresh} together with the pro-
 277 portionality constant relating τ^{-1} to σ_{ind} will change between different sets of experi-
 278 ments. Both depend on the selected location of the barrier within the interface (Gvirtsman
 279 & Fineberg, 2021). Currently, we do not know how to properly express this relation in
 280 dimensionless form.

281 Results

282 Nucleation Evolution

283 When a nucleation event takes place, a nucleation patch will start to slowly expand.
 284 The nucleation patch’s center, x_{nuc} , is located within the damaged region created by the
 285 shear wave launched with each arrest event. As shown in the sequence of contact area
 286 measurements presented in Fig. 4b, nucleation patch expansion is a *2D* phenomenon;
 287 the patch simultaneously expands (at about the same velocity) in both the x and z di-
 288 rections within the xz interface plane. When normalizing $A(x, z, t)$ to study rupture nu-
 289 cleation, contact area normalization was performed using $A(x, z, t_0)$, for t_0 immediately
 290 *after* the initial shear wave passage. We denote contact area measurements that are nor-
 291 malized in this way by $\hat{A}(x, z)$.

292 As in the example presented in Fig. 4b, the edge of the nucleation patch expands
 293 via a slow and nearly constant velocity, v_{nuc} . We characterize the size of the patch, $L(t)$,
 294 by the distance from its edge to x_{nuc} along the x direction (see Fig. 4b - lower panel).
 295 In terms of $\xi(t)$, defined by (Gvirtsman & Fineberg, 2021) as the distance of the edge
 296 of the nucleation patch from the barrier edge, $L(t) + r^* = \xi(t) + \delta$, where δ is the bar-
 297 rier width (see Fig. 1a). Here, we choose to use the more natural quantity, $L(t)$, to de-
 298 scribe nucleation patch extension and dynamics.

299 Nucleation front propagation velocities, v_{nuc} , are extremely slow. These velocities
 300 are typically 2-3 orders of magnitude lower than the velocities of the dynamic ruptures
 301 that they excite. Moreover, v_{nuc} are constant for each nucleation event. Nucleation patches
 302 do not accelerate as they propagate, in strong contrast to typical dynamic ruptures. The
 303 value of v_{nuc} is determined by σ_{ind} . Moreover, Fig 4c demonstrates that beyond loca-

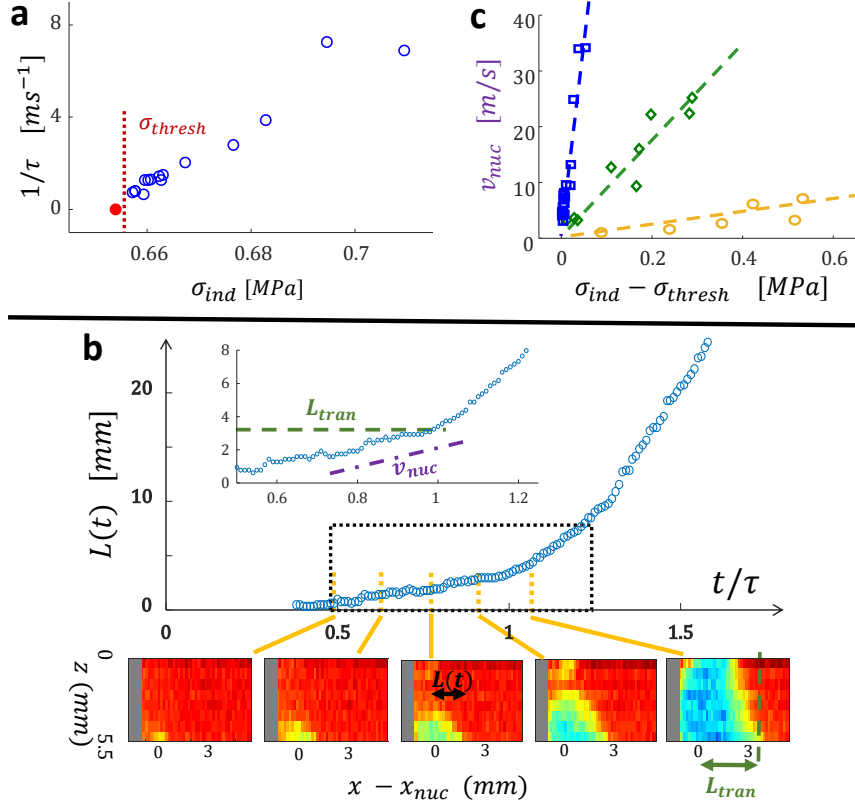


Figure 4. Slow expansion during the nucleation stage. (a) The inverse timescale of nucleation, $1/\tau$, as a function of the induced stress $\sigma_{ind} = \Delta\sigma_{xy}(r^*)$ at the nucleation point, x_{nuc} (as denoted in Fig. 2a). Open points: nucleation events from the same stick-slip sequence. The intercept with the x axis reveals a finite stress threshold, σ_{thresh} , below which nucleation does not occur (Gvirtsman & Fineberg, 2021). σ_{thresh} is determined by linearly extrapolating values of $\sigma_{ind}(v_{nuc})$ to the point where $v_{nuc} = 0$. Red point: an event where no nucleation occurred. (b) Measurements of $\tilde{A}(x, z, t)$ show how nucleation patches slowly expand in both x and z . We use $L(t)$ to characterize the 1D length of the nucleation patch in x . Time is normalized by τ of this event. The patch edge propagates with a slow and constant velocity, v_{nuc} , and the transition at $t = \tau$ is defined by the onset of the patch’s acceleration at length $L_{tran} \equiv L(t = \tau)$. Inset: Enlargement of the region denoted by the dashed square on the main figure. Bottom: 2D snapshots of the nucleation expansion. $L(t)$ is defined as the distance from the edge of the patch to the nucleation point. (c) v_{nuc} as a function of σ_{ind} for 3 sequences of stick slip experiments. In each sequence the barrier was placed at a different location along the interface, producing nucleation in different topographic areas. Every point corresponds to a single nucleation event and each color represents a different sequence. Each sequence is characterized by a different value of the stress threshold, σ_{thresh} . While in all sequences $v_{nuc} \propto \sigma_{ind} - \sigma_{thresh}$, suggesting a non-inertial process, the constant of proportionality as well as σ_{thresh} vary significantly with the nucleation location. Dashed lines are guides to the eye.

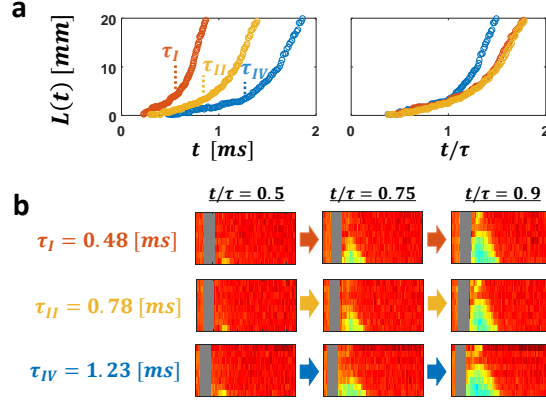


Figure 5. Self similar evolution. (a) The expansion in x , $L(t)$, of 3 different representative nucleation events from the stick-slip sequence presented in Fig. 3. Dashed lines denote the timescale, τ , of each event; indices correspond to labeled events in Fig. 3. When time is scaled by τ , all of the over 20 events in Fig. 3 collapse onto a single curve (here, only 3 are shown explicitly). Note that the collapse is valid only for $t < \tau$, during the nucleation phase. After reaching the transition point, L_{tran} , the dynamics will vary according to LEFM. (b) Snapshots of $\tilde{A}(x, z, t)$ within the nucleation region, showing the 2D self-similarity in the xz plane for the same events. When scaled by their respective values of τ , the geometric expansions of all events are nearly identical.

304 tion dependent thresholds, σ_{thresh} , v_{nuc} linearly depend on σ_{ind} for sets of experiments
305 performed under varying conditions and barrier locations. Such linear behavior suggests
306 a non-inertial process.

307 LEFM predicts that a rupture should be stable below the Griffith length, L_G (Freund,
308 1998). Moreover, beyond L_G , for the stress conditions within nucleation regions, LEFM
309 would predict that a rupture should rapidly accelerate to nearly sonic velocities, as de-
310 scribed by (Freund, 1998). We find that the slow and constant evolution characterized
311 by v_{nuc} continues until $L(t)$ reaches a critical length, L_{tran} (see Fig. 4b). For $L(t) >$
312 L_{tran} nucleation patches accelerate sharply, exhibiting the dynamic behavior expected
313 by LEFM. Beyond L_{tran} rupture fronts therefore enter the dynamic rupture regime (Latour
314 et al., 2013). As demonstrated by (Svetlizky, Kammer, et al., 2017), the rapid accelera-
315 tion that takes place when $t > \tau$ ($L > L_{tran}$), as well as the subsequent rupture motion,
316 beyond possibly a short transition region, is wholly described by fracture mechan-
317 ics.

318 We therefore identify L_{tran} with L_G . The nucleation time, τ , corresponds to the
319 time when $L(t) = L_{tran}$. Moreover, the 3-5mm size of L_{tran} corresponds well (see Dis-
320 cussion section) to the calculated values of L_G for these conditions, as shown in (Gvirtzman
321 & Fineberg, 2021).

322 Self Similarity of Nucleation Patches

323 For each sequence of rupture events, as we saw in Fig. 4a, the nucleation time, τ ,
324 is determined by the shear stress σ_{ind} at the nucleation location. Surprisingly, for each
325 given sequence, both the nucleation patch dynamics and patch shapes are self-similar,
326 when t is scaled by τ . A typical example is presented in Fig. 5, where 3 representative

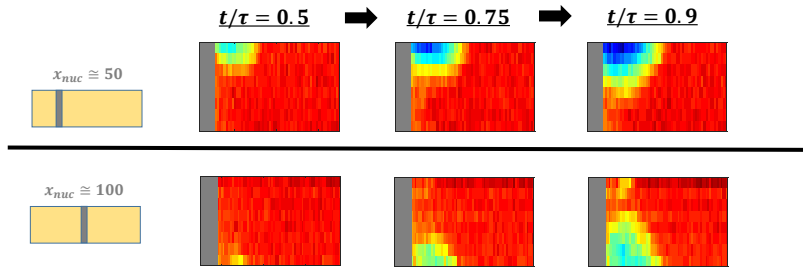


Figure 6. Different nucleation sites provide different 2D evolution. Nucleation patch dynamics at two *different* locations, x_{nuc} , selected within the interface. In both cases, the nucleation dynamics were self-similar as in Fig. 5. Each row is representative of a single sequence of events, where each column shows a snapshot of the 2D nucleation patch at a different stage of the normalized time, t/τ . The location of the barrier in each event is presented schematically at the left.

327 nucleation events with very different timescales, τ , are considered. Compared are the dy-
 328 namics of both $L(t)$ and the entire 2D nucleation patch for these events. As a function
 329 of t/τ , the nucleation dynamics of all 3 events are nearly identical, until the dynamic rup-
 330 ture takes place, at L_{tran} . As values of L_{tran} are quite similar for each nucleation, we
 331 have $v_{nuc} \approx L_{tran}/\tau$.

332 Note that the self-similarity demonstrated in Fig. 5 takes place solely throughout
 333 the nucleation phase, before reaching the critical point at L_{tran} ($t = \tau$). Beyond this
 334 point, rupture propagation takes place according to fracture mechanics. In general, rup-
 335 ture evolution is quite different for different events, as seen when comparing event *c* in
 336 Fig. 5a, for $t > \tau$, with the other events.

337 Although nucleation events are self-similar, the specific *details* of their evolution
 338 depend on the local conditions at the site of nucleation. Fig. 6 demonstrates the vari-
 339 ation of the 2D shapes and expansion rates when comparing different stick-slip sequences
 340 that nucleated at different locations within the interface. For each sequence, we chose
 341 the nucleation location by placing the barrier at different positions along the interface.
 342 Although each sequence was, itself, self-similar, the variation of the local conditions at
 343 each barrier gave rise to different nucleation patch shapes, nucleation velocities, σ_{ind} de-
 344 pendence and nucleation site locations relative to each barrier.

345 The Influence of Nucleation site Topography

346 $A(x, t)$, when *not* normalized, provides us with a measure of the interface ‘topog-
 347 raphy’, as shown in Fig. 1b. These non-normalized intensity values of the 2D surface
 348 measure the absolute amount of light transmitted through the interface that reflects the
 349 total area of the micro-contacts encompassed within each (x, z) pixel of our camera (Rubinstein
 350 et al., 2004). While interface topography of newly established interfaces can vary sig-
 351 nificantly with each rupture event (Morad et al., 2022), mature interfaces can stabilize
 352 their topography (Sagy et al., 2007). Indeed, throughout stick-slip sequences consisting
 353 of numerous rupture events, we find that the topography of the interface does not qual-
 354 itatively change. As Fig. 7 shows, the passage of rupture fronts leaves the qualitative
 355 character of the interface topography intact. The rupture fronts, of course, do cause quan-
 356 titative variations of $A(x, z, t)$, but the overall topological features (peaks, valleys, ridges)
 357 are retained by the interface. The small section described in Fig. 7 is wholly represen-
 358 tative. Even in this single example, the different rupture events that sequentially tra-
 359 verse this section are wholly different in character. These events range from precursory

360 events, where the interface only partially ruptures (Rubinstein et al., 2007) to highly en-
 361 ergetic events (e.g. the 4th panel in Fig. 7b) in which the overall changes in $\langle A \rangle$ of
 362 this patch and the stress release (F_S/F_N) precipitated by a rupture were quite large.

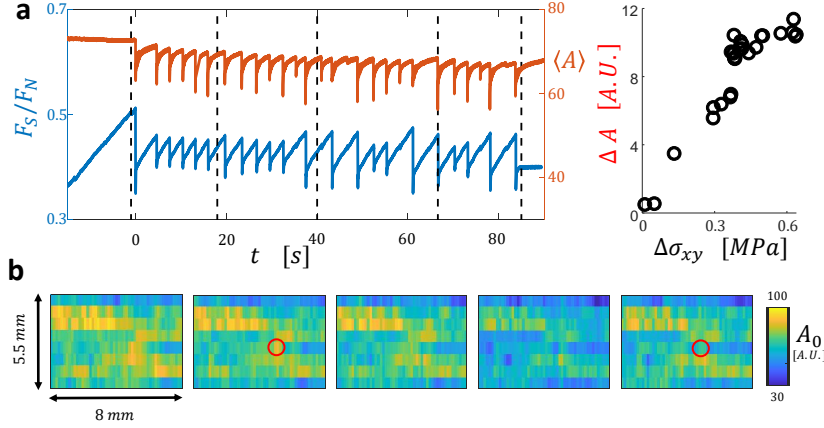


Figure 7. Overall surface topography is largely retained throughout rupture events. The overall surface topography does not change despite numerous rupture fronts that traverse the interface. Shown is the evolution of the topographical map of a nucleation region, whose dynamics are described in the lower panel of Fig. 6. The interface topography remained largely invariant throughout numerous stick-slip sequences. (a) (left) A typical stick-slip sequence consisting of 27 full and partial events. Compared are, $\langle A \rangle$, the mean values of the non-normalized contact area (red) of the typical section of the interface presented in (b) (units are arbitrary, but constant throughout the sequence) and the respective values of the applied stress ratio (blue) F_S/F_N . By both measures, the magnitudes of the different events in the sequence vary considerably. The events include system-wide as well as partial rupture (Rubinstein et al., 2007) events. (right) The corresponding contact area drops, ΔA , as a function of the peak stress drops, $\Delta\sigma_{xy}$, measured by the strain gage closest to the contact area measurements. (b) Snapshots of the non-normalized real area of contact in the section of the interface, whose mean values were quantified in (a). Colors represent the intensity of light that is transferred through each pixel, color coded from 0 to 100. The color codes describe the relative area of the asperities within each pixel that are making contact. Each pixel encompasses an area of $165 \times 688 \mu\text{m}$. As single contacts are approximately $5 \times 5 \mu\text{m}$, each pixel contains over 1000 contacts. Therefore, a value of 50 roughly tells us that 50% of the asperities within a given pixel are in contact relative to the areas of maximal contact (100). These statistics don't significantly depend of the value of the light threshold used. As the light intensity corresponds to the total area of the contacts within each pixel, this picture provides us the 'topography' of the real contact area, $A(x, z, t)$. The different panels correspond to the times noted by the dashed lines in (a). Despite the numerous rupture events that traversed this section, the topography has not qualitatively changed, although the overall quantitative differences are evident in $\langle A \rangle$. Some qualitative changes do occur, however, as noted by the section circled in red, which largely disappeared over the rupture sequence.

363 The invariance of the overall topology over long sequences of ruptures is, perhaps,
 364 surprising to one who would expect a large event to 'reset' the contact area. This ob-
 365 servation is, however, consistent with predictions (Tabor, 1977) as well as recent exper-

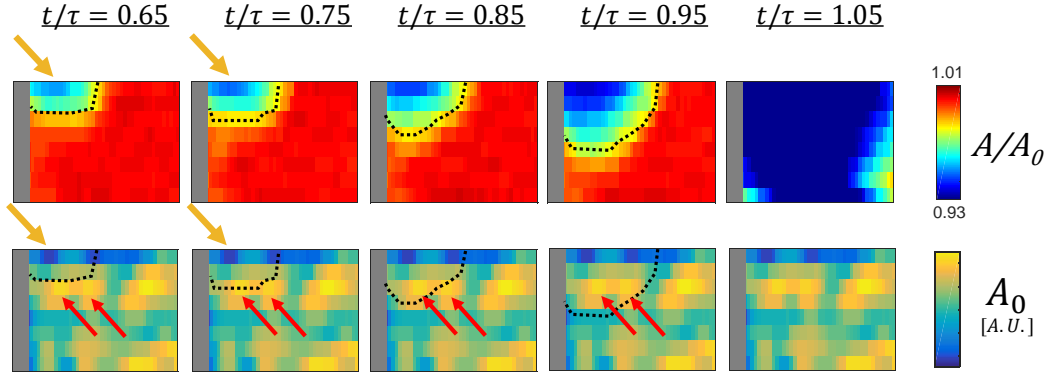


Figure 8. Local Topography guides 2D Evolution. Each column presents a snapshot during the expansion of a nucleation patch. At each stage, the top panel presents the *normalized* picture, $\tilde{A}(x, z)$, where colors corresponds to changes from the initial contact area, while the bottom panel presents the *non-normalized* intensity measurement, $A_0(x, z)$, where colors correspond to the absolute amount of contacts at this point, or the local ‘toughness’. Dashed curves represent the edge of the nucleation patch on both panels. Weak (tough) points are denoted by an orange (red) arrows, respectively.

366 imental (Chen et al., 2020) and numerical (Pham-Ba et al., 2020) observations that sug-
 367 ggest that the contact surface remains invariant when the slope of typical asperities be-
 368 comes sufficiently small. These observations are also consistent with the numerous ob-
 369 servations of ‘repeating’ earthquakes within natural faults whose seismic signatures are
 370 fairly repetitive (Sammis & Rice, 2001).

371 Despite the overall approximate invariance of $A(x, z, t)$, some contact area features
 372 indeed change as more and more rupture fronts traverse them. For example, in Fig. 7b,
 373 the encircled high spot connecting the two high ‘ridges’ in $A(x, z, t)$ was largely erased
 374 by the 27 successive ruptures that passed through the section presented. Such relatively
 375 ‘small’ changes in local barrier topography can precipitate very large effects, such as pos-
 376 sibly observed in the Tohoku 2011 earthquake in which a pre-existing barrier was appar-
 377 ently broached (Scholz, 2014).

378 Let us now consider the effect of the topography of the contact area near a given
 379 barrier. The nucleation site’s influence on dynamics comes to play through the local dis-
 380 tribution of contacts - the local ‘topography’. We can take advantage of our raw mea-
 381 surement of light passing through the interface to obtain a picture of this topography.
 382 A typical example of how topography influences nucleation front dynamics is presented
 383 in Fig. 8. In the figure we compare the topographical map of the nucleation area to the
 384 path selected by a nucleation front at different moments in time, in the region of the bar-
 385 rier imposed at 50mm (the event described in the upper panels of Fig. 6). Here we fol-
 386 low the evolution of a nucleation patch in the normalized and non-normalized represen-
 387 tations of the contact area where, in the latter, we sketch the nucleation patch bound-
 388 aries at each instant. There are two factors that influence the selection of the nucleation
 389 point. (1) The proximity to the barrier is important; the closer to the barrier, the higher
 390 the magnitude of the induced stress field, σ_{ind} , generated by the arrested crack location
 391 (see Fig. 2a). (2) An additional key factor for selection of the nucleation location is the
 392 local topography; the lower the contact area (weaker the location) the easier it will be
 393 to initiate rupture. We see the influence of both of these factors in the example presented

in Fig. 8. The initiation of the patch (denoted by an orange arrow) is, indeed, located at one of the weakest points in the near vicinity of the barrier.

Once nucleation takes place, we find that the local topography will serve to guide the path and geometry of the nucleation front. As Fig. 8 demonstrates, local ‘tough’ points - hotter colors - indeed influence the nucleation evolution. A ridge of tough points (red arrows) seems to both guide the nucleation front and ‘delay’ its 2D expansion. These rather small heterogeneities have a large impact on the dynamics of the very slow nucleation fronts; after overcoming the tough points, patches will often ‘jump’. On the other hand, local toughness variations along the interface will generally not influence the propagation of a fast crack. At the onset of propagation, we expect that the *physical* heterogeneity of the surface itself will have a much larger influence on guiding a crack front than, for example, the heterogeneity of the stress field (Ripperger et al., 2007). The reason for this expectation is that the fracture energy (contact strength) is a local quantity. At each location, a rupture front must ‘compare’ the energy flux into its tip (an integral quantity) with the fracture energy, a local one. Whereas local strengths can fluctuate widely, as the map in Fig. 8 suggests, the energy flux is determined by the mean values of the surrounding fields. The energy flux is, therefore, a smooth function of the position and is not heavily influenced by small-scale fluctuations of the stress field.

We also find that, once the nucleation front expands over the entire span of the interface, its dynamics are significantly affected (Fukuyama et al., 2018; McLaskey, 2019; Gvirtzman & Fineberg, 2021), and it will generally increase its propagation speed.

The buildup of the cohesive zone

As the nucleation patch expands beyond the critical point, L_{tran} , it undergoes a rapid acceleration and propagates rapidly according to fracture mechanics (Svetlizky, Kammer, et al., 2017). Cracks are, in LEFM, considered to be singular objects whose tip is described by putatively singular stress fields, $\sigma_{i,j} \propto K/r^{1/2}$. Nature, however, will not countenance mathematically singular fields and real cracks are more complex objects. In particular, the singular fields at their tips must be regularized or ‘blunted’. The region in which rupture-tip singularities are blunted is generally called a ‘cohesive zone’ (Freund, 1998; K. Bertram Broberg, 1999). This region bridges the gap between the (approximately constant) shear stresses along the interfaces at the tail of a rupture and the approximate singularity at its tip, and essentially describes the region in which the fracture process is taking place at the rupture tip. Only recently (Svetlizky & Fineberg, 2014; Berman et al., 2020) has this rather elusive region been experimentally observed for a dynamic rupture, in cases of ‘fully developed’ frictional ruptures.

The cohesive zone, however, needs to be formed by a crack. In fully developed cracks, the form and scale of this region are dependent on the propagation velocity (Freund, 1998; K. Bertram Broberg, 1999), but for frictional cracks (for the conditions in these experiments) the extent of the cohesive zone of relatively slow cracks has been shown to extend over a few mm (Svetlizky & Fineberg, 2014; Berman et al., 2020) in the vicinity of the rupture tip. At $t = \tau$, the size of the nucleation patch is at this scale. The cohesive zone, however, is not necessarily fully developed.

We now are in a position to study how the cohesive zone development takes place, by monitoring the development of $A(x, t)$ for $t > \tau$. Prior studies have shown (Svetlizky & Fineberg, 2014) that the near-tip shape of $A(x, t)$ is a way to experimentally characterize the cohesive zone, as this is precisely the region in which a propagating rupture transforms the initially intact interface ($\tilde{A} = 1$) to one in which the contact area is both reduced and conceivably weakened. This region is where contact ‘fracture’ is taking place. The reduction of A at a rupture tip is a weak function of the propagation velocity (Svetlizky, Bayart, et al., 2017). For the slowest rupture velocities, (Svetlizky, Bayart, et al., 2017) found that $\tilde{A} \approx 0.9$ behind the rupture tip, yielding $\Delta A \equiv (A - A_{res})/A_0 > 0.1$,

445 where A_{res} is the residual contact area of a fully developed rupture front. Rupture prop-
 446 agation velocity, moreover, influences more than simply the magnitude of the drop in the
 447 contact area. For example, recent studies of frictional ruptures (Berman et al., 2020) in
 448 these materials have revealed that the cohesive zone structure, in fact, also changes dra-
 449 matically with the propagation velocity.

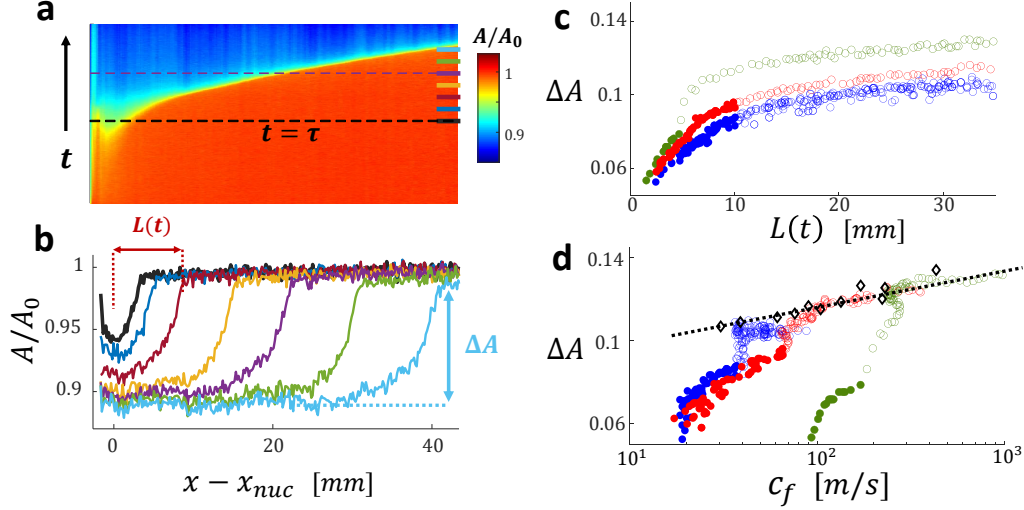


Figure 9. Cohesive zone growth: from nucleation to propagation. (a) Spatio-temporal pic-
 ture of the evolution in x (integrated over z) of $A(x, t)/A_0$ throughout the transition from the
 nucleation stage to dynamic rupture. (b) Profiles of the 1D normalized contact area, $\tilde{A}(x - x_{nuc})$,
 from the data presented in (a) for $t \geq \tau$. The time of each profile is denoted by the bars on the
 right edge of (a); plot colors correspond to respective bar colors. The profile at $t = \tau$ is marked
 in black. Rupture lengths, $L(t)$, are defined with respect to x_{nuc} , and ΔA is defined as the max-
 imum drop in contact area within the tail of the rupture front. (c) ΔA as a function of $L(t)$, for
 3 different events from the same stick-slip sequence, where $t \geq \tau$. Red symbols correspond to the
 event presented in (a,b). (d) ΔA as a function of rupture front velocities, $c_f = \dot{L}$, in a semi-log
 representation. Black symbols denote the contact area drop of all spontaneous *fully developed*
 rupture fronts belonging to this sequence, prior to their arrest at the barrier. The dashed line is a
 guide for the eye. The logarithmic dependence of the values of $\Delta A(c_f)$ was previously described
 in (Svetlizky, Bayart, et al., 2017). Full circles in (c-d) denote the propagation stage prior to the
 final jump to fully developed values of ΔA .

450 In Fig. 9 we present the evolution of ΔA as a function of the distance, $x - x_{nuc}$,
 451 at times beyond $t = \tau$. Fig. 9 demonstrates that at the onset of dynamic rupture $\Delta A \approx$
 452 0.05, and has not yet reached its fully developed value. As the rupture progresses and
 453 accelerates, ΔA increases until, only at $x - x_{nuc} \sim 20\text{mm}$ does ΔA attain the value
 454 compatible with a fully developed rupture front. It is interesting that the growth of ΔA
 455 is not ‘continuous’. For each of the 3 examples described in Figs. 9c,d, ΔA undergoes
 456 a nearly discontinuous ‘jump’ to its steady-state value as seen in Fig. 9d; after τ , ΔA
 457 grows continuously from ≈ 0.05 to $\Delta A \approx 0.075 - 0.09$ before making a rapid ‘jump’
 458 to its fully developed value. The ΔA values for a fully developed ruptures were deter-
 459 mined from the rupture fronts prior to their arrest at the barriers (black symbols in Fig.
 460 9d), which correspond to the measurements of (Svetlizky, Bayart, et al., 2017). In sum-
 461 mary, we find that the full development of the cohesive zone does not take place within

462 the nucleation stage. During the slow evolution of the nucleation patch ΔA grows to a
 463 value of ~ 0.05 . After the transition at $t = \tau$, ΔA continues to develop until reach-
 464 ing its velocity dependent value for a fully developed rupture front.

465 Discussion

466 Here, we have focused on characterizing the nucleation process, which, in contrast
 467 to dynamic rupture, is not a process that is described in our current understanding of
 468 fracture mechanics. To achieve this, we have utilized our knowledge of fracture mechan-
 469 ics to establish well-defined conditions that enable a detailed study of rupture nucleation;
 470 we used ‘artificially’ arrested ruptures to provide quantitative characterizations of both
 471 the location and conditions for which nucleation takes place.

472 This technique for generating nucleation events can, essentially, be considered a demon-
 473 stration of ‘remote triggering’ of earthquakes.

474 As shown in (Gvirtsman & Fineberg, 2021), arrested ruptures generate a wave prop-
 475 agating at approximately the shear wave speed, C_S , that slightly reduces $A(x, z, t)$ along
 476 its path. This slight damage creates a well-defined starting point in time for nucleation.
 477 We surmise that these processes are similar to remote triggering processes that are some-
 478 times observed upon the arrest of large natural earthquakes (Brodsky & van der Elst,
 479 2014). This mechanism may also be involved in the excitation of aftershocks in the close
 480 vicinity of an arrested earthquake.

481 Despite this ‘dynamic’ triggering of nucleation, we believe that the ensuing nucle-
 482 ation dynamics are actually quite general. As shown in Fig. 4b, stresses at the nucle-
 483 ation site must surpass a location-dependent threshold value. In the work described here,
 484 we have ‘artificially’ created these conditions, where the high stress induced by an ar-
 485 rested rupture on one side of the barrier generated a high-stress region beyond the bar-
 486 rier’s other side.

487 By this method, we have demonstrated that the process of nucleation is a well-defined
 488 continuum process whose dynamics are qualitatively similar at each barrier location. In
 489 particular:

- 490 • The nucleation process is the vehicle that will create an initial rupture – bring-
 491 ing the initial nucleation patch to a size (corresponding to the Griffith length) at
 492 which a frictional system loses stability to dynamic fracture.
- 493 • All nucleation patches observed have a distinct 2D character. Nucleation fronts
 494 are well-defined entities that propagate at velocities, v_{nuc} that are 2-3 orders of
 495 magnitude below typical rupture velocities. The values of v_{nuc} are determined by
 496 the shear stress values, σ_{ind} , surrounding the nucleation location. Below a thresh-
 497 old value for σ_{ind} nucleation can not take place.
- 498 • For nucleation to occur, we require a combination of a high stress concentration
 499 and a locally weak region.
- 500 • While nucleation is a continuum process, heterogeneity along an interface (of ei-
 501 ther stresses, local values of the fracture energy, or even prior rupture history) will
 502 be a deciding factor in where, when and in what form nucleation will take place. Thus,
 503 local topography highly influences the details of the progression of nucleation patches
 504 (Fryer, B., Giorgetti, C., Passelègue et al., 2022).
- 505 • Self-similar evolution of nucleation patches will only be evident so long as the im-
 506 portant topographical features at nucleation site remains invariant. We have shown
 507 that, in the ‘mature faults’ described in these experiments, the interface topog-
 508 raphy indeed remains approximately invariant - despite numerous rupture fronts
 509 that traverse a given area.

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Relation to previous work

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The nucleation process described here is, essentially, a quantitative extension of Ohnaka’s description (Ohnaka & Shen, 1999) of rupture nucleation. One difference is, perhaps, that we have clearly shown that there are two distinctly different processes that describe frictional ruptures; a distinct nucleation phase that couples to the phase of rapid rupture that is described by fracture mechanics. These observations complement and expand the observations of Latour (Latour et al., 2013), Gori et al (Gori et al., 2021), and McLaskey (McLaskey, 2019; Cebry & McLaskey, 2021).

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Rupture nucleation, of course, can also be initiated by means other than those used here, for example by fluid injection. Fluid injection both locally decreases the normal pressure, thereby weakening the interface, as well as increases the background shear stress, as in our experiments. The effects of nucleation via fluid injection can be seen in the enhanced seismic activity in locations where hydrofracture is implemented (Schultz et al., 2020). Nucleation by fluid injection has also been studied in recent experiments (Cebry & McLaskey, 2021; Gori et al., 2021) whose qualitative observations were, in many respects, very similar to the nucleation process demonstrated here. Fluid injection created slow nucleation patches that expanded well beyond the region that the injected fluid could reach, suggesting that the stresses created by the injection process dominate the expansion. Slow patch expansion took place until patches reached lengths (we surmise the Griffith length) where rapid dynamic ruptures were triggered. From this point, ruptures accelerated and propagated at dynamic velocities.

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The retention of local topography

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The retention of the local topography in our experiments (e.g. Fig. 7) may appear rather surprising; one could expect that significant sliding along the interface that takes place in multiple successive events should crush or at least largely modify any significant topological features. Such topography retention will certainly not *always* take place. Recent experiments (Morad et al., 2022) have indeed demonstrated that ‘new’ interfaces, created by the controlled fracture of intact rock, evolve dramatically with slip. We might understand these apparent disparities in light of recent work (Pham-Ba et al., 2020) that has demonstrated the validity of the ‘Tabor parameter’ (Tabor, 1977) as a condition for contact rupture. Above a critical roughness, asperities will fracture and surfaces will be ground down. Below this value faults are ‘mature’, plastic deformation of asperities is expected for which interface topography will be fairly well conserved. This situation had been previously noticed in early simulations (Ben-Zion & Rice, 1995) where repeatability of earthquakes was seen to be related to grid-size dependence. Further support for this condition has also been recently observed in rock friction (Chen et al., 2020). The retention of interface topography should be a necessary condition for the numerous instances of repeating earthquakes that are routinely observed within natural faults (Sammis & Rice, 2001; Uchida & Bürgmann, 2019).

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Rupture beyond nucleation

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The strong dependence of the details of the nucleation process on the local topography is, we believe, due to the fact that nucleation is a threshold process; any perturbation of the conditions around the requisite threshold will have a strong influence. Once, however, the transition to dynamic fracture occurs, the influence of topography will lessen considerably.

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In dynamic fracture the ensemble dynamics do not react to individual asperities, but instead relate to a continuous mean (and mesoscopic) quantity that characterizes the fracture surface, the fracture energy. That said, sufficiently large heterogeneity at

558 large scales will influence fracture dynamics – this property was utilized here to cause
 559 the rupture arrests that triggered the subsequent nucleation process.

560 We would like to add a few words regarding interface rupture as described by dy-
 561 namic fracture. When a frictional rupture propagates, it leaves in its wake a residual stress
 562 residual shear stress at each point, $\sigma_{xy}^{res}(x)$. So long as σ_{xy}^{res} is not a strong function of
 563 the slip velocity (Roch et al., 2022; Barras et al., 2019, 2020), the description of rupture
 564 can be perfectly mapped to the fracture mechanical description for shear fracture (Palmer
 565 C. & Rice R., 1973). The stability of an initial ‘interface crack’ is determined by the in-
 566 stantaneous balance of energy; the global amount elastic energy within the *entire* sys-
 567 tem (above the shear stress level determined by σ_{xy}^{res}) is balanced by the local dissipa-
 568 tion at the interface defined by the fracture energy. There is no ‘threshold value’ for the
 569 imposed elastic energy in a system. The greater the imposed shear stress beyond σ_{xy}^{res}
 570 prior to nucleation (or, equivalently, the larger the amount of existent elastic energy prior
 571 to rupture nucleation) the larger a rupture’s acceleration towards either the Rayleigh wave
 572 speed or beyond (Kammer & McLaskey, 2019; Passelègue et al., 2020), once nucleation
 573 takes place. Moreover, all of consequent rupture dynamics are described (Svetlizky, Kam-
 574 mer, et al., 2017; Kammer et al., 2018) by fracture mechanics. This picture has been val-
 575 idated by numerous experiments over the past decade (Ben-David, Cohen, & Fineberg,
 576 2010; Ben-David & Fineberg, 2011; Svetlizky, Kammer, et al., 2017; Passelègue et al.,
 577 2013; Wu & McLaskey, 2019; Xu et al., 2019; Passelègue et al., 2020; Rubino et al., 2017;
 578 Yamashita et al., 2018; Chen et al., 2021).

579 In general, this balance of energy enables dynamic ruptures to possess an energy
 580 ‘buffer’ that they can draw on to overcome local barriers. This energy buffer comes at
 581 the expense of their propagation velocity. Depending on a rupture’s instantaneous prop-
 582 agation speed, the rupture can, to a point, increase the energy flux at its tip by slow-
 583 ing down, if required to overcome a fracture energy barrier. As ruptures approach C_R ,
 584 this dynamic energy buffer becomes huge. As $C_f \rightarrow C_R$, even the large barriers cre-
 585 ated here are easily overcome.

586 This entire description of rupture dynamics is, however, contingent on an impor-
 587 tant condition; *that an initial crack is in existence*. Without an initial flaw (or crack)
 588 the singular focusing of energy to the rupture tip that enables the LEFM description can
 589 not take place. The fracture mechanics framework can not describe how this initial rup-
 590 ture is created. This framework can only describe its stability and ensuing dynamics as-
 591 suming that it exists. For this reason, the nucleation process is critical as, without it,
 592 an interface will always be stable to fracture - no matter what the level of initially im-
 593 posed energy in the system.

594 In a sense, rupture nucleation is analogous to the conditions leading to a forest fire.
 595 Forest conditions must be sufficiently dry - the drier the forest the larger the potential
 596 for a large conflagration. Whether a forest fire will initiate or not at a given time, how-
 597 ever, depends on whether someone ‘lights a match’ (nucleation).

598 **The relation between L_{tran} and the Griffith length**

599 In the section entitled ‘Nucleation Evolution’ (Fig. 4 and 5) we demonstrated that
 600 nucleation patch growth is the vehicle that brings a frictional system a well-defined tran-
 601 sition length, L_{tran} after rapid rupture dynamics initiate. How does L_{tran} compare to
 602 the Griffith length, L_G , the relevant scale for the nucleation of dynamic fracture? Gvirtz-
 603 man et al. (Gvirtzman & Fineberg, 2021) calculated L_G , via energy balance, for the ge-
 604 ometric and stress conditions in these experiments. These calculations yielded values of
 605 3–5mm, in good agreement with our measurements of L_{tran} . The close agreement be-
 606 tween L_{tran} and L_G indeed supports the idea that the nucleation process is indeed the
 607 vehicle that brings the system to LEFM-described dynamics. (The nucleation process
 608 in essence *replaces* the concept of a critical static friction coefficient).

In the literature there are two other, quite different, criteria for the stability of a frictional interface. Below, we demonstrate that each of these criteria predict values for L_{tran} that are much larger than the measured values.

Let us first consider the calculated stability of a frictional interface separating two rigidly sliding bodies that are coupled by the friction law described by Rate and State friction. This calculation (Rice et al., 2001) yielded a critical nucleation length, at which the steady slide becomes unstable to be:

$$L_c^{R/S} = \mu D_c \pi / [(b - a) \cdot \sigma_0] \quad (3)$$

In our experiments, the shear modulus and normal load are, respectively, $\mu = 2GPa$ and $\sigma_0 = 4.5MPa$. The slip distance $D_c \approx 5\mu m$ was obtained from direct measurements of the cohesive zone size (Berman et al., 2020). Values of the coefficients $a \approx 0.008$ and $b \approx 0.015$ for PMMA under our experimental conditions were measured separately (Ben-David, Rubinstein, & Fineberg, 2010; Baumberger & Caroli, 2006). Applying these values to Eq. 3 yields $L_c^{R/S} \approx 1m$, which is 2-3 orders of magnitude larger than the measured value of L_{tran} .

Let us now consider the critical length for instability suggested by Uenishi and Rice (Uenishi & Rice, 2003; de Geus et al., 2019; Schär et al., 2021), which assumes a slip weakening friction law in the vicinity of a near-critical value of shear stress. Here, the estimated critical length was predicted to be:

$$L_c^{UR} = 1.158\mu D_c / [\sigma_{peak} - \sigma_{res}] \quad (4)$$

where σ_{peak} and σ_{res} are, respectively, the peak and residual stress values of a frictional interface. These can be related to the fracture energy, Γ , by means of the slip weakening definition of $\Gamma = \frac{1}{2} \cdot D_c \cdot [\sigma_{peak} - \sigma_{res}]$. Inputting the measured value of $\Gamma = 1J/m^2$ (Svetlizky & Fineberg, 2014; Gvirtzman & Fineberg, 2021) and $D_c \sim 5\mu m$ (as above) into Eq.4 yields a value of $L_c^{UR} \approx 3cm$, which is about an order of magnitude larger than L_{tran} .

What is ahead

What are we still missing? Our fundamental understanding of rupture nucleation processes within frictional interfaces and earthquake dynamics is very much in its infancy. In this work, we have provided a first step, the empirical characterization of this critical process. In particular, we have presented evidence that nucleation is not clearly related to the standard fracture mechanics framework; nucleation is stress driven, occurs below L_G , and is characterized by propagation velocities that are considerably below typical rupture speeds. We currently lack a theoretical understanding and description of this important process. At present, we do not even possess the requisite insight to properly non-dimensionalize expressions such as $v_{nuc} \propto \sigma_{ind}$. We believe that a theoretical description of these processes should also be able to properly relate the mesoscopic scales (e.g. nucleation patch size and shape) to the microscopic topographical features of the interface in the nucleation region. These goals still remain as significant theoretical challenges.

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Data Availability

All data used in this work is publicly available on 4TU Centre for Research Data. Dataset: <https://doi.org/10.4121/21523512.v2>.

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