Stiff directed lines in random media

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We investigate the localization of stiff directed lines with bending energy by a short-range random potential. We apply perturbative arguments, Flory scaling arguments, a variational replica calculation, and functional renormalization to show that a stiff directed line in $1 + d$ dimensions undergoes a localization transition with increasing disorder for $d > 2/3$. We demonstrate that this transition is accessible by numerical transfer matrix calculations in $1 + 1$ dimensions and analyze the properties of the disorder-dominated phase in detail. On the basis of the two-replica problem, we propose a relation between the localization of stiff directed lines in $1 + d$ dimensions and of directed lines under tension in $1 + 3d$ dimensions, which is strongly supported by identical free-energy distributions. This shows that pair interactions in the replicated Hamiltonian determine the nature of directed line localization transitions with consequences for the critical behavior of the Kardar-Parisi-Zhang equation. We support the proposed relation to directed lines via multifractal analysis, revealing an analogous Anderson transition-like scenario and a matching correlation length exponent. Furthermore, we quantify how the persistence length of the stiff directed line is reduced by disorder.

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I. INTRODUCTION

Elastic manifolds in random media, especially the problem of a directed line (DL) or directed polymer in a random potential, are one of the most important model systems in the statistical physics of disordered systems [1]. DLs in random media are related to important nonequilibrium statistical physics problems such as stochastic growth, in particular the Kardar-Parisi-Zhang (KPZ) equation [2]. Burgers turbulence, or the asymmetric simple exclusion model (ASEP) [3]. Furthermore, there are many and important applications of DLs in random media such as kinetic roughening [3], pinning of flux lines in type II superconductors [4,5], domain walls in random magnets, or wetting fronts [1,6].

Directed lines have a preferred direction and no overhangs with respect to this direction. The energy of DLs such as flux lines, domain walls, wetting fronts is proportional to their length; therefore, the elastic properties of directed lines are governed by their line tension, which favors the straight configuration of shortest length. Both thermal fluctuations and a short-range random potential (point disorder) tend to roughen the DL against the line tension. As a result of the competition between thermal fluctuations and disorder, DLs in a random media in $D = 1 + d$ dimensions exhibit a disorder-driven localization transition [7] for dimensions $d > 2$, i.e., above a critical dimension $d_c = 2$. These transitions have been studied numerically for dimensions up to $d = 4$ [8–18]. At low temperatures, the DL is in a disorder-dominated phase and localizes into a path optimizing the random potential energy and the tension energy. Within this disorder-dominated phase, the DL roughens, and there are macroscopic energy fluctuations and a finite pair overlap between replicas [19–21] (introduced below). At high temperatures, the disorder is an irrelevant perturbation, and the DL exhibits essentially thermal fluctuations against the line tension. It has been suggested that the critical temperature for the localization transition of a DL in a random medium and the binding of two DLs by a short-range attractive potential coincide [11,14].

DLs in a random medium map onto the dynamic KPZ equation for nonlinear stochastic surface growth with the restricted free energy of DLs in $1 + d$ dimensions satisfying the KPZ equation of a $d$-dimensional dynamic interface. The localization transition of DLs with increasing disorder corresponds to a roughening transition of the KPZ interface with increasing nonlinearity. In the context of the KPZ equation, it is a long-standing open question (recently discussed, for example, in Ref. [22]) whether there exists an upper critical dimension where the critical behavior at the localization transition is modified. Therefore, the critical behavior of lines in random media can eventually also shed light onto the critical properties of the KPZ equation.

In the present paper, we study the localization transition of stiff directed lines (SDLs). We define SDLs as directed lines with preferred orientation and no overhangs with respect to this direction but with a different elastic energy as compared to DLs: SDLs are governed by bending energy, which penalizes curvature, rather than line tension, which penalizes stretching of the line. This gives rise to configurations which are locally curvature free, i.e., straight but straight segments can assume any orientation even if this increases the total length of the line. We investigate the disorder-induced localization transition of SDLs for a short-range random potential and the scaling properties of conformations in the disordered phase. A typical optimal SDL configuration in the presence of an additional short-range random potential at zero temperature is shown in Fig. 1(a), in comparison to a typical optimal DL configurations in Fig. 1(b).

There are a number of applications for SDLs in random media. SDLs describe semiflexible polymers smaller than their persistence length, such that the assumption of a directed line is not violated. Our results apply to semiflexible polymers such as DNA or cytoskeletal filaments like F-actin in a random environment as it could be realized, for example, by a porous medium [23]. Moreover, SDLs are closely connected to surface growth models for molecular beam epitaxy (MBE) [24].
presence of surface diffusion. MBE can be described by a

domain-wall motion equation [25,26], which is

equivalent to the overdamped equation of motion of a SDL.

As a result, SDLs exhibit the same super-roughness as MBE

interfaces. More generally, the zero-temperature problem of a

SDL in disorder can be formulated as a generic optimization

problem for paths in an array of randomly distributed favorable

"pinning" sites (as indicated by points in Fig. 1), which

minimize their bending energy at the same time as maximizing

the number of visited favorable pinning sites.

There is an important relation between the statistical

physics of DLs and SDLs, which stems from the return prob-

abilities of pairs of lines: the contact probability

\( p_{\text{contact}}(L) \) of two thermally fluctuating DLs in

1 + 3d dimensions, i.e., the probability of two DLs with common starting point to

meet again after length \( L \), decays with the same power

law \( p_{\text{contact}} \sim L^{-3d/2} \) as the contact probability of SDLs in

1 + d dimensions [27,28]. We will provide strong numerical

evidence for \( d = 1 \) that this relation between DLs in 1 + 3d

dimensions and SDLs in 1 + d dimensions holds not only for

purely thermal fluctuations but also in the presence of a

random medium with a short-range disorder potential. This

relation then allows us to address the localization transition of

DLs from a different perspective [28]. Because the proposed

relation between DLs and SDLs is based on return probabilities

of pairs of replicas, our results also suggest that the critical

properties of DLs in a random potential and, thus, the

critical properties of the KPZ equation are determined by

the corresponding two-replica problem.

In particular, the relation between DLs and SDLs implies

that the critical dimension for the localization of SDLs

is \( d_c = 2/3 \) rather than \( d_c = 2 \) for DLs. Therefore, SDLs

exhibit a localization transition already in \( D = 1 + 1 \)
dimensions, which is easily amenable to numerical transfer matrix

studies, whereas it requires numerical studies in \( D = 1 + 3 \)
dimensions to investigate the localization transition of DLs.

Therefore, we can verify several concepts that have been

proposed or found for the localization transition of DLs, e.g.,

the pair overlap as order parameter or the multifractal properties of the

localization transition, by simulations of SDLs in \( D = 1 + 1 \)
dimensions.

The paper is structured as follows. In the next section, we

introduce the model for SDLs and comment on its relation

to DLs. The paper is then divided into two parts, analytical

considerations and numerical findings. First, we apply

scaling analysis to determine the lower critical dimension for the

localization transition as well as estimates for the

roughness in the localized phase. We then present analytical

treatments using variations in replica space and the functional

renormalization group and outline difficulties associated with

both techniques. At last, we introduce a numerical transfer

matrix algorithm for a SDL in a random medium and present

numerical results, which validate the existence of a disorder-

driven phase transition into a localized, roughened phase at low

temperatures. We present numerical results for the roughness,

the disorder-induced persistence length, Derrida-Flyvbjerg

singularities, and the free-energy distribution. Furthermore,

we investigate numerically the pair overlap order parameter

and multifractal properties of the transition.

A short account of some of these results has already

appeared as a Rapid Communication [29].

II. MODEL OF THE STIFF DIRECTED LINE

The configuration of a general directed line, i.e., one

without overhangs or loops and without inextensibility con-

straint, can be parametrized by \( (x, z(x)) (0 \leq x \leq L) \) with a

d-dimensional displacement \( z(x) \) normal to its preferred

orientation. In the following, we call the fixed projected length

\( L \) the length of the line. The contour length of the line is given

by \( L_c \approx L + 1/2 \int_0^L dx [\delta(\zeta(x))]^2 \) (to leading order in \( z \)); it is not

fixed and always larger than the length \( L \). Each configuration

of a SDL is associated with an energy,

\[
\mathcal{H} = \int_0^L dx \left[ \kappa \left( \frac{\delta^2 z(x)}{\delta x^2} \right)^2 + V(x, z(x)) \right],
\]

where the first term is the bending energy (to leading order in \( z \)). The second term is the disorder energy with a Gaussian

distributed quenched random potential \( V(x, z(x)) \) with zero mean

\( \overline{V} = 0 \) and short-range correlations,

\[
\overline{V(x, z)V(x', z')} = g^2 \delta^d(z - z') \delta(x - x'),
\]

where \( \overline{\cdot} \) denotes the quenched disorder average over realizations of \( V \), whereas \( \overline{\cdot} \) denotes thermal averaging.

The SDL model (1) is often used as a weak-bending approximation to the so-called worm-like chain or Kratky-

Porod model [30,31]

\[
\mathcal{H}_{\text{WLC}} = \int_0^L ds \left[ \frac{\kappa}{2} \left( \frac{\delta^2 r(s)}{\delta s^2} \right)^2 + V(r(s)) \right],
\]

which is the basic model for inextensible semiflexible polymers, such as DNA or cytoskeletal filaments like F-actin.

The chain is parametrized in arc length, leading to a local

inextensibility constraint \( \delta r(s) = 1 \), which is hard to account for, both numerically and analytically [23,32]. For thermally

fluctuating semiflexible polymers, the approximation (1) only

applies to a weakly bent semiflexible polymer on length scales

below the so-called persistence length, which is [32,33]

\[
L_p = \frac{2 \kappa}{D - 1} = \frac{2 \kappa}{d - 1} T,
\]

in \( D = 1 + d \) dimensions. We use here and throughout the

following energy units with \( k_B \equiv 1 \). Above the persistence
length, a semiflexible polymer loses orientation correlations and starts to develop overhangs.

Also in a quenched random potential the SDL model describes semiflexible polymers in heterogeneous media, only as long as tangent fluctuations are small such that overhangs can be neglected, which is the case below a disorder-induced persistence length, which we will derive below.

We consider the SDL model also in the thermodynamic limit beyond this persistence length, because we find evidence for a relation to the important problem of DLs in a random medium in lower dimensions. This relation is based on replica pair interactions and shows that pair interactions also determine the nature of the DL localization transition. Moreover, this relation can make the DL transition in high dimensions computationally accessible. We will now outline the idea behind this relation.

III. RELATION TO DIRECTED LINES

The difference between SDLs and DLs [1] is the second derivative in the first bending energy term in Eq. (1) for SDLs, which differs from the tension or stretching energy.

pel = ∫ dx (∂xz)2/2 of DLs with line tension τ. This results in different types of energetically favorable configurations (see Fig. 1): large perpendicular displacements z of SDLs as shown in Fig. 1(a) are not unfavorable as long as their "direction" does not change, i.e., as long as they are locally straight and, therefore, do not cost bending energy. Such configurations increase, however, the length of the line and are suppressed for DLs by the tension or stretching energy.

The statistics of displacements z is characterized by the roughness exponent ξ, which is defined by ⟨z^2(L)⟩ ∼ L^2ξ. The thermal roughness is ξ_{th,τ} = 1/2 for DLs and ξ_{th,κ} = 3/2 for SDLs: Equating the thermal energy T with the respective elastic energies gives T ∼ τζ^2/L for DLs and T ∼ κζ^2/L^3 for SDLs. Here and in the following we use subscripts τ (tension) and κ (bending stiffness) to distinguish between the two systems.

Although typical configurations differ markedly, a SDL subject to a short-ranged (around z = 0) attractive potential V(z) can be mapped onto a DL in high dimensions d' = 3d [27, 28]. This equivalence is based on the probability that a free line of length L starting at [z(0) = 0] "returns" to the origin, i.e., ends at z(L) = 0. This return probability is characterized by a return exponent χ: Prob[z(L) = 0] ∼ L^-χ. The same return exponent characterizes the contact probability p_{contact}(L) ∼ L^-χ, i.e., the probability of two lines with a common starting point to meet again after length L, as follows from considering the relative coordinate. For DLs, which are essentially random walks in d transverse dimensions, the return exponent is χ_d = d/2 [35], whereas it is χ_{3d/2} for a SDL (after integrating over all orientations of the end) [34]; they are related to the roughness exponents by χ = ξ_d [28]. The return exponent χ governs the critical properties of the binding transition to a short-range attractive potential or, equivalently, of the binding transition of two lines interacting by such a potential [27, 28]. This follows, for example, directly from a necklace model treatment [35]. The relation

χ_{τ}(3d) = χ_{κ}(d) implies that the binding transition of two DLs in 3d dimensions maps onto the binding transition of two SDLs in d dimensions.

In the replica formulation of line localization problems such as in (1), the random potential gives rise to a short-range attractive pair interaction (see below). Therefore, pairwise interactions of DLs play a prominent role also for the physics of a single DL in a random potential. Furthermore, the critical temperature T_{c,τ} for a DL in a random potential is believed to be identical to the critical temperature T_{c,κ} for a system with two replicas [11,14]. In Sec. VII D, we will show numerically that also for SDLs in a random medium T_{c,κ} = T_{c,τ} holds. The important role of pairwise interactions suggests that not only the binding transition of two DLs in 3d dimensions maps onto the binding transition of two SDLs in d dimensions but that the same dimensional relation holds for the localization transitions of SDLs and SDLs in a short-range random potential.

One aim of this work is to support this conjecture by providing strong numerical evidence that the d → 3d analogy between SDLs and SDLs in a random potential holds for the entire free-energy distributions for DLs in 1 + 3 and SDLs in 1 + 1 dimensions. Because this analogy is rooted in the binding transition of replica pairs, we can conclude that critical properties of the localization transition are determined by pair interactions in the replicated Hamiltonian, which has been previously suggested in Refs. [36,37].

Moreover, it has been proposed that pair interactions can be used to formulate an order parameter of the disorder-driven localization transition of DLs in terms of the overlap q ≡ lim_{L→∞} ∫_0^L dx dθ[z_1(x) − z_2(x)] [19–21], i.e., the average number of sites per length, that two lines in the same realization of the disorder have in common. Localization by disorder gives rise to a finite value of the pair overlap q. This coincides with the binding energy per length that characterizes binding of two polymers by a pair potential. We will also show that the pair overlap indeed provides a suitable order parameter for the localization transition of SDLs in 1 + 1 dimensions. A schematic summary of the relation of DLs and SDLs together (together with some relevant numeric results) is shown in Fig. 2.

IV. SCALING ANALYSIS

A. Lower critical dimension

We start with a scaling analysis by use of a Flory-type argument. For displacements z ~ the bending energy in Eq. (1) scales as E_b ∼ z^2L^{−3}, which also leads to [z^2] ∼ L^3/L^2, and the thermal roughness exponent ξ_{th,κ} = 3/2. The disorder energy in Eq. (1) scales as E_d ∼ √E_b ~ L^{d/2}. The disorder energy we get E_d ∼ L^{(2−3d)/2}, from which we conclude that the disorder is relevant below a lower critical dimension d_{c,κ} with

d_{c,κ} = 2/3.

Above this critical dimension for d > 2/3 and, thus, in all physically accessible integer dimensions, the SDL should exhibit a transition from a thermal phase for low g to a disorder-dominated phase above a critical value g_c of the
disorder (see Fig. 3). Of course, the same distinction could be made in terms of the temperature with a high-temperature phase for \(T > T_c\) and a disorder-dominated low-temperature phase for \(T < T_c\). In the disorder-dominated phase, the SDL becomes localized or “pinned” into a configuration favored by the random potential and assumes a roughened configuration; see Fig. 1. We note that the lower critical dimension for DLs is \(d_c,\kappa = 2\), which is in accordance with the relation between DLs in 1 + 3d dimensions and SDLs in 1 + d dimensions proposed in the previous section. We also point out that for both types of lines the return exponent \(\omega\) assumes the universal value \(\chi = 1\) at the critical dimension because of \(\chi_r = d_{c,\tau}/2 = 1\) and \(\chi_\kappa = 3d_{c,\kappa}/2 = 1\) \([27,28]\).

With \(d_{c,\tau} = 2/3 < 1\) the localization transition of SDLs can be studied numerically in 1 + 1 (or higher) integer dimensions, whereas for DLs with \(d_{c,\tau} = 2\), the localization transition is only accessible in simulations in 1 + 3 (or higher) integer dimensions. Our numerical study of the line localization transition presented below focuses on SDLs in 1 + 1 dimensions, which are computationally better accessible using transfer matrix techniques as compared to a 1 + 3-dimensional system of DLs. This allows us to verify important concepts for the localization transition, such as the overlap order parameter in Sec. VII E, which have not been accessible for DLs in 1 + 3 dimensions until now.

**B. Roughness**

Balancing the Flory estimates, \(E_b \sim E_d\), gives a roughness estimate \(z \sim L^{\omega}\). When disorder is relevant, this leads to

\[
\xi_{\text{Fl},\kappa} = \frac{7}{4 + d} \quad \text{for} \quad d < d_{c,\kappa} = \frac{2}{3}.
\]

This result is only applicable below the critical dimension \(d < d_{c,\kappa}\), where \(\xi_{\text{Fl},\kappa} > \xi_{\text{th},\kappa}\). Above the critical dimension, the Flory result would give \(\xi_{\text{Fl},\kappa} < \xi_{\text{th},\kappa}\), which contradicts our expectation that the SDL roughens as it adjusts to the random potential. Furthermore, the exponent \(\omega\) related to the sample to sample free-energy fluctuations \(\Delta F^2 \equiv (F - \overline{F})^2\) via \(\Delta F \sim L^{\omega}\) would be negative because of the general scaling relation

\[
\omega = 2\xi_\kappa - 3.
\]

This scaling relation follows from the scaling of the bending energy \(E_b \sim z^2L^{-3} \sim L^{2\xi_{\text{th}}-3}\) and the assumption \(\Delta F \sim E_b\) that bending and free energy have the same scale dependence. Note that we do not subscript \(\omega\) in Eq. (8) as we believe that \(\omega_\tau = \omega_\kappa\); see Sec. VII. An exponent \(\omega < 0\) contradicts the existence of large disorder-induced free-energy fluctuations in the low-temperature phase \([12,38]\), for which there is also strong numerical evidence \([9-11]\). We conclude that this kind of argument is not applicable above the critical dimension. The same problems occur in Flory arguments for the DL for \(d > d_{c,\tau} = 2\), where one finds \(\xi_{\text{Fl},\tau} = 3/(4 + d)\) and \(\omega = 2\xi_\tau - 1\). The Flory approach underestimates the roughness exponent for all dimensions both for DLs and SDLs, i.e., \(\zeta > \xi_{\text{th}}\), as it features only one length scale on which the line can adjust to the disorder. Therefore, the Flory results \(\xi_{\text{Fl}}\) should provide a lower bound for the roughness exponent, i.e., \(\zeta > \xi_{\text{Fl}}\). Moreover, the thermal roughness \(\xi_{\text{th},\kappa} = 1/2\) for DLs and \(\xi_{\text{th},\kappa} = 3/2\) for SDLs should provide another lower bound for the disorder-induced roughness, i.e., \(\zeta > \xi_{\text{th}}\), because of the existence of disorder-induced free-energy fluctuations in the low-temperature phase, i.e., \(\omega > 0\).

Furthermore, we can give an upper bound for the disorder-induced roughness by an argument which relates the line to the zero-dimensional problem of a single particle in a harmonic and a random potential \([39]\). In this argument we divide the line in its middle into two straight and rigid segments separated by the midpoint \((L/2, z)\). The single (i.e., zero-dimensional) coordinate \(z\) describes the restricted shape fluctuations of the two-segment line. The bending energy of these shape fluctuations is \(\mathcal{H}_b(z) = \frac{1}{2}(kL^\delta)z^2\), and the disorder energy of the two straight segments is \(\mathcal{H}_d(z) = V(z)\) with \(V(z) = 0\) and \(V(z) = (2g^2L^\delta)(z - z')\). In Ref. [39], a roughness \(z^2 \sim (g^2L^\delta)^{1/2}(kL^\delta) \sim L^{1/2}\) (plus logarithmic corrections) has been obtained for this zero-dimensional problem with the single degree of freedom \(z\) by an Imry-Ma argument and a more detailed replica calculation, which implies a roughness
for the stiff two-segment line. For the directed line, an anal-
ogous argument gives ζ0,σ = 3/4. In the framework of a
functional renormalization group (FRG) calculation both
roughness exponents can be written as ζ0 = ε/4 in a di-

tensional expansion with the appropriate ε = 4 − d for DLSs
and ε = 8 − d for SDLs (see Sec. VI below). Using the scaling
relations (8), this results in 00η = 1/2 for two-segment lines
both for SDLs and DLSs), which should be considered an
upper bound to the energy fluctuation exponent, because the
adaptation of the line to the potential must not lead to larger
fluctuations as compared to the trivial case of summing up
random numbers [40]. Therefore, ζ0 also should provide an
upper bound for the roughness exponent, i.e., ζ < ζ0. The
upper bound ζ < ε/4 is also found in the FRG calculation [40].

All in all, we have obtained bounds

max(ζ0,ζη) < ζ < ζ0, (10)

which apply both for SDLs and DLSs. For SDLs, this gives a
relatively small window of possible roughness exponents,

max(3/4, 7/4) < ζ < 7/4, (11)

which, for example, limits ζc for SDLs in 1+1 dimensions to
1.5 < ζc < 1.75.

V. VARIATION IN REPLICA SPACE

To go beyond scaling arguments we use the replica tech-
technique [41] following the treatment of directed manifolds
by Mezard and Parisi [42]. For the sake of convenience, we
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PHYSICAL REVIEW E 88, 012103 (2013)

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will comment on how to adapt this to higher dimensions. The
quenched average of the free energy F = −β−1ln Z is treated in
the representation

In Z = lim n→∞ Zn, (12)

calculating averages Zn of an n-times replicated system in the
limit n → 0. For the calculation we introduce an additional
parabolic potential or “mass” term, f = dxμζ′(x), in Eq. (1) as
an infrared regularization and a finite correlation length λ in
the disorder correlator,

V(x,z)V(x′,z′) = g2 f0(ε−x/z−x′)δ(x−x′), (13)

where we use f0(x) = √2πλ2 exp(−x/(2λ2)) to retain
the original δ correlator [compare Eq. (2)] for λ ≈ 0. We
write the replicated and averaged partition function as Zrep =

Πa(∫ dζa exp(−βHrep)) with the following replica Hamilton-

ian in Fourier space:

Hrep = 1/2L n a=1 ∑ k (κ k4 + μ)ζa2

− βg2/2 ∑ a,β=1 L dζ a,β (ζa zβ)2, (14)

As mentioned before, Hrep is related to a pair binding
problem: In the limit λ ≈ 0 the second term becomes

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which is an attractive short-

range interaction of two replicated lines.

We use variation in replica space with the quadratic, i.e.,
Gaussian, trial Hamiltonian

H V = (2L)−1 ∑ k a,β=1 L dζ a,β G−1 a,β ζa zβ, (15)

with G−1 a,β = (κ k4 + μ)ζa,β, with the self-energy matrix Σ
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dimensions, we interpret this as an indication for the existence of a critical disorder strength or a critical temperature. The replica approach leads, however, to the thermal roughness exponent also in the low-temperature phase.

VI. FUNCTIONAL RENORMALIZATION GROUP

There has been some success studying elastic manifold problems in disordered potentials using FRG analysis [40,45–49]. This method can be adapted to generalized elastic energies,

$$\mathcal{H}_m \sim \int d^Dx \left( \nabla_x^m \mathbf{z} \right)^2,$$  \hspace{1cm} (23)

for \(D\)-dimensional elastic manifolds with \(d\) transverse dimensions (in the FRG literature, the number of transverse dimensions is frequently denoted by \(N\)). The case \(m = 1\) corresponds to elastic manifolds as they have been already extensively studied [40,45–49], whereas \(m = 2\) corresponds to manifolds dominated by bending energy. Lines are manifolds with \(D = 1\), i.e., DLs correspond to \(m = 1\) and \(D = 1\) and SDLs with a bending energy to \(m = 2\) and \(D = 1\). In the FRG approach we take the Gaussian distributed random potential to have zero mean and a correlator of the general form

$$\langle \mathbf{V}(\mathbf{x}, \mathbf{z}) \mathbf{V}(\mathbf{x}', \mathbf{z}') \rangle = R(\mathbf{z} - \mathbf{z}')\delta^{(D)}(\mathbf{x} - \mathbf{x}'),$$  \hspace{1cm} (24)

where the whole function \(R(\mathbf{z})\) is renormalized under a change of scale.

In renormalization, we integrate out short-wavelength fluctuations in a shell \(\Lambda/b < |k| < \Lambda\) in momentum space and perform a subsequent infinitesimal scale change (SC) by a factor \(b = e^{dl}\),

$$x \rightarrow bx, \hspace{1cm} (25)$$

$$z \rightarrow b^\kappa z,$$  \hspace{1cm} (26)

in order to restore the high-momentum cutoff \(\Lambda\). This leads to the following FRG flow equations:

$$\frac{dT}{dl}_{\text{SC}} = 2(\zeta_0 - \zeta)T,$$ \hspace{1cm} (27)

$$\frac{dR}{dl}_{\text{SC}} = (\epsilon - 4\zeta)R + \zeta(\mathbf{z} \cdot \nabla \mathbf{z})R + O(R^2) + O(R^3),$$ \hspace{1cm} (28)

with

$$\epsilon = 4m - D$$ \hspace{1cm} (29)

and \(\zeta_0 = (2m - D)/2\).\(^1\) The flow equation (27) for the temperature is believed to be exact due to a Galileian invariance of the Hamiltonian. For a disorder-dominated phase with \(\zeta > \zeta_0\) corresponding to \(\omega > 0\), the system is characterized by a \(T = 0\) RG fixed point, at which we want to determine the roughness exponent \(\zeta\).

\(^1\)For \(D > 2k\) the manifold does not have a macroscopic roughness and \(\zeta_0\) can no longer be interpreted as the thermal roughness exponent.

The terms \(O(R^2)\) and \(O(R^3)\) in the RG flow equation (28) of the disorder correlation function \(R(z)\) are additional one-loop [46] and two-loop [40,47–49] contributions, respectively. In Ref. [40], a generalized elasticity with a general parameter \(m\) has already been discussed up to the two-loop level for \(d = 1\). The one-loop contributions are independent of \(m\) and assume exactly the same form for \(m = 2\) as for standard elastic manifolds with \(m = 1\) and as they have been calculated in Refs. [40,45,48] for \(d = 1\) and Refs. [46,49] for general \(d\). For \(d = 1\), it has been shown that the two-loop contributions, however, acquire a \(m\)-dependent numerical prefactor [40,48]. For general \(d\) and \(m\), the two-loop contribution has not been calculated. The exponent \(\zeta\) is determined from the FRG equation (28) for \(R(z)\) by requiring a fixed-point solution for short-range disorder to be positive and vanish exponentially for large \(z\). Therefore, in one-loop order results for the roughness exponent \(\zeta\) depend on \(m\) only through the dimensional expansion parameter \(\epsilon = 4m - D\).

For \(d = 1\), we can adapt the final results achieved in Ref. [45] in one loop, which have been extended in Refs. [40, 48] to two loops,

$$\zeta_{\text{FRG}} = 0.20830\epsilon + 0.000686X_m \epsilon^2,$$ \hspace{1cm} (30)

with the \(m\)-dependent numerical prefactor \(X_m = [X_1 = 1, X_2 = -1/6\text{ leading order in }\epsilon\] for a SDL \((D = 1, m = 2, \epsilon = 7)\) with \(d = 1\) transverse dimensions, we obtain a roughness exponent \(\zeta_{\text{FRG},\kappa} \approx 1.4571\) in one loop, which is close to the Flory estimate \(\zeta_{\text{FL,\kappa}} = 7/2\) but also violates the lower bound set by the thermal roughness, \(\zeta_{\text{FRG,\kappa}} < \zeta_{\text{th,\kappa}} = 3/2\), implying \(\omega < 0\). On the two-loop level, we find a negative prefactor \(X_2 < 0\) and, thus, still \(\zeta_{\text{FRG,\kappa}} < \zeta_{\text{th,\kappa}} = 3/2\).

In the literature, the existence of an upper critical dimension \(d_u\), above which \(\zeta < \zeta_0\) applies, has been discussed, in particular for DLs \((D = 1, m = 1)\) and as a candidate for the upper critical dimension of the KPZ equation to which the DL problem can be mapped. Our above finding, \(\zeta_{\text{FRG,\kappa}} < \zeta_{\text{th,\kappa}} = 3/2\), in two-loop order, indicates that for SDLs with \(m = 2, d = 1\) is already above the upper critical dimension \(d_{u,\kappa}\), i.e., \(d_{u,\kappa} < 1\). Using the one-loop result for general \(d\), we can estimate this upper critical dimension \(d_{u,\kappa}\) for SDLs. The approximate formula

$$\zeta_{\text{FRG}}(d) = \frac{\epsilon}{4 + d} \left[ 1 + \frac{1}{4\epsilon^2} \frac{2}{d+1} (d+2)^2 \right]$$ \hspace{1cm} (31)

from Ref. [46] gives a critical dimension \(d_{u,\kappa} = 0.937669 < 1\). Solving the FRG fixed-point equation for \(R(z)\) numerically, we find \(d_{u,\kappa} \approx 0.84 < 1\) using the one-loop equations and the numerical methods outlined in Ref. [40] (we use Taylor expansions up to order 12, which allows us to determine \(\zeta\) to four digits). The result, \(d_{u,\kappa} < 1\), remains valid using the two-loop equations, where the additional two-loop terms contain the same factor \(X_m\) as for \(d = 1\). However, the two-loop result for the critical dimension is lower because, analogously to the DL results of Ref. [49], the two-loop corrections are negative for \(d > d_{u,\kappa}\). In the following section, we present numerical results which show that there exists a disorder-dominated phase with \(\zeta_c > \zeta_{\text{th,\kappa}} = 3/2\) for SDLs in \(d = 1\) below a critical temperature, which shows that
the FRG results are questionable for lines with $D = 1$ and, correspondingly, large values for the expansion parameter $\epsilon$.

**VII. NUMERICAL RESULTS**

Returning from $(D + d)$-dimensional manifolds to the problem of $(1 + d)$-dimensional SDLs in disorder, further progress is possible by numerical studies using the transfer matrix method [1,50] for $T = 0$ (see Fig. 1) and for $T > 0$.

Previous numerical studies for DLs in $1 + 3$ dimensions offer the opportunity for a comparison of the exponents $\omega$ (energy fluctuations), describing the low-temperature phase, and $v$ (correlation length), describing the transition, in the low-temperature phase, to test the aforementioned analogy to SDLs in disorder, further evaluated. We are using the transfer matrix approach and divide the line into straight segments which are connected by the transfer matrix. This allows for an iterative computation of the restricted partition function

$$Z_L(z', v) = \sum_{v, z} \exp(-\beta \Delta E_L(v - v', z')) Z_{L-1}(z, v);$$

for numerical stability reasons, the $Z_L(z, v)$ are normalized in each length iteration such that $\sum Z_L(z, v) = 1$. The normalization constant is a useful quantity as it is the total partition function and, therefore, gives the free energy. Additionally, the normalized restricted partition function is used in the computation of thermal averages,

$$\langle X \rangle = \left( \sum_{z, v} Z_L(z, v) \right)^{-1} \sum_{z, v} X(z, v) Z_L(z, v).$$

This averaging procedure is correct only for quantities $X$ that are measured at the end of the line; moments of the energy (potential or total) are accumulated along the contour of the line and have, therefore, to be computed in an iteration scheme [50] very similar to Eq. (35). Finally, for all observables, the quenched average over realizations of the disorder has to be performed.

**B. Existence and nature of the disorder-dominated phase**

1. Roughness

The most natural observable to analyze looking for a localization transition is the roughness $\langle z^2 \rangle$. One expects to see two different regimes, a high-temperature phase with the thermal roughness $\langle z^2 \rangle \sim T L^3$ and a low-temperature phase with $\langle z^2 \rangle \sim T^0 L^2$. Also, the influence of numerical details onto the roughness should be smaller than their influence on the free energy. Figure 4 shows the roughness $\langle z^2 \rangle$ as a function of temperature and demonstrates that these expectations are met. In order to determine the roughness exponent $\zeta_\chi$ we measure

\[ \langle X \rangle = \left( \sum_{z, v} Z_L(z, v) \right)^{-1} \sum_{z, v} X(z, v) Z_L(z, v). \]
a “local” roughness exponent \([18]\),
\[
2\xi(L) = \log_{10}(\zeta^2(L)/\zeta^2(L/5)).
\]
(37)
The data for \(\zeta\) as a function of temperature presented in Fig. 5 exhibits two distinct high- and low-temperature regimes and a significant “dip” of the local roughness exponent around \(T \approx 1.4\). This method of determining \(\zeta\) gives better results than fitting \((\zeta^2(L))\).

Via the scaling relation (8), \(\omega = 2\xi - 3\), we obtain the exponent \(\omega\) from the roughness exponent \(\xi\). As in the context of DLs \([54]\), it can be argued that \(\omega\) should vanish at the disorder-induced localization transition, resulting in a roughness exponent \(\xi = 3/2\). This seems to hold, even though the numerical value for high temperatures is slightly above \(\xi = 3/2\). This is strong evidence for a phase transition at \(T_c \approx 1.4\). For low temperatures, the values \(2\xi \approx 3.11\) give \(\omega \approx 0.11\) according to Eq. (8), which is slightly lower than \(\omega = 0.186\), which is the literature value for DLs in 1 + 3 dimensions \([10,11,51,52]\).

2. Disorder-induced persistence length

The roughness is closely related to the averaged tangent directions \(\langle \Delta \zeta \rangle \equiv \langle (\Delta \zeta)^2 \rangle\), which should scale as
\[
\langle \Delta \zeta^2 \rangle / L^2 \sim L^{2(\xi-1)} \sim L^{1+\omega}. \tag{38}
\]
We define an effective disorder-induced persistence length \(\tilde{L}_p\) for the SDL as the length scale at which the tangent fluctuations become equal to 1,
\[
\langle \Delta \zeta^2 \rangle (\tilde{L}_p) = 1. \tag{39}
\]
This generalized definition for the disordered system is consistent with the standard definition for the persistence length \(L_p\) of a thermally fluctuating SDL in the absence of disorder, where we expect \(\langle \Delta \zeta^2 \rangle \approx L / L_p\) with \(L_p = \beta \kappa\). Apart from numerical prefactors this gives the standard persistence length of the WLC model, see Eq. (4), which is defined as the decay length of tangent correlations.

In the low-temperature phase, we expect a disorder-dominated roughness and, therefore, temperature-independent tangent fluctuations \(\langle \Delta \zeta^2 \rangle\), which results in a temperature-independent disorder-induced persistence length \(\tilde{L}_p \sim \beta \kappa\).

at low temperatures according to the criterion (39). For SDLs in 1 + 1 dimension, we expect \(\tilde{L}_p \propto g^{-1}\).

To determine the generalized persistence length from Eq. (39) numerically, we used the tilt symmetry of the replicated Hamiltonian \([55]\), according to which the “connected” average
\[
\left\langle (\Delta \zeta)^2 \right\rangle - \left\langle \zeta \right\rangle^2 \approx L / \beta \kappa \tag{41}
\]
is independent of the disorder strength. Therefore, we can use a fit to the sample-to-sample tangent fluctuations of the form
\[
\left\langle (\Delta \zeta)^2 \right\rangle (T, L) = a(T) L^{1+\omega(T)}, \tag{42}
\]
with an amplitude \(a(T)\) and an exponent \(\omega(T)\), which should agree with the free-energy exponent \(\omega\). We then can rewrite \(\langle \Delta \zeta^2 \rangle\) as
\[
\langle \Delta \zeta^2 \rangle = \left(\langle \zeta^2 \rangle - \langle \zeta \rangle^2\right) + \langle \zeta \rangle^2 = L / \beta \kappa + a(T) L^{1+\omega(T)} \tag{43}
\]
determine \(\tilde{L}_p\) using Eq. (39). Our results for \(\tilde{L}_p\) are shown in Fig. 6. In the low-temperature phase for \(T < T_c \approx 1.4\), the persistence length becomes indeed disorder induced, i.e., almost independent of temperature and our results are consistent with the above scaling result (40). We consider this nondivergent persistence length at low temperatures also to be consistent with previous results for the nondirected version of the SDL, the WLC in disorder \([23]\). In the high-temperature phase, our results approach the standard thermal persistence length \(L_p \sim \beta \kappa\).

The fit results for \(a(T)\) and \(\omega(T)\) are shown in Fig. 7. For \(T < T_c\), our results are consistent with \(\omega \approx 0.186\), which is the literature value for DLs in 1 + 3 dimensions \([10,11,51,52]\). In fact, we get very similar results for \(L_p\) if we fix \(\omega(T) = \omega = 0.186\).
Derrida and Flyvbjerg, the distributions rather than details of the Hamiltonian involved. According to statistics and the distribution used for the random potential specific point $z$ partition $w$, to the literature value for DLs in 1

3. Derrida-Flyvbjerg singularities

One of the features expected in a disorder-dominated phase are Derrida-Flyvbjerg singularities [15,56]. These are features of the statistical weights, here the normalized restricted partition $w(z) = Z(z)/Z$ function for the SDL to end at a specific point $z$. As the phase is disorder dominated, some characteristics of these weights should originate from mere statistics and the distribution used for the random potential rather than details of the Hamiltonian involved. According to Derrida and Flyvbjerg, the distributions $P_1(w_1)$ and $P_2(w_2)$ of the largest and the second largest weight, respectively, should exhibit singularities at $1/n$ ($n = 1, 2, 3, \ldots$) in a disorder-dominated phase with a multivalley structure of phase space [56]. For SDLs in disorder, we calculated the distribution of the value of the largest and the second-largest weight numerically as shown in Fig. 8 and indeed find singularities at 1 and 1/2 for $T < T_c$. We are not able to clearly resolve higher singularities at $1/n$ with $n \geq 3$, which might be due to the underlying (Gaussian) distribution or the number of samples used. Analogous singularities can be found in the distribution of the information entropy $s = -\sum_z w(z) \ln w(z)$ at values $-\ln(1), -\ln(2), -\ln(3)$, whereas nothing similar can be observed at high temperatures, where the entropy distribution is Gaussian and the distribution of the (second) largest weight is sharply peaked around zero. We see this as a confirmation that the SDL indeed undergoes a transition to a disorder-dominated phase at $T_c \approx 1.4$.

C. The free-energy distributions of SDL in 1 + 1 and DL in 1 + 3 dimensions are identical

The exponent $\omega$ can alternatively be determined in a more direct way by fitting $\Delta F = (F - T)^{1/\omega} \propto L^{\omega}$ at temperatures $T \ll T_c$, giving values of $\omega \approx 0.15-0.16$ (cf. Fig. 9).

We can go one step further and consider not only the second moment but the whole distribution of the free energy as shown in Fig. 10, which is obtained by computing the free energy for every sample and rescaling to zero mean and unit variance,

$$G_F(X) = \text{Prob}(F - \bar{F})/\Delta F = X.$$ (44)

This rescaling should make $G_F$ more robust against the influence of numerical details. For DLs in $d = 1$ it has been found that this distribution is of a universal form [57]. The asymptotic behavior of the negative tail of the rescaled free-energy distribution for low temperatures is of the form

$$\ln G_F(X) \sim -|X|^\eta \quad (X < 0, |X| \geq 1).$$ (45)

This allows us to determine the energy fluctuation exponent $\omega$ via the Zhang argument [1,11,13]: A saddle-point integration gives $\ln Z \sim -nF/T - (n\Delta F/T)^{\eta/(\eta-1)}$; on the other hand,
In $\mathbb{Z}^d \sim L$ should be extensive, resulting in $\Delta F \sim L^{1-1/\eta}$ or
\[ \eta = (1 - \omega)^{-1}. \]  
(46)
We find $\eta \approx 1.23$ (dashed black line in Fig. 10) or $\omega \approx 0.18$. This is in agreement with the values reported for DLs in 1 + 3 dimensions.

For a direct comparison of the rescaled free-energy distributions of a SDL in 1 + 1 and a DL in 1 + 3 dimensions we simulated both systems (the DL up to lengths $L = 60$) and find that the rescaled free-energy distributions in the low-temperature phases have to be considered identical within numerical accuracy. This could hint towards a new seemingly “universal” distribution for certain random systems, much like the Tracy-Widom distribution that is found for the DL in 1 + 1 dimensions and various other systems [57]. For finite system sizes this universal behavior can only be expected for free energy fluctuations $|X|$ small compared to an upper threshold [58,59]; we believe, however, that our simulation does not cover the very rare fluctuations that induce the nonuniversal part for very large $|X|$. In Fig. 11 we show the distributions for $T < T_c$ (DL and SDL) in a manner where the exponent $\eta$ becomes more apparent. Here we introduce the exponent $\eta'$, which is the analog to $\eta$ for the positive tail,
\[ \ln G_F(X) \sim -|X|^{\eta'} \quad (X > 0, |X| > 1), \]  
(47)
where the Zhang argument is not applicable. We find a value $\eta' \approx 1.84$.

Based on an exact renormalization on the diamond lattice [17] and an optimal fluctuation approach [59], it has been previously suggested for DLs that $\eta'$ and $\omega$ are related via [cf. Eq. (46)],
\[ \eta' = d_{\text{eff}}/(1 - \omega), \]  
(48)
with $d_{\text{eff}} = 1 + d$ for the hypercubic lattice. This is found to be valid for the DL in 1 + 1 dimensions, where $\omega = 1/3$ and $\eta' = 3$ [17], and references cited therein]. For the problem at hand, the literature value $\omega = 0.186$ for DLs in $d = 3$ would lead to $\eta' \approx 5$, which is far from the value $\eta' \approx 1.84$ we find; thus, the ratio $\eta'/\eta$ does not match the prediction $\eta'/\eta = (1 + d)$. In Ref. [59], $\eta = 2$ and $\eta' = 3$ were found to be superuniversal (independent of $d$) for the DL in dimensions $d > 2 = d_c$ at temperatures $T > T_c$. We can neither confirm nor deny this result, as we are not able to cover the “most distant” part of the distribution, but our interpretation that $G_F$ is Gaussian for $T > T_c$ would lead to $\eta = \eta' = 2$.

The finite-size corrections to the free energy [1,60] should also scale as $L^{\alpha}$ leading to
\[ F/L \approx a + bL^{\omega - 1}. \]  
(49)
We did not succeed in determining a precise, consistent $\omega$ in this way, possibly because our systems are too small.

The free-energy distributions in Fig. 10 seem to be identical for $T < T_c$ and $T \approx T_c$ (as it is also the case for the DL [11]), which suggests $\omega_{T=T_c} \approx 0.186$ contradicting the $\omega_{T=T_c} = 0$ argument [54], but one has to bear in mind that the saddle-point integration in the course of the Zhang argument is only applicable if $\eta > 1$ or $\omega > 0$.

A more distinct difference between the behavior at criticality and at low temperatures can be found in the distribution of the potential energy as shown in Fig. 12 [also the potential energy is rescaled using $X = (E - \bar{E})/\Delta E$ analogously to Fig. 10 for the free-energy distribution]. For the potential energy distribution, the behavior at the critical temperature clearly differs from the behavior at both $T > T_c$ and $T < T_c$ (which are not identical for the DL) and exhibits a decay,
\[ \ln G_E(X) \sim \exp(-|X|) \quad (X > 0, |X| \gg 1), \]  
(50)
resembling extreme value distributions of the Gumbel type.

A tentative explanation might be that the transition occurs because of extreme values of the potential at which the otherwise thermally fluctuating line localizes. The random potential has a Gaussian distribution; thus, its extreme values are distributed according to a Gumbel distribution or Fisher-Tippett type I extreme value distribution [61],
\[ P_{\text{Gumbel}}^{\alpha,\beta}(X) = \frac{1}{\beta} e^{-z(X)/\beta}, \]  
(51)
with $z(X) = (X - \alpha)/\beta$, the location parameter $\alpha$, and the scale parameter $\beta$ (the parameters depend on the number of

FIG. 11. (Color online) Double logarithmic plot of the negative logarithm of the rescaled free-energy distribution $G_F(X) = (F - \bar{F})/\Delta F$ at low temperatures $T < T_c$ for the DL (light blue stars for $X < 0$ and yellow squares for $X > 0$) and the SDL (red plus signs for $X < 0$ and dark blue Xs for $X > 0$). We see identical behavior for SDL and DL consistent with exponents $\eta \approx 1.23$ and $\eta' \approx 1.84$; see text.

FIG. 12. (Color online) Rescaled potential energy distribution for a SDL in 1 + 1 dimensions. Plotted are distributions for three finite temperatures, $T < T_c$ (black thin solid line), $T \approx T_c$ (red thick solid line), and $T > T_c$ (light green solid line). Results for a DL in 1 + 3 dimensions are shown in blue (medium solid line). Here the brown (dark dashed) curve is an approximation using a one-parameter Gumbel distribution [cf. Eq. (52a) with $m = 1.7$], whereas the yellow (light dashed) curve is the normal distribution.
\( \text{D. The localization transition temperature } T_c \text{ equals the replica pair transition temperature } T_2 \text{ for SDLs} \)

We have checked for the SDL via numerical transfer matrix calculations that the localization temperature in disorder \( T_{c,k} \) equals transition temperature \( T_{2,k} \) for replica pair binding, \( T_{c,k} = T_{2,k} \). As stated before, we use the same transfer matrix algorithm for the replica pair system with a short-range binding potential by exploiting that the binding of two SDLs can be rewritten as a binding problem of one effective SDL in an external potential using relative coordinates \([63]\). This SDL has a binding stiffness of \( \kappa' = \kappa/2 \) and the “potential energy” we are interested in \([\text{cf. Eq. (14)}]\) for the replica pair binding is \( -\beta g^2 \int dx \delta (z) \). For the interpretation of the simulation results, one has to keep in mind that the energy functional is temperature dependent and, therefore, derivatives of the free energy with respect to the inverse temperature \( \beta \) are not given directly by cumulants of the internal energy. Using \( E_b = \int dx (\partial_x z)^2 \) and \( V = -g^2 \int dx \delta (z) \) the partition function is given by \( Z = f \int Dz \exp (-\beta E_b - \beta^2 V) \) and the free energy by \( F = -\beta^{-1} \ln Z \), implying that

\[
\frac{\partial (\beta F)}{\partial \beta} = \langle E_b + 2\beta V \rangle = U + \beta \langle V \rangle, \tag{53}\]

where \( U \) is the total internal energy (treating \( \beta V \) as a potential). Thus, the derivative of the difference of the free energies with and without the adsorption term with respect to the inverse temperature, which should give the divergent correlation length \( \beta \delta F = F_{V} - F_0 \sim L(T_c - T)^{\nu/2} \), is identical to the “potential energy” \( \delta E_{pot} = -\beta V \) and usual finite-size scaling should be applicable (cf. Fig. 14). We retain the known correlation length exponent \( \nu = 2 \) for the adsorption problem \([63]\). We see matching curves for the used system sizes \( L = 100, 200, \ldots, 500 \) around \( T - T_c \approx 0 \) for \( T_{2,k} = 1.44 \), which equals \( T_{c,k} \approx 1.4 \) for the SDL in disorder.

\( \text{E. Pair overlap as order parameter} \)

Finally, we identify an order parameter of the localization transition. For DLs, the disorder-averaged overlap \( q = \lim_{L \to \infty} \int_0^L dx \int_0^L dx' \langle z_1(x) - z_2(x) \rangle \) of two replicas has been

FIG. 13. (Color online) The reduced free energy \( \delta F = F - F_{\text{ann}} \) rescaled by \( \ln^{1/2} L \) for lengths \( L = 50, 60, \ldots, 100 \) as a function of the temperature \( T \). There is a pseudocrossing of the lines around \( T \approx 1.38 \).

FIG. 14. (Color online) Finite-size scaling for the SDL adsorption problem that corresponds to the two-replica binding; see text for a detailed explanation. The scaling uses \( T_c = 1.44 \) and \( \nu = 2 \). As only one sample is needed for the calculation we used larger system lengths \( L = 100 \) (red plus signs), 200 (green Xs), and 500 (light blue solid squares).
proposed as order parameter [19,20]. Until now, it has been numerically impossible to verify this order parameter for DLs in \( d > 2 \) dimensions where a localization transition exists because the relevant 2d-dimensional two replica phase space is too large. For SDLs, on the other hand, the transition is numerically accessible already in 1 + 1 dimensions and we show that the overlap \( q \) is indeed a valid order parameter using an adaptation of the transfer matrix technique from Ref. [19]; see Fig. 15. This involves simulating two interacting SDLs; therefore, we can only use lengths up to \( L = 30 \) and \( 10^3 \) samples. For DLs, it has been found that the overlap at criticality decays as \( q \sim |T - T_c|^{-\beta} \) with an exponent \( \beta' \approx 1.36 \). Nevertheless, the connection between DLs and SDLs provides the first system to test the proposed order parameter in a localization transition numerically and to determine the otherwise inaccessible exponents \( \beta' \) or \( \Sigma \).

### F. Multifractal properties at the transition

For DLs in \( 1 + 3 \) dimensions, some insight into the underlying structure of the transition has been achieved within the multifractal formalism [64]. As we conjecture the transitions for the DL and SDL to be essentially analogous, we expect to find similarities to the analysis that has been done before for DLs [14] but also deviations where the obvious geometrical differences become important. The idea is that different moments \( Y_q \) of the statistical weights

\[
Y_q(L) = \sum_i \text{Prob}(z(L) = z_i)^q
\]

are dominated by different regions of the support and, thus, show a different scaling with the system size \( L \). The probability is given by the ratio of restricted and unrestricted partition function \( \text{Prob} [z(L) = z_i] = w_L(z_i) / Z_L(z_i) \). The sum goes over all the possible ending points in the numerical simulation, whereas for the continuous analytical problem \( Y_q \) would be defined by an integral over \( z \).

In the high-temperature phase, the weights \( w_L(z) \) obey the scaling form \( w_L(z) = L^{-\eta} \Omega(z / L^\eta) \) with the return exponent \( \chi \) [defined by \( w_L(0) \sim L^{-\chi} \) as introduced above]. It is straightforward to see that the \( Y_q \) then scale according to

\[
Y_q(L) \big|_{T > T_c} \sim L^{-(q-1)\chi} = L^{-(q-1)\beta^d}
\]

in the high-temperature phase. In the low-temperature phase, the line is localized and, therefore, the \( Y_q \) remain finite as \( L \to \infty \). At criticality the \( Y_q \) are diverging, but the quenched disorder is relevant, and it becomes important how the (necessary) average over realizations of the disorder is computed. A common question regarding systems with disorder is whether a certain quantity is self-averaging. If so its typical and average values, the latter of which could be dominated by rare events, should be identical. For the \( Y_q \) a reasoning like this motivates the definition of

\[
Y_q^{av} = \frac{1}{T \approx T_c} \sum_{T \approx T_c} L^{-\tau(q)} = L^{-(q-1)D(q)}, \quad (56a)
\]

\[
Y_q^{hyp} = \exp \ln Y_q \big|_{T \approx T_c} \sim L^{-\tau(q)} = L^{-(q-1)D(q)}, \quad (56b)
\]

where the definition of \( D(q) \) is such that it includes the obvious case of \( q = 1 \), where \( Y_1 = 1 \) and \( \tau(1) = 0 \) by definition. The \( D(q) \) are referred to as generalized dimensions [64], and the function \( D(q) \) discriminates between monofractal \( D(q) = \text{const.} \) as for \( T > T_c \) and multifractal behavior \( D(q) \neq \text{const.} \). The interpretation of the \( D(q) \) as (fractal) dimensions of subsets is rather peculiar as we are dealing with a probability distribution or measure. Nonetheless, it is useful as it requires \( D(q) \) to be monotonically decreasing because none of the subsets can have a larger dimension than their union. There are at least two special values of \( D(q) \) with an obvious meaning: \( D(0) \) is the Hausdorff dimension of the support and,
thus, directly related to the geometry of the system and \( D(1) \) is called the information dimension as it appears like a dimension in the Shannon information entropy.

\[
s = - \sum_{i} w(z_i) \ln w(z_i) = - \partial_{q} Y_{q} \bigg|_{q=1} \approx D(1) \ln L. \quad (57)
\]

However, \( D(1) \) cannot be computed directly (\( Y_{1} \equiv 1 \)) but only as an analytic continuation of \( D(q) \). A measure is called fractal if and only if \( D(0) > D(1) \) [65].

In Fig. 17, we present numerical results for \( D(q) \) and \( \tilde{D}(q) \) for a SDL in disorder at criticality. As a control for the numerics one can use the information dimension \( D(1) \), which should coincide with its high-temperature value \( D(1)_{T=0} = 3/2 \) at criticality. The data appear to resemble the Anderson transition-like scenario that has been reported for the DL [14] with \( D(q) = \tilde{D}(q) \) for \( q < q* \approx 1.5 \). The separation of \( D(q) \) and \( \tilde{D}(q) \) indicates different behaviors of typical and average values of \( Y_{q>q*} \), from which one is tempted to conclude that these quantities are not self-averaging. Furthermore, both \( Y_{q}^{av} \) and \( Y_{q}^{yp} \) are diverging faster than exponential for \( q < 0 \), which leads to \( D(q < 0) = D(q < 0) = \infty \). The information dimension \( D(1) \) is measured to be about 1.4 and, therefore, does not coincide with the expected high-temperature value 3/2. However, this could be a numerical artifact from limited system sizes.

For Anderson transitions, the finite-size scaling of the \( Y_{q} \) at criticality does involve the multifractal spectrum but only one correlation length exponent \( v \) [11,66],

\[
Y_{q} = L^{-\frac{q}{v}} f \left[ \left( \frac{T_{c} - T}{L^{1/v}} \right) \right].
\quad (58)
\]

Thus, it allows for a completely independent validation (cf. Fig. 18) of the critical temperature and the correlation length exponent, yielding a more exact value \( T_{c} \approx 1.44 \) compatible with our result from Secs. VII B and VIII D and, unambiguously, a value \( 1/v \approx 0.5 \) for the correlation length exponent.

\[3\]The Hausdorff dimensions for the measures related to DLs and SDGs with \( d \) transverse dimensions are \( D_{r}(0) = d \) and \( D_{r}(0) = 2d \) (\( z \) and \( v = \partial_{z} \)), respectively, which gives \( D_{r}(0) = 3 \) for DLs in \( 1 + 3 \) dimensions and \( D_{r}(0) = 2 \) for SDGs in \( 1 + 1 \) dimensions.

An equivalent description of the multifractal nature is related to the Legendre transform \( f(\alpha) \) of \( \tau(q) \) given by

\[
q = f'(\alpha), \quad (59a)
\]

\[
\tau(q) = \alpha q - f(\alpha). \quad (59b)
\]

The function \( f(\alpha) \) is called the singularity spectrum, because it gives the number \( N(\alpha) \sim L^{f(\alpha)} \) of points \( z \), where the weight \( w(z) \) has a singularity \( w \sim L^{-\alpha} \). A Legendre transform of the measured \( \tau(q) \) and \( \tilde{\tau}(q) \) would require an analytical continuation and, thus, be very error prone. Fortunately, it is possible to directly measure \( \alpha(q) \) and \( f(q) \) [67] and, thus, the singularity spectrum \( f(\alpha) \) parametrically:

\[
f(q) = - \lim_{L \to \infty} \sum_{i} \mu(q,z_{i}) \ln \mu(q,z_{i}) / \ln L, \quad (60a)
\]

\[
\alpha(q) = - \lim_{L \to \infty} \sum_{i} \mu(q,z_{i}) \ln w(z_{i}) / \ln L, \quad (60b)
\]

with

\[
\mu(q,z_{i}) = w(\xi) w(\xi_{j}) \left( \sum_{j} w(\xi_{j}) \right). \quad (60c)
\]

This method of computing \( f(\alpha) \) gives the Legendre transform of \( \tau(q) \), because Eq. (59) implies (omitting the limits)

\[
\tau(q) = \alpha(q) q - f(q)
\]

\[
= - \ln \sum_{i} w^{q}(z_{i}) / \ln L
\]

\[
= - \ln Y_{q}(L) / \ln L, \quad (60d)
\]

in agreement with Eq. (56b), and, therefore, \( f(\alpha) \) corresponds to typical values of \( Y_{q} \). Note that \( w(\xi) = \tau(q) \) is fulfilled by construction. This is the common definition of the (multifractal) singularity spectrum that is also applicable for nondisordered systems.

Here disorder is relevant and we need to capture not only the typical but also the (differing, cf. Fig. 17) average behaviors. Analogously to Eq. (60), we derive the following computation of the Legendre transform \( \tilde{f}(\tilde{\alpha}) \) of \( \tilde{\tau}(q) \) (we distinguish
measured). We use Eq. (56) and the inverse transform of (59) 

\[ f(\alpha) \] (red) from which Fig. 19 was created. The contact point is at \( \alpha = f(\alpha) \approx 5.5 \) and 5. It shows lines corresponding to \( D(\alpha) = D(2) = 2, f(\alpha) = D(1) \), and \( f(\alpha) = \alpha \); see text. The bisector and \( f(\alpha) \) touch at an \( \alpha \) that is slightly smaller than \( \alpha = 1.5 \). We consider this to be wrong, but it is consistent with the determined generalized dimensions; see also Fig. 20.

between \( \alpha \) and \( \tilde{a} \) to avoid ambiguities as both are directly measured. We use Eq. (56) and the inverse transform of (59)

\[ \tilde{f}(q) = -\ln \sum_i w^q(z_i)/\ln L, \] (61a)
\[ \tilde{a}(q) = \tilde{f}(q) = -\sum_i w^q(z_i)\ln w(z_i)/(\ln L) \sum_i w^q(z_i), \] (61b)
\[ \tilde{f}(\tilde{a}) = \tilde{a}q - \tilde{f}(q) = -\sum_i w^q(z_i)(\ln w^q(z_i)/\sum_j w^q(z_j) - 1)/\ln L. \] (61c)

The spectrum \( f(\alpha) \) is shown in Fig. 19. Its shape matches the expectations that origin from general properties and the known results for the DL [14] and is consistent with the results for the Legendre transform \( \tau(q) \). Our results show that \( f(\alpha) \) is a monotonic function starting at a finite \( \alpha_{\min} = D_{\min}(q) \approx 0.8 \), which is close to the DL value \( \alpha_{\min} \approx 0.77 \) and ending at \( \alpha_{\max} = \infty \), which corresponds to the infinitely large values \( \tau(q) \) for \( q < 0 \). The maximum value of \( f(\alpha) \) is \( f(\alpha \to \infty) \to D(0) \) and it touches the bisector \( f = \alpha \) around \( \alpha = D(1) \approx 1.4 \), thus confirming the previously found deviation \( D(1) \neq 3/2 \). We see no indication that \( f(\alpha) \) becomes negative somewhere, which would describe rare events (number decreases exponentially in \( L \)), but this is to be expected as \( f(\alpha) \) contains the typical behavior and should not become negative [14]. The average behavior that leads to \( f(\alpha) \) does include rare events and does not seem to have a finite \( \alpha_{\min} \), implying \( D(\infty) = 0 \). We back this by noting that the we can achieve a good fit of the data for \( \tau(q) \) and \( \tilde{f}(q) \) at large \( q \) (we used \( 5 < q < 20 \)) with an monomial Ansatz \( f(q) = aq^b \). For an approximate analysis we round \( b \) to one decimal, giving

\[ \tau(q \gg 1) \approx aq, \] (62)
\[ \tilde{f}(q \gg 1) \approx aq^{2/5}, \] (63)
with some constants \( a, \tilde{a} \). We apply the Legendre transform and get for \( q \gg 1 \)

\[ \alpha(q) \approx a \approx 0.81 \]
\[ f(\alpha) \approx 0 \]
\[ a(q) \sim q^{-3/5} \to 0 \]
\[ f(\alpha) \sim -\tilde{a}^{-2/3} \to -\infty. \]

In summary, we see a (Anderson transition-like) scenario in the multifractal analysis of the localization transition of a SDL in \( 1 + 1 \) dimensions which is very similar to the findings in Ref. [14] for a DL in \( 1 + 3 \) dimensions despite obvious differences due to the different geometry. In particular, the multifractal structure of the statistical weights differs between typical and average values both for DLs and SDLs. This becomes apparent in the generalized dimensions \( D(q), \tilde{D}(q) \), which differ for \( q \approx 1.5 \). We also find a matching critical correlation length exponent \( v \). Additionally, we showed that the average behavior is significantly influenced by rare events, leading to negative values in \( f \), the Legendre transform of \( D(q) \). The significance of rare (extreme) events at criticality is in accordance with our findings for the energy distribution.

VIII. CONCLUSION

We studied SDLs in \( 1 + d \) dimensions subject to quenched short-range random potential analytically and numerically. Using Flory-type scaling arguments and a replica calculation, we show that, in dimensions \( d > 2/3 \), a localization transition exists from a high-temperature phase, where the system is essentially annealed, to a disorder-dominated low-temperature phase. The low-temperature phase is characterized by large free-energy fluctuations with an exponent \( \omega > 0 \) and a roughness exponent \( \zeta \), which slightly exceeds the thermal value \( \zeta_{\text{th}} = 3/2 \) for a SDL. Flory arguments suggest \( \zeta = 7/(4 + d) \) for a SDL. Both exponents are related by \( \omega = 2\zeta - 3 \).

We find a reduction of the persistence length of a stiff directed line by disorder. In the low-temperature phase, the persistence length is disorder induced and temperature independent.

We also performed variational replica FRG calculations. The replica approach gives no conclusive results on the exponents for \( d > 2/3 \) but supports the existence of a localization transition. The FRG calculation is performed for \( D + d \) dimensional manifolds governed by bending energies with the SDL corresponding to \( D = 1 \) and employs an expansion in \( \epsilon = 8 - D \) dimensions. For a SDL in \( 1 + d \) dimensions, the FRG result suggests the existence of an upper critical dimension \( d_u < 1 \), which we can rule out by numerical calculations.
This strongly supports a relation between DLs in $1+3d$ and SDLs in $1+d$ dimensions, which is based on identical return exponents $\chi$ for two replicas to meet. The validity of a relation based on properties of a single replica pair suggests that the critical properties of DLs in a short-range random potential are governed by replica pair interactions. The mapping can make DL transitions in high dimensions computationally accessible, which we demonstrated in showing that the two-replica overlap provides a valid order parameter across the localization transition of SDLs in $1+1$ dimensions. Furthermore, the importance of pair interactions suggests that the critical temperature for DLs in random potentials is indeed identical to the temperature below which the ratio of the second moment of the partition function and the square of its first moment diverges, which implies that the localization transition temperature $T_c$ in a random potential equals the replica pair transition temperature $T_2$ for replica pair binding. The equality $T_2 = T_c$ has been originally put forward and verified numerically for DLs in $1+3$ dimensions [11,14]. Using the numerical transfer matrix approach, we could verify this conjecture also for SDLs in $1+1$ dimensions. Our findings are summarized schematically in Figs. 2 and 21.

The binding transition of DL pairs becomes discontinuous for $d > 4$ and, analogously, the binding of SDL pairs for $d > 4/3$ [28,63]. Because DLs in random potentials are equivalent to the KPZ equation [2], the validated relation to the SDL suggests that the roughening transition of the KPZ problem could acquire similar discontinuous features for $d > 4$ dimensions. Thus, $d = 4$ would remain a special dimension, although it is not the upper critical dimension [68].

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