Scaling in a two-component surface-growth model

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(Received 7 May 1998; revised manuscript received 26 June 1998)

We study scaling in kinetic roughening and phase ordering during growth of binary systems. We use a 1+1-dimensional single-step solid-on-solid model with Ising-like interaction between two components. We observed that the model exhibits a crossover from an intermediate regime, with effective scaling exponents for kinetic roughening significantly larger than for the ordinary single-step growth model, to an asymptotic regime with exponents of the Kardar-Parisi-Zhang class. The crossover time and length are exponentially increasing with the strength of interaction K. For a given large K, scaling with enhanced exponents is valid over many decades. The effective scaling exponents are continuously increasing with K. Surface ordering proceeds up to the crossover. The average size of surface domains increases during growth with the exponent close to 1/2; the spin-spin correlation function and the distribution of domains obey scaling with the same exponent.

I. INTRODUCTION

Growth by vapor deposition is an effective process for producing high-quality materials. The microscopic mechanisms of growth were intensively studied in the past mainly in the case of homoepitaxial growth.1–3 However, a common situation in nature as well as in modern technologies is the growth of binary or more component systems. Due to the nonequilibrium nature of growth the properties of the resulting film can be very different from the properties of equilibrium bulk material (surface alloys that have no bulk analog can be formed, highly anisotropic structures can be prepared, etc.). The problem of growth in a system with two or more components is of great practical importance but it is also interesting from a purely statistical-mechanical point of view, because the growth process may belong to a new universality class4,5 and such a system might exhibit a nonequilibrium phase transition between low- and high-temperature phases.

There are two interfering phenomena in the growth of binary systems: kinetic roughening and phase ordering. During growth the initially flat surface is becoming rough. This is called kinetic roughening. It has been found that this process often fulfills the invariance with respect to scaling in both time and length. Let us consider a surface in a d-dimensional space given by a single-valued function h(\(r,t\)) of a \(d'\)-dimensional (\(d = d' + 1\)) substrate coordinate \(r\). The surface roughness is described by the surface width \(w(t,L) = \langle \sqrt{h^2 - L^2} \rangle\), where \(t\) is the time, \(L\) is a linear size, and the bar denotes a spatial average, \(\langle \cdots \rangle\) a statistical average. The surface width often obeys the dynamical scaling law \(w(t,L) \sim L^z f(t/L^\alpha)\), with the scaling function \(f(x)\) fulfilling \(f(x) = \text{const.}, x \gg 1\) and \(f(x) \propto x^\beta, x \ll 1\) (\(\beta = \alpha/z\)). Dynamical scaling allows us to classify growth processes into dynamical universality classes according to values of exponents \(\alpha\) and \(z\) (or \(\alpha\) and \(\beta\)).5,7 This scaling has been observed in a wide variety of growth models and many of them belong to the Kardar-Parisi-Zhang (KPZ) universality class.5 There has been considerable effort in order to find different possible universality classes.

On the other hand, the process of ordering in ordinary binary systems with a constant number of particles can lead to phase separation. In the case of phase separation dynamical scaling exists as well, e.g., in the Ising model at low temperatures.9 In phase ordering, the characteristic length \(D\) is the average size of domains formed by particles of one type. It increases with time as a power law, \(D \propto t^\psi\). The dynamics can be classified according to values of the exponent \(\psi\). Phase ordering is usually a bulk process. However, one can also study growth-induced ordering at surfaces. In this case the evolution of the average domain size on the surface is of interest.

On the microscopic level, growth is usually investigated using discrete growth models. Although several growth models for binary systems were introduced in various contexts, e.g., for the study of phase separation during molecular-beam epitaxy,10 or growth of binary alloys,11 our understanding of growth of composite systems is still at the beginning. In particular, little is known so far about kinetic roughening in two-component growth models. This problem was probably first considered by Ausloos, Vandewalle, and Cloots.4 They introduced a generalization of the Eden model and coined a magnetic Eden model, which contains two types of particles with the probabilities of growth given by the Ising-like interaction. Ausloos, Vandewalle, and Cloots suggested that the
magnetic Eden model does not belong to the KPZ universality class. Recently El-Nashar, Wang, and Cerdeira\textsuperscript{5} studied kinetic roughening in a ballisticlike two-component growth model with a varying probability for deposition of a given type of particle. They observed that the exponent $\beta$ changes with the probability and argued that kinetic roughening no longer follows the KPZ scaling law. Although the phase ordering was apparently present it was not studied in these works.

In this paper we concentrate on the situation where both processes, kinetic roughening as well as phase ordering, are important and affect each other. We investigate scaling in both roughening of the surface and phase ordering. We use a one-dimensional two-component single-step solid-on-solid growth model that we recently introduced.\textsuperscript{12} It is particularly convenient for the study of asymptotic scaling behavior. Here we present results of extensive numerical simulations that complement preliminary results published elsewhere.\textsuperscript{12,13}

The paper is organized as follows. In Sec. II, our model and the measured quantities are defined. In Sec. III, results of our Monte Carlo simulations are described. Section IV contains a discussion. Finally, a summary is given in Sec. V.

II. MODEL AND MEASURED QUANTITIES

A. Modeling two-component growth

There is a large variety of single-component growth models that can be potentially generalized to the multicomponent case. Moreover, there are different possible means of generalization. One usually tries to use a model which is as simple as possible and still contains important features. Our aim is to find such a model for the study of scaling during two-component growth.

The commonly used approximation is a discrete model with the so-called solid-on-solid (SOS) condition. It means that the surface is described by a single-valued discrete function $h(i)$. The index $i$ is the horizontal coordinate that labels sites of the substrate. The rates for elementary growth processes depend usually only on the values of $h$ in a neighborhood of the initial and, in the case of diffusional barrier, possibly also of the final position of a particle. The situation is more complex for a two-component system because the rates of elementary growth processes depend not only on the geometry but also on the local composition. In practice it means that we need to store more information, e.g., the composition in an additional data array. Let us denote the type of a particle by a variable $\sigma$ that can have value $+1$ or $-1$. The geometry of the surface at time $t$ is described by the function $h(i,t)$ and the composition of the deposit is represented by a function $\sigma(i,y)$, where the two variables $i$ and $y$ are horizontal and vertical coordinate of a particle, respectively. The variable $y$ is restricted only to values from 1 to $h(i,t)$.

Storing the composition of the whole deposit is possible only for relatively small sizes of the substrate and not too many monolayers (ML) of deposited particles. When we study scaling phenomena where the behavior for very large samples and long times (i.e., many ML) is investigated, too much memory would be required. However, when bulk processes can be neglected, it is sufficient to remember only the composition within a certain finite depth under the surface, because deeper layers cannot affect surface growth. The complication is that the depth that should be stored is in general not well defined and, in principle, it may be unlimited. For example, to describe the rate for a process in which a particle is moving from or to the bottom of a step we need to store the composition in the depth equal to the maximum step size. But it is known that in some models of MBE growth\textsuperscript{14} based on an unrestricted SOS model steps of an arbitrary size can be present. Natural solution of this technical obstacle is to use so-called restricted SOS model in which possible configurations are limited by an additional constraint $|h(i) - h(j)| \leq N$; $i$ and $j$ being nearest neighbors and $N$ a given integer.

B. Two-component single-step model

Our model is based on the simplest restricted SOS model, the so-called single-step solid-on-solid model. The difference of heights between two neighboring sites is restricted to $+1$ or $-1$ only. The advantage of this choice is that if we restrict ourselves to nearest-neighbor interactions between particles then we can define rates for elementary moves of particles using only the composition on the surface. Hence, the rates at any time are given by the surface profile $h(i,t)$ and the composition \textit{only} on the surface, which is described by the field $\sigma(i,t)$ of the same dimensionality as $h(i,t)$. We call such a model the \textit{two-component single-step} (TCSS) model.

The growth rules depend in general on the physical situation under study. We consider a rather simple case that, however, allows us to evaluate the effect of ordering on kinetic roughening. As indicated above, we do not allow processes that lead to exchange of particles in bulk. This is well justified because rates for such processes are usually several orders of magnitude lower than for processes on the surface. We also do not include surface diffusion. This is a serious restriction from the point of view of application to epitaxial growth. However, it is well known that the study of scaling in models with diffusion is demanding on computer power already in the case of one-component growth\textsuperscript{14} and that it is difficult to obtain results with good statistics. We consider, rather, the condensation-evaporation dynamics. Moreover, we restrict ourselves here to the pure growth situation. Evaporation can be included but we expect that it will not change the scaling behavior provided deposition occurs more frequently than evaporation, i.e., the average surface height is not constant.

Hence, particles are only added during the evolution. Due to the single-step constraint a particle can be added only on a site at a local minimum of height. We call such site the growth site. Once the position and the type of the particle are selected, they are fixed forever. The probability of adding a particle of type $\sigma$ to a growth site $i$ depends only on its local neighborhood and is controlled by the change of energy of the system after deposition of a new particle. The energy is given by Ising-like interaction. The probability is proportional to $\exp\left(-\Delta E(i,\sigma)/k_B T\right)$, where $k_B$ is Boltzmann’s constant, $T$ is temperature, and $\Delta E(i,\sigma)$ is the change of energy.\textsuperscript{15}

We describe our growth model for simplicity in \textit{1+1} dimensions but it can be straightforwardly generalized to any dimension. Several realizations of the single-step geometry
The sum can be deposited.

are possible in 1+1 dimensions (Fig. 1) leading to three different variants of the TCSS model. They differ in the number of nearest neighbors of a new particle. While in variant B [Fig. 1(b)] there is an ambiguity in the type of a newly deposited particle if two neighbors are of opposite types, this is not the case in variant A [Fig. 1(a)] in which the number of nearest neighbors is odd. We expect that the effect of ordering dynamics is stronger for variant A than for variant B. Variant C [Fig. 1(c)] is technically slightly more complicated to simulate due to the varying number of nearest neighbors of a deposited particle. Therefore, we consider variant A with three nearest neighbors, which is represented as stacking of rectangular blocks with the height equal to double the width [Fig. 1(a)]. Nevertheless, we expect similar asymptotic scaling behavior for all three variants.

The expression for the energy change is

$$\frac{\Delta E(i, \sigma)}{k_B T} = -K\sigma[\sigma(i-1) + \sigma(i) + \sigma(i+1)] - H\sigma.$$

(1)

Here, $K$ is a dimensionless coupling strength and $H$ is the bias leading to preferential deposition of particles of a selected type (+1 for positive, −1 for negative $H$). In analogy with magnetic systems we will call $H$ an external field. The sum $\sigma(i-1) + \sigma(i) + \sigma(i+1)$ contains types of particles on the surface within nearest neighbors of the growth site (which are three in the chosen variant: left, bottom, and right); see Fig. 2.

C. Measured quantities and simulation procedure

The evolution of surface morphology is affected by composition of the surface and vice versa. We measure quantities describing both aspects. The surface morphology is characterized by the surface width $w(t, L)$ defined in the Introduction, and by the height-difference correlation function $G(r, t) = (1/L) \sum_{i=1}^{L} [(h(i+r,t) - h(i,t)]^2$, which is expected to obey a scaling relation $G(r, t) \sim r^{2\alpha} g(r/t^{1/2})$, with the scaling function $g(x)$ constant for $x \ll 1$ and $g(x) \sim x^{-2\alpha}$ for $x \gg 1$. In the case of phase ordering we restrict ourselves to the composition of the surface, because only it affects the evolution of the surface profile. We measure two quantities defined on the surface: (i) the average of surface domain sizes, and (ii) the correlation function $S(r, t)$ analogous to the spin-spin correlation function used in magnetic systems. We call the surface domain a compact part of the surface composed of particles of the same type. The size of the domain $d$ is measured along the surface. The average size of the surface domains depends on time, on the strength of coupling, on the external field, and also on the initial composition of the substrate. We denote the statistical average of this quantity by $D = \langle d \rangle$. The correlation function $S(r, t)$ is defined as follows: $S(r, t) = (1/L) \sum_{i=1}^{L} \langle \sigma(i+r,t) \sigma(i,t) \rangle$.

We performed simulations for various values of the coupling strength $K>0$ and mainly for zero external field $H$. System sizes ranged from $L = 250$ to $L = 80 \, 000$, and the number of monolayers deposited was up to $3 \times 10^5$, and for small systems up to $4 \times 10^6$ ML. We measured time $t$ of the simulation in ML. A statistical average was obtained by averaging over a varying number of independent runs [from ten for $L = 80 \, 000$ up to several thousand for $L = 250$].

Growth starts on a flat surface, however, in two-component models the evolution strongly depends on the initial composition of the substrate. Here we consider two possibilities: (i) a neutral substrate, i.e., substrate without any interaction with deposited particles (in this case the system orders spontaneously from the beginning), and (ii) an alternating substrate, with the alternating types of particles. The case of a homogeneous substrate composed of one type of particles is reported elsewhere.

III. RESULTS

A. Evolution of morphology

Figure 3 shows examples of time evolution of the surface morphology and composition for selected coupling strengths.
and external fields. Note that the times for which surface profiles are shown increase as a power law. Visual inspection of many configurations leads to the following observations. In the case of zero external field (upper panel), we can see that with increasing coupling the surface is becoming more and more rough (faceted) and at the same time more and more clean columnar structures are formed; the anisotropy induced by growth is more pronounced. The average width of the columns increases with time. We can also see that, for a given time, the average width of the column is decreases with the coupling strength. This seems to be in contradiction with the expectation that ordering should be more pronounced for stronger coupling. Notice also that for larger coupling we have observed that the surface width exhibits a crossover in time. At first the width increases with an effective exponent $\beta_{\text{eff}} > \frac{1}{3}$, but after certain time $t_{\text{cross}}$ it crosses over back to $\beta_{\text{KZ}} = \frac{1}{3}$. This can be clearly seen in Fig. 4 (curves for $K = 0.7, 1.0$, and $1.3$) where we plotted the time dependence of the quantity $w(t)/t^{1/3}$ for several couplings in order to compare evolution of the surface width with the KZ behavior. Notice also that the absolute value of $K$ at a given time increases with $K$. The crossover from the regime with the enhanced $\beta_{\text{eff}}$ to the regime with $\beta = \frac{1}{3}$ is definitely not a finite-size effect. We checked that the time $t_{\text{cross}}$ is the same for different system sizes.

From Fig. 4 we can also see that $t_{\text{cross}}$ is increasing with the coupling. We were not able to observe the crossover to the KZ behavior for coupling $K \geq 2$ on the time scale of our simulations.\(^{18}\) In order to estimate the time needed we plotted the dependence of $t_{\text{cross}}(K)$ on $K$ (Fig. 5). It can be fitted as $t_{\text{cross}}(K) \propto e^{0.8K}$. When we extrapolate this data to $K = 2$ we get $t_{\text{cross}}(K = 2) > 10^9$, which is longer then the time of our simulation.

We found that for any $K > 0.7$ there is a time interval extending over several decades in which we can well fit our data as a power law with the exponent $\beta_{\text{eff}} > \frac{1}{3}$, before there is the crossover to the KZ behavior or our simulation stops. The question remains as to whether $\beta_{\text{eff}}$ goes to a specific value for large $K$. We did not find any indication that $\beta_{\text{eff}}$
saturates to a certain value, at least for the investigated range of the coupling constant $0 < K < 3$. The effective exponent $\beta_{\mathrm{eff}}$ is an increasing function of $K$ (see inset in Fig. 4). We attribute the rather large value of $\beta_{\mathrm{eff}}$ for strong coupling to pinning of the surface at domain boundaries.

2. Height-difference correlation function

The second exponent for kinetic roughening is the roughness exponent $\alpha$. It can be calculated from the dependence of the saturated surface width on the system size. We used here an alternative and often more accurate way. We calculate $\alpha$ from the spatial dependence of the height-difference correlation function $G(r,t) \propto r^{2\alpha}$ in the long-time limit.

The obtained exponents $\alpha$ also depend on coupling. Exponent $\alpha$ has a value close to $\alpha(KPZ) = \frac{1}{2}$ for weak coupling. For stronger coupling, we have found that when the system is sufficiently large there is a crossover behavior in the spatial dependence of the height-difference correlation function (see Fig. 6). On a small length scale it increases faster than $r$, and for a sufficiently large $K$ we can fit it as a power law $r^{2\alpha_{\mathrm{eff}}}$ with an effective exponent $\alpha_{\mathrm{eff}} > \frac{1}{2}$. However, if the distance is larger than a certain length $l_{\text{cross}}$, the form of the function crosses over to power law with the exponent which is close to $\frac{1}{2}$, i.e., to the KPZ behavior (cf. data for $K = 1$ in Fig. 6).

If the coupling is strong ($K \geq 2$) we do not see the crossover but only the larger exponent $\alpha_{\text{eff}} = 1$. We expect that this is because even the system size $L = 80 \, 000$ and the time 262 144 ML are not large enough to get into the crossover regime. We have found that $\alpha_{\text{eff}}$ is increasing smoothly with the strength of coupling from $\alpha(KPZ) = \frac{1}{2}$ to a value slightly less than 1 (inset in Fig. 6). The value $\alpha = 1$ is a natural limit imposed by the single-step constraint.

3. Scaling

Having both exponents $\alpha$ and $\beta$, we can try to verify scaling. For weak coupling, the exponents are close to the KPZ exponents and scaling with these exponents is satisfied. For larger coupling, when the crossover is observed, we cannot get scaling for all times and lengths, nevertheless for long times and large lengths the KPZ scaling is valid. When coupling is sufficiently strong then the behavior characterized by the enhanced exponents extends over many decades of time and length and looks practically as asymptotic. Then we can ask ourselves if there is scaling with the new exponents that is satisfied on this scale. In order to show that there is scaling, we should get a data collapse. We have found that indeed we get the data collapse over many decades in the strong-coupling regime ($K \geq 2$). As an example we show in Fig. 7 the data collapse of the height-difference correlation function $G(r,t)$ obtained for $K = 2$ with exponents $\alpha = 0.97$ and $z = 1.98$. For different $K$ we need different exponents, e.g., for $K = 3$ we get the best data collapse for $\alpha = 0.99$ and $z = 1.7$. The question remains what is the true asymptotic scaling in the strong-coupling regime. We expect that for any $K$ there is a crossover to the KPZ scaling (although $t_{\text{cross}}$ may be astronomically large), and the asymptotic behavior will belong to the KPZ class.
The time evolution of the average surface domain size $D$ for several values of the coupling constant is shown in Fig. 8. We can see that the behavior again depends on $K$. For small coupling, $D$ at first increases; however, after some time $t^{(D)}(K)$ the surface domain size saturates to a value $D_{\text{sat}}(K)$. We have checked that the saturation is not a finite-size effect (see Ref. 12—inset in Fig. 3), but it is an intrinsic property of the model. Both the time $t^{(D)}(K)$ and the saturated value $D_{\text{sat}}(K)$ rapidly increase with $K$ (Fig. 5). We have found that the dependence on $K$ can be well fitted by exponential functions: $D_{\text{sat}}(K) \sim e^{\gamma_1 \pm 0.00} K$ and $t^{(D)}(K) \sim e^{\gamma_2 \pm 0.00} K$. We were not able to observe saturation for $K=2$ because of a prohibitively long simulation time needed. Hence, for $K \geq 2$ the domain size increases over the time of our simulation. But we believe that the evolution of the surface domains is analogous to the evolution of domains in the one-dimensional Ising model and that in the long-time limit it will saturate for any $K$ (Ref. 19).

The evolution of domains is strongly affected by the initial conditions and a certain transition time is needed before a value independent on the initial state is reached. This phenomenon is similar to what we have observed in the case of the surface width. The transition time is increasing with coupling and can be quite long, e.g., for $K=3$ it is several hundred ML.

We have found that for $K \geq 1$ there is a time interval in which the increase of the average domain size can be fitted by a power law, $D(t,K) \sim t^{\psi_{\text{eff}}(K)}$ with an exponent $\psi_{\text{eff}}$ depending on $K$. For large $K$, the exponent $\psi_{\text{eff}}$ seems to saturate to a value slightly smaller than $\frac{1}{2}$ (see inset in Fig. 8). This is the same exponent as for the Ising model with a nonconserved order parameter. 

2. Distribution of surface domain sizes

The average surface domain size contains information about the formation of domains during growth. However, it is not clear solely from that quantity whether the domains form a kind of periodic structure with a typical domain length or if the domain sizes are more random. In order to obtain more information about the domains, we measured the probability distribution $P(d,t)$ of the domain size $d$ as a function of time. We performed simulations for three values of coupling $K=1, K=2$, and $K=3$, the system size $L = 1000$, and times up to 32 768 ML. To get good statistics we had to make the average over 10 000 independent runs. We observed that at initial times there is a rather sharp asymmetric peak with the position shifting to higher values of $d$ with increasing time. During the time evolution the amplitude of the peak decreases and the peak becomes broader and eventually disappears. The time scale for this behavior depends on $K$.

We try to rescale our data extending over many orders of magnitude and to see whether or no there is scaling. We applied a scaling formula of the form

$$P(d,t) = F(d/D(t))/D(t),$$

with a scaling function $F$. The average domain size $D(t)$ is in fact equal to the average computed from the distribution $P(d,t)$.

In Fig. 9 the rescaled distribution of domain sizes for $K=1$ is shown as a function of the variable $x=d/D$, for times from 4 to 32 768. We can see that the peak at around $x = 0.4$ exists only for short times and for a time of approximately $t=64$ it changes to a small plateau that vanishes for longer times. The distribution of domain sizes at long times converges to a function that we found to be well fitted by an exponential. Moreover, we found that the peak vanishes around the time $t^{(D)}$ when the average domain size saturates.

The results for $K=2$ are shown in Fig. 10. We can see that the scaling formula (2) is satisfied from times 8 to 6144. The data for longer times are not shown here because after 10 000 independent runs they still have too much noise. The scaling function $F(x)$ has a maximum again near $x = 0.4$. This indicates the creation of a quasiregular domain structure, but the distribution has a very broad tail for larger $x$. 

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**FIG. 8.** Time evolution of average surface domain size for coupling constants $K= 0.7$ ( ), 1.0 (□), 1.3 (○), 2.0 (△), 3.0 (○) and zero external field, $L=80000$. Inset: Effective exponent $\psi_{\text{eff}}$ as function of coupling $K$.

**FIG. 9.** Distribution of domain sizes for $K=1$ and times 4 (■), 8 (●), 16 (○), 32 (○), 64 (□), 128 (○), 256 (×), 512 (▽), 1024 (+), 6144 (△), 32768 (▲). Inset: Detail of the distribution. The symbols are the same as in the main graph.
extending up to $x \approx 4$, which makes the domain structure irregular. We compared the scaling for different values of $K$. For times shorter than the saturation time the scaling function does not depend on $K$, which is demonstrated in the inset in Fig. 10 for $K=1.0, 2.0$, and $3.0$.

Behavior similar to that in the strong-coupling regime was observed for the kinetic Ising model in one dimension. Numerical simulations of the Ising model at zero temperature\cite{20} show that the distribution of domain sizes obeys the form (2) with a peak around $d/D \approx 0.4$. Also, the form of the function $F(x)$ found in Ref. 20 looks very similar to our results. On the contrary, for any nonzero temperature it may be shown analytically that the distribution of domain sizes in equilibrium, i.e., in the infinite time limit, is exponential (see, e.g., Ref. 21). Analytical results for the zero-temperature kinetic Ising model\cite{22} give asymptotic results $F(x) \approx \pi x$ for $x \rightarrow 0$ and $F(x) \approx \exp(-A x + B)$ for $x \rightarrow \infty$, with $A = \frac{1}{2} \alpha(3/2)$ $= 1.30618 \ldots$, and $B = 0.5971 \ldots$, which is in good agreement with our results, as Fig. 10 shows.

Summing up, we should again distinguish two time regimes. At initial times, the form of the distribution of domain sizes as a function of $x$ does not change and is characterized by a pronounced peak. Scaling (2) holds and the only change during the time evolution is the increase of the average domain size $D$. When $D$ begins to saturate, the peak vanishes and, in the saturated regime, the distribution of domain sizes is exponential. The reason that, in the case of strong coupling ($K = 2.0, 3.0$), the peak remained for all times observable is simply that the duration of the simulation is still much shorter than the saturation time.

3. Correlation function

The correlation function $S(r,t) = (1/L) \sum_{i=1}^{L-1} \langle \sigma(i+r,t) \sigma(i,t) \rangle$ decays nearly exponentially with the distance $r$. The decay is characterized by the correlation length $\xi(t,K)$, which we computed by fitting the data to the function $\exp[r/\xi(t)]$ in the interval $(0,r_s)$ in which the decay is well described by an exponential. According to our experience, a good recipe for fixing the interval $(0,r_s)$ is to find the distance $r_s(t)$ as the minimal distance for which $S(r_s(t),t)$ $\approx 0.05$. This is done for a given time and coupling.

The time behavior of the correlation length $\xi(t,K)$ is similar to that of the average domain size $D(t,K)$. It increases in time as a power law $\xi(t,K) \propto t^{k_{v_{\text{eff}}}(K)}$ (we found $k_{v_{\text{eff}}}=0.5$ for $K = 2$). For a small coupling constant $K$, $\xi(t,K)$ saturates to a finite value $\xi_{\text{sat}}(t)$ at about the same time $t_{\text{sat}}(D)$ as the saturation of the average surface domain size occurs. The saturation for large values of $K$ cannot be seen because the simulation time is insufficient. The dependence of the saturated correlation length $\xi_{\text{sat}}(t)$ on $K$ is shown in Fig. 5. It can be fitted by an exponential, $\xi_{\text{sat}}(t) \approx e^{(3.29 \pm 0.03)K}$. This behavior is similar to what was observed for $D_{\text{sat}}$ (cf. Fig. 5).

For times shorter than $t_{\text{cross}}$ we observed the following scaling: $S(r,t) \approx S(r/t^\nu)$, the value of exponent $\nu$ for $K = 2$ is $\nu = 0.465$ (Fig. 11). The function $\tilde{S}(x)$ was found to have the form $\tilde{S}(x) = \exp(-bx-ax^2)$ where the parameters $a,b$ were fitted as $a = 2.6 \pm 0.1$ and $b = 1.86 \pm 0.09$. This behavior agrees with analytical results for the zero-temperature kinetic Ising model.\cite{22} The correlation functions $S(r,t)$ at times larger than $t_{\text{cross}}$ do not depend on time but they depend on $K$. These functions $S_{\text{sat}}(r,K)$ can be also scaled into a universal form using the saturated correlation length $S_{\text{sat}}(r,K)$ $= S(r/t_{\text{cross}}^{\nu}(K))$.

IV. DISCUSSION

The crossover in the surface width as well as in the height-difference correlation function is clearly related to the stopping of phase ordering on the surface. We observed that the time for saturation of a domain size $t_{\text{cross}}(D)$ is approximately proportional to the time $t_{\text{cross}}$ for a crossover to the KPZ
The most striking features in the surface morphology are the pyramids or teeth observed for a sufficiently strong coupling ($K > 1.5$). This can be understood from the growth rules of the model. Let us consider a growth site with all nearest neighbors of the same type. The probability that it will be occupied by a new particle is $e^{-3K} + e^{3K}$. The first term $e^{-3K}$ gives the probability that a new particle will be of the same type as the old ones. This is much larger than the second term corresponding to the probability that a new particle will be of the opposite type. The creation of new domain walls is thus strongly inhibited and growth proceeds preferably by adding the particles of the same type. On the other hand, the probability of occupying a growth site next to the boundary between two domains is $e^{-K} + e^{K}$, i.e., smaller than the growth probability inside the domain. Hence, growth inside the domain is preferable. This leads to the formation of pyramidalike features composed of one type of particles with facets of maximal slope and domain walls in the bottoms of the valleys. In other words, growth is pinned by domain walls. This is in accord with the results on nonhomogeneous growth, where the presence of the inhomogeneity leads to the formation of a dip in the surface.24,25

At the same time, if we observe the deposition of particles next to a domain wall, we can see that the wall movement is due to deposition of a particle of the opposite type, which has probability $e^{-K}$, while no movement has probability $e^{K}$. That is why the wall movement is very slow for large $K$. The surface domains with slowly moving walls result in long vertical lamellae (cf. Fig. 3). This leads to the seemingly surprising fact that the width of lamellae of different types of particles for a given time decreases with the coupling, as we observed for growth with initial conditions fixed by neutral substrate.26 This is also the reason for longer transient times for larger $K$.

All reported exponents were obtained for zero external field. A nonzero external field leads to surface (as well as bulk) magnetization, or in the context of alloy growth, to changing stoichiometry. We can still define exponent $\psi$ for growth of the dominant domain size as well as exponents for kinetic roughening. The effect of this symmetry breaking on the values of exponents remains to be studied.

V. CONCLUSION

We have investigated the interplay between phase ordering and kinetic roughening using the 1+1-dimensional two-component single-step SOS growth model. We examined the validity of scaling for both phenomena and measured the effective scaling exponents.

We observed two situations depending on the strength of coupling $K$ between two types of particles. For a moderate, sufficiently strong ($K > 0.3$) but not very large ($K < 1.8$) coupling, there is a crossover in time and spatial behavior of geometrical characteristics of the surface profile. The effective exponents $\alpha$ and $\beta$ for times shorter than $t_c$ are significantly larger than the KPZ exponents. After a crossover the KPZ exponents were observed. Surface ordering proceeds only up to a finite time $t_{\text{cross}}$, after which it stops. Crossover time $t_{\text{cross}}$ is proportional to $t_{\text{sat}}^D(K)$ and both are exponentially increasing with the coupling strength. For strong coupling ($K = 2$), we observed only enhanced exponents and ordering continued up to the longest simulation time. However, we believe that this difference is only due to the finite time of our simulation, and that ordering will eventually stop and the crossover to the KPZ behavior will occur for any coupling.

The intermediate growth regime is connected with surface phase ordering. It results in enhanced and more rapid kinetic roughening. There is also a crossover in geometrical characteristics with increasing coupling for a fixed time and length scale. Scaling exponents are continuously increasing with $K$. We found that, for sufficiently strong coupling, scaling with enhanced exponents is satisfied over many decades.

During phase ordering, the average size of the surface domains $D$ increases in time as $D \propto t^\psi$ with the exponent $\psi$ close to 1/2. The spin-spin correlation function and the distribution of domain sizes obey scaling with the same exponent. Our results for the surface ordering in the intermediate regime are in agreement with the known results for a one-dimensional kinetic Ising model with the nonconserved order parameter at zero temperature. We expect that the phase ordering on the surface is essentially described by the kinetic Ising model for any coupling. Domain growth stops when the average domain size reaches the equilibrium correlation length. This is reflected in turn by a crossover in effective exponents for kinetic roughening.

Our results lead to conclusion that in 1+1 dimensions there is no new universal behavior and that the TCSS model belongs to the KPZ universality class for any value of coupling. However, since the crossover time and the correlation length are increasing exponentially with coupling, the new behavior in the intermediate regime can be dominant for practically relevant times and length scales.

It is of interest to study the TCSS model in 2+1 dimensions. If the analogy with the kinetic Ising model is valid also here, the size of surface domains will not be restricted and it will diverge for some critical value of $K$. Then a new universal behavior may be observed. Furthermore, one can expect that in 2+1 dimensions a phase transition in kinetic roughening exists. It would be also desirable to explore scaling in different growth models for binary systems, in particular, in models with the surface diffusion.

ACKNOWLEDGMENTS

We thank P. Šmilauer for critical comments. This work was supported by Grant Nos. A1010513 of the GA AV ČR and 202/96/1736 of the GA ČR.
We consider the simplest expression for the energy change leading to phase ordering. The real process of vapor deposition is much more complicated and depends on the local chemical potential, which in turn depends on both the surface morphology and the composition on the surface. The contribution from the surface morphology enters into our simple model only by the single-step constraint. The energy barrier for deposition a particle is finite only for growth sites and infinite for all other sites. The barrier is the same for all growth sites. This could be modified by taking into account interactions not only to nearest neighbors but to next nearest neighbors. The effect of the composition is described by the expression (1).

To be more precise the initial configuration is $h_i^{(0)} = a \cdot (i \mod 2)$ because of the single-step constraint. Here $a$ is the lattice constant. The initial configuration is not strictly flat but it has the intrinsic width $w_{\text{int}} = a/4$.

The additional structure seen for initial times in the case of large couplings is a transient caused by the initial conditions. When we start from a different initial composition of the substrate the dependence of $w$ on $i$ is different for some initial time interval. This interval is increasing with coupling, for $K = 3$ it is about 100 ML (cf. Fig. 4).

The kinetic Ising model evolves towards a stationary state that is an equilibrium state of the thermodynamical Ising model. The domain size $D$ in the Ising model increases as $D \propto e^{2K}$.

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