

# Analysis of Island Dynamics in Epitaxial Growth of Thin Films \*

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## Abstract

This work is concerned with analysis and refinement for a class of island dynamics models for epitaxial growth of crystalline thin films. An island dynamics model consists of evolution equations for step edges (or island boundaries), coupled to a diffusion equation for the adatom density, on an epitaxial surface. The island dynamics model with irreversible aggregation is confirmed to be mathematically ill-posed, with a growth rate that is approximately linear for large wavenumbers. By including a kinetic model for the structure and evolution of step edges, the island dynamics model is made mathematically well-posed. In the limit of small edge Péclet number, the edge kinetics model reduces to a set of boundary conditions, involving line tension and one-dimensional surface diffusion, for the adatom density. Finally, in the infinitely fast terrace diffusion limit, a simplified model of one-dimensional surface diffusion and kink convection is derived and found to be linearly stable.

**Keywords:** epitaxial growth, island dynamics, step edges, normal velocity, adatom diffusion, linear stability, step-edge kinetics, line tension, surface diffusion.

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## 1 Introduction

Epitaxy is the growth of a thin film on a substrate in which the crystal properties of the film are inherited from those of the substrate. Since an epitaxial film can (at least in principle)

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grow as a single crystal without grain boundaries or other defects, this method produces crystals of the highest quality. In spite of its ideal properties, epitaxial growth is still challenging to mathematically model and numerically simulate because of the wide range of length and time scales that it encompasses, from the atomistic to the continuum.

Burton, Cabrera and Frank [2] developed the first detailed theoretical description for epitaxial growth. In this “BCF” model, the adatom density solves a diffusion equation with an equilibrium boundary condition ( $\rho = \rho_{eq}$ ), and step edges (or island boundaries) move at a velocity determined from the diffusive flux to the boundary. Modifications of this theory were made, for example in [7, 8, 10, 11], to include additional effects such as curvature of the boundary, but these are all for near equilibrium. Numerical simulation of island dynamics, using a level set method, was implemented in [4, 5, 9, 12]. A fully nonequilibrium modeling approach was formulated in [3] using an atomistic, kinetic model for the structure and evolution of a step edge (island boundary). All of these are continuum models in that they involve coarse graining in the lateral directions, but they retain atomistic discreteness in the growth direction. They are “island dynamics” models, since they describe an epitaxial surface by the location and evolution of the island boundaries and step edges.

The present article is concerned with the analysis and refinement of island dynamics models. The models are developed here in the context of step-flow growth for an epitaxial surface consisting of a periodic step-train, and their stability and asymptotic properties are analyzed.

The first and simplest island dynamics model involves *irreversible aggregation*. The corresponding boundary condition is that the adatom density vanishes on island boundaries or step edges. This follows from the assumption that an adatom sticks irreversibly to a boundary immediately after it hits the boundary. This model is mathematically equivalent to the BCF model, since  $\rho^{BCF} - \rho_{eq}$  satisfies the equations of this model.

As expected, we find that the island dynamics model with irreversible aggregation is mathematically ill-posed and that a Mullins-Sekerka type dendritic instability occurs in the motion of step edges for large wavenumbers [13, 14]. The instability is weak, since the growth rate in the dispersion relation is asymptotically linear as a function of wavenumber, and it has small coefficient due to two-sided attachment.

The next model is the island dynamics model with *edge kinetics*. It involves not only the island boundaries and the adatom density but also the density of edge-adatoms—atoms that diffuse along island boundaries—and the density of kinks along island boundaries. Both the attachment and detachment of adatoms to and from the boundaries are included in the model. The underlying equations include a diffusion equation for the adatom density together with a mixed type boundary condition, a diffusion equation for the edge-adatom density, and a convection equation for the kink density.

For the linear stability analysis of the island dynamics model with edge kinetics, we only consider the steady and quasi-steady systems. This is justified by the large magnitude of terrace adatom diffusion constant  $D_T$  for most practical applications. In the range of realistic parameters, both systems are linearly stable, with decay rate proportional to  $l^2$ , for

both small and large wavenumbers  $l$ . The constants of proportionality are solely characterized by the “edge Péclet number”  $P_E$ , an input parameter that is proportional to both the constant deposition flux rate and the step width but inversely proportional to the edge diffusion constant  $D_E$ . In fact, for small edge Péclet numbers, both systems are stable for any wavenumbers. Note that the Bales-Zangwill type instability of a step edge [1, 7, 10, 11, 15] is not present due to the exclusion of asymmetry in boundary attachment and the inclusion of edge-adatom diffusion and kink convection in the model.

The edge kinetics model can be directly solved in the limit of small edge Péclet number, to yield new formulas for the normal velocity and for the boundary conditions on the adatom density. These include line tension and one-dimensional surface diffusion (i.e. surface diffusion along the step edge) terms of the usual Gibbs-Thomson type, but with parameters that come from a kinetic steady state rather than from thermodynamic equilibrium. To our knowledge, this is the first derivation of line tension and surface diffusion from a kinetic, atomistic approach. Stability analysis for the resulting system is contained in [7, 10, 11].

Alternatively, in the infinitely fast terrace diffusion limit, the island dynamics model with edge kinetics reduces to a simplified model that involves only the edge-adatom density and kink density on a step edge (but not the adatom density on the terraces). We find that the dispersion relation for the original island dynamics model with edge kinetics is qualitatively recaptured by this simplified model.

In Section 2, the general framework of the island dynamics models is formulated. Section 3 contains the linear stability analysis for the island dynamics model with irreversible aggregation. In Section 4, the island dynamics model with edge kinetics and its linear stability are described. Section 5 describes the limit of small edge Péclet number, including boundary conditions and normal velocity that involve line tension and surface diffusion terms. In Section 6, a simplified model of surface diffusion and kink convection is derived under the assumption of the infinitely fast terrace diffusion, and its linear stability is analyzed. Finally, various details are gathered in four Appendices.

## 2 Island dynamics models

This section provides a general formulation of an island dynamics model applied to step-flow growth of an epitaxial thin film.

Consider a simple cubic crystal with lattice spacing  $a$  and with crystallographic directions parallel to the  $x$ ,  $y$  and  $z$  axes. For notational convenience, nondimensionalize in space for spatial coordinate  $(x, y, z)$ , diffusion coefficient  $D$ , number density per area  $\rho$ , and number density per length  $\phi$  as

$$(x, y, z) \rightarrow (x/a, y/a, z/a) \quad \text{spatial coordinate} \quad (2.1)$$

$$D \rightarrow D/a^2 \quad \text{diffusion coefficient} \quad (2.2)$$

$$\rho \rightarrow a^2 \rho \quad \text{2D density} \quad (2.3)$$

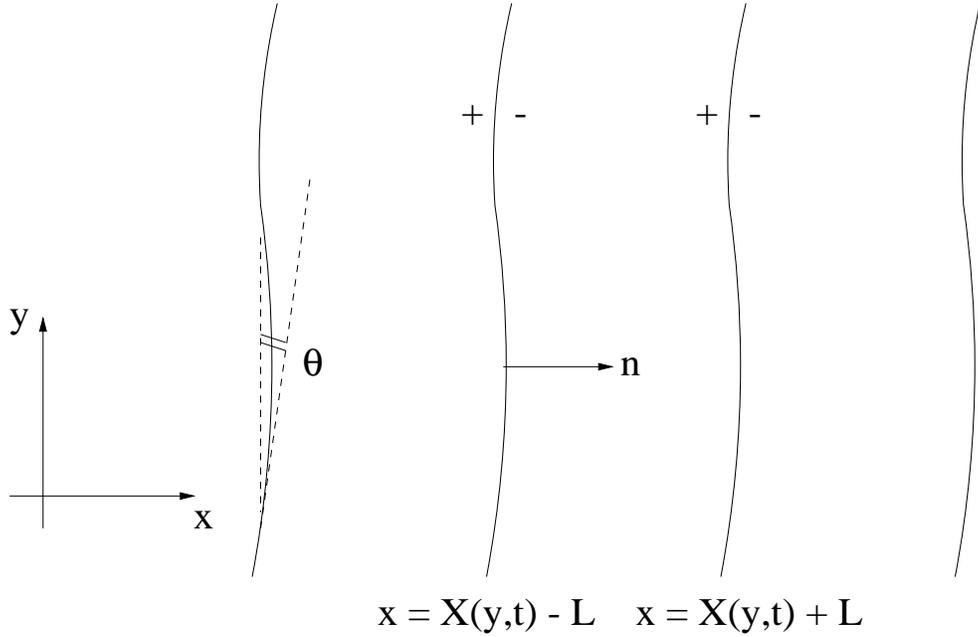


Figure 2.1. The geometry of step-flow growth of a crystal.

$$\phi \rightarrow a\phi \qquad \text{1D density.} \qquad (2.4)$$

This is equivalent to measuring all distances in units of  $a$  or to setting  $a = 1$ .

Assume that the crystal surface consists of a periodic sequence of step edges that are approximately straight and parallel to the  $y$  axis, as in Figure 2.1. Each step is one atomic layer lower than the preceding one to its left, and the steps move to the right during crystal growth. These step edges are represented as

$$x = X(y, t) + (2j + 1)L, \quad j = 0, \pm 1, \dots,$$

in which  $X(y, t)$  is a smooth function and  $L > 0$  is half of the step width. For each point  $(X(y, t), y)$  on a step edge, let  $\theta = \theta(y, t)$  denote the signed angle between the tangent of the step edge and the  $y$  axis; i.e.,

$$\tan \theta = -\partial_y X, \quad -\pi/2 < \theta < \pi/2. \qquad (2.5)$$

The curvature  $\kappa = \kappa(y, t)$ , the unit normal  $\mathbf{n} = \mathbf{n}(y, t)$  pointing from the upper terrace into lower terrace, and the normal velocity  $v = v(y, t)$  of the step edge are given by

$$\kappa = \partial_s \theta, \quad \mathbf{n} = (\cos \theta, \sin \theta), \quad v = (\partial_t X) \cos \theta, \qquad (2.6)$$

in which  $\partial_s$  is the tangential derivative in the  $y$  direction. Note that the sign of  $\theta$  has been changed from that used in [3] to avoid a minus sign in the definition of curvature  $\kappa$ . From

(2.5) and (2.6), the formulas for  $\mathbf{n}$  and  $v$  are

$$\mathbf{n} = \frac{1}{\sqrt{1 + (\partial_y X)^2}}(1, -\partial_y X) \quad \text{and} \quad v = \frac{\partial_t X}{\sqrt{1 + (\partial_y X)^2}}. \quad (2.7)$$

Epitaxial growth involves deposition, diffusion and attachment of adatoms on the surface. Deposition is from an external source, such as a molecular beam. The adatoms on the surface can be described by an adatom density function  $\rho = \rho(x, y, t)$ , which is assumed to be periodic in  $x$  with period  $2L$ . Adatom diffusion on the epitaxial surface is described by a diffusion equation of the form

$$\partial_t \rho - D_T \nabla^2 \rho = F \quad \text{for } X(y, t) - L < x < X(y, t) + L, \quad (2.8)$$

in which  $D_T$  and  $F$  are the terrace diffusion constant and the deposition flux rate, respectively, after spatial nondimensionalization as in Eqs. (2.1)–(2.4). For simplicity, we have left out desorption and nucleation terms on the right-hand side of the equation (2.8). Attachment of adatoms to the step edges and the resulting motion of the step edges are described by boundary conditions at the step edges for the diffusion equation and a formula for the step-edge velocity  $v$ .

Denote the restriction of any function  $w$  to the step edge from upper and lower terraces as  $w_+$  and  $w_-$ , respectively, as indicated in Figure 2.1. The net flux to the step edge from upper and lower terraces is denoted as  $f_+ = f_+(y, t)$  and  $f_- = f_-(y, t)$ , respectively, in which

$$v \rho_+ + D_T \mathbf{n} \cdot \nabla \rho_+ = -f_+ \quad \text{at } x = X(y, t) + L, \quad (2.9)$$

$$v \rho_- + D_T \mathbf{n} \cdot \nabla \rho_- = f_- \quad \text{at } x = X(y, t) - L. \quad (2.10)$$

The total flux is  $f = f_+ + f_-$  given by

$$f = -v[\rho] - D_T[\mathbf{n} \cdot \nabla \rho],$$

in which the square bracket  $[\ ]$  is defined as the difference across the step; i.e., for any  $w$

$$[w] = w_+ - w_-$$

The principal dimensionless parameters for epitaxial growth are the ratio of flux and diffusive, which we refer to as ‘‘Péclet numbers’’ by analogy with fluid mechanics. In steady state the flux to an edge and the edge velocity are  $f = 2FL$  and  $v = f$ . Let  $P_T$  be the terrace Péclet number and  $P_E$  be the edge Péclet number, defined as

$$P_T = v_0/D_T = 2FL/D_T, \quad (2.11)$$

$$P_E = 2FL/D_E, \quad (2.12)$$

in which  $D_E$  is the edge diffusion constant. An additional frequently used dimensionless parameter is

$$R = F/D_T = P_T/2L.$$

Different island dynamics models are distinguished by having different diffusive boundary conditions and different normal velocity. The following is a summary of the four island dynamics models that are analyzed in this paper, of which the fourth does not involve an adatom density:

1. *The island dynamics model with irreversible aggregation (Section 3):*

$$\begin{aligned}\rho &= 0 & \text{at } x = X(y, t) \pm L, \\ v &= f.\end{aligned}$$

2. *The island dynamics model with step-edge kinetics (Section 4):*

$$\begin{aligned}f_+ &= D_T \rho_+ - D_E \phi & \text{at } x = X(y, t) + L, \\ f_- &= D_T \rho_- - D_E \phi & \text{at } x = X(y, t) - L, \\ v &= k w \cos \theta,\end{aligned}$$

in which  $f_{\pm}$  are defined by (2.9) and (2.10),  $D_E$  is the (spatially nondimensionalized) diffusion coefficient for both edge diffusion and edge detachment,  $\phi$  and  $k$  are the densities of edge-atoms and kinks, and  $w$  is the kink velocity, defined in Section 4. The velocity law given in the last equation above was first derived in [8].

3. *The island dynamics model with line tension and surface diffusion (Section 5):*

$$\begin{aligned}f_+ &= D_T(\rho_+ - \rho_*) - \gamma\kappa, & \text{at } x = X(y, t) + L, \\ f_- &= D_T(\rho_- - \rho_*) - \gamma\kappa, & \text{at } x = X(y, t) - L, \\ v &= D_T \mathbf{n} \cdot [\nabla \rho] + \beta \rho_{*yy} + (\gamma/D_E) \kappa_{yy},\end{aligned}$$

in which  $\kappa$  is curvature and  $\rho_*$ ,  $\beta$  and  $\gamma$  are defined in Section 5.

4. *The infinitely fast terrace diffusion limit (Section 6):*

$$v = l_1 D_E \phi k \cos \theta,$$

in which  $l_1$  is a constant defined in Section 6.

### 3 Island dynamics with irreversible aggregation

The island dynamics model with irreversible aggregation for the periodic step-train problem consists of the following diffusion equation, boundary conditions and velocity law:

$$\partial_t \rho - D_T \nabla^2 \rho = F \quad \text{for } X(y, t) - L < x < X(y, t) + L, \quad (3.1)$$

$$\rho = 0 \quad \text{at } x = X(y, t) \pm L, \quad (3.2)$$

$$v = -D_T \mathbf{n} \cdot [\nabla \rho], \quad (3.3)$$

in which the adatom density  $\rho = \rho(x, y, t)$  is assumed to be periodic in  $x$  with period  $2L$ , and  $v$  is the normal velocity defined in (2.6).

A steady planar solution of (3.1)–(3.3) is given by

$$X_0(y, t) = v_0 t, \quad (3.4)$$

$$v_0 = 2FL, \quad (3.5)$$

$$\rho_0 = b_0 + b_1 \tilde{x} + b_2 e^{-P_T \tilde{x}}, \quad (3.6)$$

$$\tilde{x} = x - v_0 t, \quad (3.7)$$

in which

$$b_0 = (1/2) \coth(P_T L), \quad b_1 = -(2L)^{-1}, \quad b_2 = -(2 \sinh(P_T L))^{-1}.$$

Next we perform linear analysis for perturbations of this planar solution. Set

$$X = X_0 + \varepsilon X_1, \quad \rho = \rho_0 + \varepsilon \rho_1, \quad \mathbf{n} = \mathbf{n}_0 + \varepsilon \mathbf{n}_1, \quad v = v_0 + \varepsilon v_1,$$

where  $\varepsilon$  is a parameter small in magnitude. By (2.7), the unit normal is  $\mathbf{n}_1 = (0, -\partial_y X_1)$  and the normal velocity is  $v_1 = \partial_t X_1$ . The linearized equations for  $\rho_1 = \rho_1(\tilde{x}, y, t)$  are

$$\partial_t \rho_1 - v_0 \partial_{\tilde{x}} \rho_1 - D_T \nabla^2 \rho_1 = 0 \quad \text{for } -L < \tilde{x} < L, \quad (3.8)$$

$$\rho_1 + X_1 \rho'_0 = 0 \quad \text{at } \tilde{x} = \pm L, \quad (3.9)$$

$$\partial_t X_1 = -D_T [\partial_{\tilde{x}} \rho_1] - D_T X_1 [\rho''_0], \quad (3.10)$$

where the jumps are defined using the limiting values at  $\pm L$ . These equations are derived using perturbation theory for a free boundary, as described in Appendix A.

Principal mode solutions  $(X_1, \rho_1)$  have the form

$$X_1 = \hat{X}_1 e^{\omega t + i l y} \quad \text{and} \quad \rho_1 = \left( \hat{\rho}_+ e^{\hat{D}_T + \tilde{x}} + \hat{\rho}_- e^{\hat{D}_T - \tilde{x}} \right) e^{\omega t + i l y},$$

where  $l$  is the wavenumber and  $\hat{X}$ ,  $\hat{\rho}_+$ ,  $\hat{\rho}_-$ ,  $\hat{D}_{T+}$ ,  $\hat{D}_{T-}$  and  $\omega$  are all constants. For these solutions, (3.9) and (3.10) are equivalent to

$$\hat{\rho}_+ e^{\hat{D}_T + L} + \hat{\rho}_- e^{\hat{D}_T - L} + \hat{X}_1 \rho'_0(L) = 0,$$

$$\hat{\rho}_+ e^{-\hat{D}_T + L} + \hat{\rho}_- e^{-\hat{D}_T - L} + \hat{X}_1 \rho'_0(-L) = 0,$$

$$\omega \hat{X}_1 = -2D_T \left( \hat{D}_{T+} \hat{\rho}_+ \sinh(\hat{D}_{T+} L) + \hat{D}_{T-} \hat{\rho}_- \sinh(\hat{D}_{T-} L) \right) - D_T \hat{X}_1 (\rho''_0(L) - \rho''_0(-L)),$$

which imply

$$\begin{vmatrix} e^{\hat{D}_T + L} & e^{\hat{D}_T - L} & \rho'_0(L) \\ e^{-\hat{D}_T + L} & e^{-\hat{D}_T - L} & \rho'_0(-L) \\ -2D_T \hat{D}_{T+} \sinh(\hat{D}_{T+} L) & -2D_T \hat{D}_{T-} \sinh(\hat{D}_{T-} L) & -D_T (\rho''_0(L) - \rho''_0(-L)) + \omega \end{vmatrix} = 0.$$

This is equivalent to

$$\left\{ \coth(\hat{D}_{T+}L) - \coth(\hat{D}_{T-}L) \right\} \left\{ D_T (\rho_0''(L) - \rho_0''(-L)) + \omega \right\} + D_T \left( \hat{D}_{T-} - \hat{D}_{T+} \right) \cdot (\rho_0'(L) + \rho_0'(-L)) + D_T (\rho_0'(L) - \rho_0'(-L)) \left\{ \hat{D}_{T+} \coth(\hat{D}_{T-}L) - \hat{D}_{T-} \coth(\hat{D}_{T+}L) \right\} = 0. \quad (3.11)$$

Similarly, insertion of  $\rho_1$  into (3.8) leads to

$$D_T \hat{D}_{T+}^2 + v_0 \hat{D}_{T+} - (\omega + D_T l^2) = 0, \quad (3.12)$$

$$D_T \hat{D}_{T-}^2 + v_0 \hat{D}_{T-} - (\omega + D_T l^2) = 0. \quad (3.13)$$

The three equations (3.11)–(3.13) determine  $\hat{D}_{T+} = \hat{D}_{T+}(l)$ ,  $\hat{D}_{T-} = \hat{D}_{T-}(l)$ , and the growth rate  $\omega = \omega(l)$ .

**Proposition 3.1** *For the island dynamics with irreversible aggregation, the growth rate  $\omega = \omega(l)$  satisfies*

$$\omega = \omega_0 l + o(l) \quad \text{as } l \rightarrow \infty, \quad (3.14)$$

where

$$\omega_0 = D_T (\rho_0'(L) + \rho_0'(-L)) > 0.$$

**Proof** Observe from (3.11) that  $\omega = \omega(l)$  is of order not greater than  $O(\hat{D}_{T\pm}(l))$ . But by (3.12) and (3.13),  $\hat{D}_{T+}$  and  $\hat{D}_{T-}$  are the two roots of the same quadratic equation. Hence,

$$\hat{D}_{T+} + \hat{D}_{T-} = -\frac{v_0}{D_T} \quad \text{and} \quad \hat{D}_{T+} \hat{D}_{T-} = -\frac{\omega + D_T l^2}{D_T}.$$

Consequently, both  $\hat{D}_{T+}$  and  $\hat{D}_{T-}$  have the leading order terms  $l$  and  $-l$ , respectively. By examining the leading order term in (3.11), we obtain the desired asymptotic expansion (3.14) by a series of calculations. Finally, a direct verification using (3.4), (3.5) and (2.11) leads to

$$\rho_0'(L) + \rho_0'(-L) = \frac{P_T L \cosh(P_T L) - \sinh(P_T L)}{L \sinh(P_T L)} > 0.$$

The proof is complete. **Q.E.D.**

Our analysis shows that the irreversible aggregation model of island dynamics is mathematically ill-posed. However, the instability of the step-edge motion predicted by the model is weak, since the growth rate  $\omega$  depends on the wavenumber  $l$  linearly up to the leading order, and since the coefficient,  $\omega_0$ , of the leading order term can be shown to be  $v_0 (P_T L v_0 / 3 + O((P_T L)^3))$  in the small terrace Péclet number asymptotic expansion. In fact, for the quasi-steady system—the system obtained by dropping the term  $\partial_t \rho$  in (3.1)–(3.3), one can show easily that the growth rate is  $\omega = 0$ . In Section 5, we propose more physically acceptable boundary conditions derived from the step-edge kinetics to replace the irreversibility condition in this model.

## 4 Island dynamics with the kinetic edge model

### 4.1 The kinetic edge model

The kinetic edge model of island dynamics was first developed in [3]. It involves a statistical description of the crystalline structure of a step edge, including the edge-atom density  $\phi = \phi(y, t)$  and the kink density  $k = k(y, t)$ . Edge-atoms are atoms with a single in-plane neighbor along the step; kinks are atoms with two in-plane neighbors. Kinks are of two types—right-facing kinks and left-facing kinks—the densities of which are denoted by  $k_r$  and  $k_\ell$ . The total kink density and the relation between the kink density and the normal angle are

$$k_r + k_\ell = k, \quad (4.1)$$

$$k_r - k_\ell = \tan \theta, \quad (4.2)$$

as determined by [2].

The kinetic edge model consists of a diffusion equation for the edge adatom density  $\phi$  and a convection equation for the kink density  $k$

$$\partial_t \phi - D_E \partial_s^2 \phi = f_+ + f_- - f_0, \quad (4.3)$$

$$\partial_t k + \partial_s (w(k_r - k_\ell)) = 2(g - h). \quad (4.4)$$

In Eq. (4.3),  $D_E$  is the edge diffusion coefficient,  $f_\pm$  are the net fluxes to the edge from terraces as defined in Eq. (2.9) and Eq. (2.10), and  $f_0$  is the net loss term due to the attachment of edge-adatoms to kinks. In Eq. (4.4),  $w$  is the kink velocity, and  $g$  and  $h$  represent, respectively, the creation and annihilation of left-right kink pairs. Note that left-facing kinks and right-facing kinks move in opposite directions with velocity  $w$  and  $-w$ , respectively.

The quantities  $f_+$ ,  $f_-$ ,  $f_0$ ,  $w$ ,  $g$ ,  $h$ , and  $v$  are determined by the following constitutive relations

$$f_+ = D_T \rho_+ - D_E \phi, \quad (4.5)$$

$$f_- = D_T \rho_- - D_E \phi, \quad (4.6)$$

$$f_0 = v(\phi \kappa + 1) = v(1 - \phi X_{yy}), \quad (4.7)$$

$$w = l_1 D_E \phi + D_T(l_2 \rho_+ + l_3 \rho_-) = l_{123} D_E \phi + l_2 f_+ + l_3 f_-, \quad (4.8)$$

$$g = \phi(m_1 D_E \phi + D_T(m_2 \rho_+ + m_3 \rho_-)) = \phi(m_{123} D_E \phi + m_2 f_+ + m_3 f_-), \quad (4.9)$$

$$h = k_r k_\ell (n_1 D_E \phi + D_T(n_2 \rho_+ + n_3 \rho_-)) = k_r k_\ell (n_{123} D_E \phi + n_2 f_+ + n_3 f_-), \quad (4.10)$$

$$X_t = v = w k \cos \theta, \quad (4.11)$$

where  $D_T$  is the (diffusion) hopping rate of an adatom on a terrace,  $D_E$  is the (diffusion) hopping rate of an edge-adatom along or off an edge, and all  $l_i, m_i, n_i$  ( $i = 1, 2, 3$ ) are nonnegative parameters. For convenience we have used the notation

$$q_{ij} = q_i + q_j \quad \text{and} \quad q_{ijk} = q_i + q_j + q_k$$

for  $q = l, m$ , or  $n$ . The constitutive laws (4.5)–(4.10) have been simplified by omission of terms that are insignificant for the kinetic steady state solutions of relevance to step-flow growth. The neglected terms are physically important, however, since they are necessary for detailed balance and for equilibrium solutions; they are included in the more complete analysis of [3].

The relations (4.5), (4.6), and (4.8)–(4.10) are determined by a mean field theory [3]. Eq. (4.7) is derived from the conservation of mass [3], and Eq. (4.11) comes as in [8] from counting adatoms as part of the crystal once they have attached to a kink. Note that for simplicity, a number of physical effects have been neglected. They include the desorption of adatoms into vapor, the nucleation of islands on steps, and the Ehrlich-Schwoebel effect [6, 16, 17]. In [3] the parameters  $l_i$ ,  $m_i$ , and  $n_i$  are fixed to be  $(l_1, l_2, l_3) = (2, 2, 1)$ ,  $(m_1, m_2, m_3) = (2, 4, 2)$ , and  $(n_1, n_2, n_3) = (2, 3, 1)$ , which follow from a simple geometric counting argument, but in the present work they are allowed to be arbitrary parameters.

## 4.2 Quasi-steady island dynamics with the kinetic edge model

For simplicity in the subsequent analysis, consider the “quasi-steady” island dynamics model consisting of the kinetic edge model (4.1)–(4.11), combined with quasi-steady adatom diffusion equation

$$-D_T \nabla^2 \rho = F \quad \text{for } X(y, t) - L < x < X(y, t) + L, \quad (4.12)$$

$$D_T \mathbf{n} \cdot \nabla \rho_+ = -f_+ \quad \text{at } x = X(y, t) + L, \quad (4.13)$$

$$D_T \mathbf{n} \cdot \nabla \rho_- = f_- \quad \text{at } x = X(y, t) - L. \quad (4.14)$$

We also consider the “steady” island dynamics model that is obtained by replacing Eqs. (4.3), (4.4) in the quasi-steady system by

$$-D_E \partial_s^2 \phi = f_+ + f_- - f_0, \quad (4.15)$$

$$\partial_s(w(k_r - k_\ell)) = 2(g - h). \quad (4.16)$$

In the steady system, dynamics is retained only in the equation for the step-edge position  $X$ .

## 4.3 Planar steady-state solution

Let  $X_0(y, t) = v_0 t$  define the step edge position, where  $v_0 > 0$  is a constant. The corresponding angle, curvature, and normal are  $\theta_0 = 0$ ,  $\kappa_0 = 0$ , and  $\mathbf{n}_0 = (0, 1)$ , respectively. Assuming that the adatom density  $\rho_0$  is independent of  $y$  and that both the edge-adatom density  $\phi_0$  and the kink density  $k_0$  are constants, we obtain then from (4.3)–(4.14) the following steady-state solution

$$X_0(y, t) = v_0 t, \quad (4.17)$$

$$v_0 = 2FL, \quad (4.18)$$

$$\rho_0(x, t) = -\frac{F}{2D_T} ((x - v_0t)^2 - L^2) + \frac{D_E\phi_0 + FL}{D_T} \quad \text{for } v_0t - L < x < v_0t + L, \quad (4.19)$$

$$f_{00} = 2f_{+0} = 2f_{-0} = 2FL, \quad (4.20)$$

$$k_0 = \frac{2FL}{w_0}, \quad (4.21)$$

$$w_0 = l_{123}D_E\phi_0 + l_{23}FL, \quad (4.22)$$

$$g_0 = \phi_0(m_{123}D_E\phi_0 + m_{23}FL), \quad (4.23)$$

$$h_0 = \frac{1}{4}k_0^2(n_{123}D_E\phi_0 + n_{23}FL), \quad (4.24)$$

$$g_0 = h_0. \quad (4.25)$$

Note that the last equation determines  $\phi_0$ .

In [3], the solution of these equations was found to scale with  $\varepsilon$  defined as

$$\varepsilon = P_E^{1/3} \quad (4.26)$$

in which  $P_E$  is the edge Péclet number defined in (2.12). This is described more precisely in the following Proposition, the proof of which is presented in Appendix B:

**Proposition 4.1** *Suppose  $l_{123} \neq 0$  and  $m_{123} \neq 0$ . Then,*

$$\phi_0 = \left( \frac{n_{123}}{4l_{123}^2 m_{123}} \right)^{1/3} \varepsilon^2 + O(\varepsilon^3) \quad \text{as } \varepsilon \rightarrow 0, \quad (4.27)$$

$$k_0 = \left( \frac{l_{123} n_{123}}{4m_{123}} \right)^{1/3} \varepsilon + O(\varepsilon^2) \quad \text{as } \varepsilon \rightarrow 0. \quad (4.28)$$

Suppose  $l_{23} \neq 0$ ,  $m_{23} \neq 0$  and  $n_{23} \neq 0$ . Let

$$\sigma_0 = \frac{n_{23}}{l_{23}^2 m_{23}} \quad \text{and} \quad \sigma_1 = 2\sigma_0^2 \left( \frac{n_1}{n_{23}} - \frac{m_1}{m_{23}} - \frac{2l_1}{l_{23}} - 2 \right). \quad (4.29)$$

Then,

$$\phi_0 \rightarrow \sigma_0 \quad \text{and} \quad k_0 \rightarrow 2/l_{23} \quad \text{as } \varepsilon \rightarrow \infty. \quad (4.30)$$

Moreover, if  $\sigma_1 < 0$ , then  $\phi_0$  is a strictly increasing function of  $\varepsilon$  in  $(0, \infty)$ ; if  $\sigma_1 > 0$ , then there exists a value  $\varepsilon_0 > 0$  such that  $\phi_0$  is a strictly increasing function of  $\varepsilon$  for  $0 < \varepsilon < \varepsilon_0$ .

We remark that the case that  $l_{23} = m_{23} = n_{23} = 0$  is included in our simplified model in Section 6 below, cf. (6.4). In Figure 3.1, we plot the graph of  $\phi_0$  and  $k_0$  as functions of  $\varepsilon = P_E^{1/3}$  with  $(l_1, l_2, l_3) = (2, 2, 1)$ ,  $(m_1, m_2, m_3) = (2, 4, 2)$ , and  $(n_1, n_2, n_3) = (2, 3, 1)$ , which are the parameters used in [3] and satisfy  $\sigma_1 < 0$ .

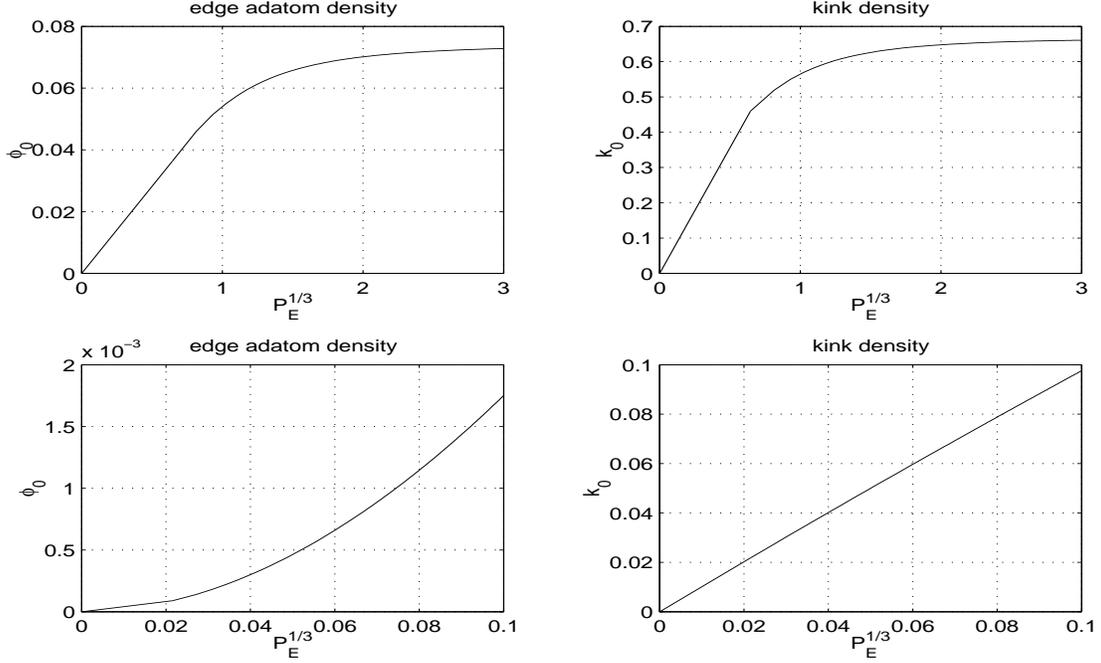


Figure 3.1. The steady-state edge-adatom density and kink density as functions of  $\varepsilon = P_E^{1/3}$  with  $(l_1, l_2, l_3) = (2, 2, 1)$ ,  $(m_1, m_2, m_3) = (2, 4, 2)$ , and  $(n_1, n_2, n_3) = (2, 3, 1)$ . The corresponding top and bottom plots differ only by scales.

#### 4.4 Linear stability analysis for island dynamics with the kinetic edge model

Consider a perturbation around the steady planar solution, defined as

$$\begin{aligned}
& (X, \rho, \phi, k, \theta, \kappa, \mathbf{n}, v, f_{\pm}, f_0, w, g, h) \\
&= (X_0, \rho_0, \phi_0, k_0, \theta_0, \kappa_0, \mathbf{n}_0, v_0, f_{\pm 0}, f_{00}, w_0, g_0, h_0) \\
&+ \varepsilon (X_1, \rho_1, \phi_1, k_1, \theta_1, \kappa_1, \mathbf{n}_1, v_1, f_{\pm 1}, f_{01}, w_1, g_1, h_1),
\end{aligned}$$

where  $\varepsilon$  is a parameter small in magnitude. Equations (2.5), (2.6), (4.1) and (4.2) imply

$$\theta_1 = -\partial_y X_1, \quad \kappa_1 = \partial_s \theta_1, \quad \mathbf{n}_1 = (0, \theta_1), \quad v_1 = \partial_t X_1, \quad (4.31)$$

$$k_{\ell} k_r = \frac{1}{4} ((k_{\ell} + k_r)^2 - (k_{\ell} - k_r)^2) = \frac{1}{4} (k^2 - \tan^2 \theta) = \frac{1}{4} k^2 + O(\varepsilon^2). \quad (4.32)$$

Now, inserting all the above expansions into Eqs. (4.3)–(4.14), and using (4.17)–(4.25), (4.31) and (4.32), we obtain the following linearized system

$$\nabla^2 \rho_1 = 0 \quad \text{for } v_0 t - L < x < v_0 t + L, \quad (4.33)$$

$$D_T \partial_x \rho_{1+} - F X_1 + f_{+1} = 0 \quad \text{at } x = v_0 t + L, \quad (4.34)$$

$$D_T \partial_x \rho_{1-} - F X_1 - f_{-1} = 0 \quad \text{at } x = v_0 t - L, \quad (4.35)$$

$$\partial_t \phi_1 - D_E \partial_y^2 \phi_1 = f_{+1} + f_{-1} - f_{01}, \quad (4.36)$$

$$\partial_t k_1 - w_0 \partial_y^2 X_1 = 2(g_1 - h_1), \quad (4.37)$$

with the constitutive relations

$$f_{+1} = D_T \rho_{1+} - D_E \phi_1 - F L X_1, \quad (4.38)$$

$$f_{-1} = D_T \rho_{1-} - D_E \phi_1 + F L X_1, \quad (4.39)$$

$$f_{01} = -v_0 \phi_0 \partial_y^2 X_1 + \partial_t X_1, \quad (4.40)$$

$$w_1 = D_T (l_2 \rho_{1+} + l_3 \rho_{1-}) + l_1 D_E \phi_1 + (l_3 - l_2) F L X_1, \quad (4.41)$$

$$g_1 = D_T \phi_0 (m_2 \rho_{1+} + m_3 \rho_{1-}) + (m_1 D_E \phi_0 + g_0 / \phi_0) \phi_1 + (m_3 - m_2) F L \phi_0 X_1, \quad (4.42)$$

$$h_1 = \frac{1}{4} D_T k_0^2 (n_2 \rho_{1+} + n_3 \rho_{1-}) + \frac{1}{4} n_1 D_E k_0^2 \phi_1 + (2h_0 / k_0) k_1 + \frac{1}{4} (n_3 - n_2) F L k_0^2 X_1, \quad (4.43)$$

$$\partial_t X_1 = w_0 k_1 + k_0 w_1, \quad (4.44)$$

where  $\rho_{1+}$  and  $\rho_{1-}$  denote the restriction of  $\rho_1$  at  $x = v_0 t + L$  and  $x = v_0 t - L$ , respectively.

The linearized system for the steady system (4.5)–(4.16), is the same except that the two time derivative terms  $\partial_t \phi_1$  and  $\partial_t k_1$  are dropped from (4.36) and (4.37), respectively.

**Dispersion relation.** We seek solutions to the linearized system (4.33)–(4.44) of the form

$$\rho_1 = (\hat{r}_1 \cosh(lx) + \hat{s}_1 \sinh(lx)) e^{\omega t + i l y}, \quad (4.45)$$

$$(X_1, \phi_1, k_1) = (\hat{X}_1, \hat{\phi}_1, \hat{k}_1) e^{\omega t + i l y}, \quad (4.46)$$

where  $l$  is the wavenumber and  $\hat{r}_1$ ,  $\hat{s}_1$ ,  $\hat{X}_1$ ,  $\hat{\phi}_1$  and  $\hat{k}_1$  are constants. The dispersion relation for this solution is analyzed in the Appendix C. In particular, its asymptotic behavior for large and small wavenumbers is described as follows:

**Asymptotic analysis.** Denote for each  $l \geq 0$  by  $\omega = \omega(l)$  the root of the linear equation (C.2) and by  $\omega_1 = \omega_1(l)$ ,  $\omega_2 = \omega_2(l)$ , and  $\omega_3 = \omega_3(l)$  the three roots of the cubic equation (C.1). Assume that  $\omega_1(l)$  is always real and that  $\omega_1(0) = 0$ .

**Proposition 4.2** *For the steady system, we have*

$$\omega(l) = -v_0 \phi_0 l^2 + O(l^4) \quad \text{as } l \rightarrow 0, \quad (4.47)$$

$$\omega(l) = -\frac{v_0 w_0}{4 h_0} l^2 + O(1) \quad \text{as } l \rightarrow \infty. \quad (4.48)$$

*For the quasi-steady system, we have*

$$\omega_1(l) = -v_0 \phi_0 l^2 + O(l^4) \quad \text{as } l \rightarrow 0, \quad (4.49)$$

and

$$\begin{aligned}
\omega_1(l) &= -D_E l^2 + O(l), \\
\omega_2(l) &= R_0 + iw_0 l + O(l^{-1}), \\
\omega_3(l) &= R_0 - iw_0 l + O(l^{-1}),
\end{aligned} \tag{4.50}$$

as  $l \rightarrow \infty$ , where  $i = \sqrt{-1}$  and

$$R_0 = \frac{1}{4} D_E k_0 ((l_3 - l_2 - n_{23}) P_E - 2n_{123} \phi_0 - 2l_1 P_E \phi_0). \tag{4.51}$$

Moreover,  $R_0 < 0$  for small edge Péclet number  $P_E$ . A sufficient condition for  $R_0 < 0$  for all  $P_E > 0$  is that

$$l_3 - l_2 - n_{23} \leq 0, \tag{4.52}$$

and a necessary condition for  $R_0 < 0$  for all  $P_E > 0$  is that

$$(l_3 - l_2 - n_{23}) l_{23}^2 m_{23} - 2l_1 n_{23} \leq 0. \tag{4.53}$$

The proof of these results is given in Appendix D.

## 5 Kinetic derivation of line tension and one-dimensional surface diffusion

As shown below, the kinetic edge model in the limit of small edge Péclet number yields a theory that is nearly as simple as the irreversible aggregation model. The asymptotic assumptions that underline this derivation are mathematically rather than physically based; i.e., they come from the “distinguished limit” rather than from a characterization of the size of the external sources that drive the problem.

### 5.1 Asymptotic Expansion for Small Péclet Number

Consider the limiting adatom densities  $\rho_{\pm}$ , the fluxes  $f_{\pm}$  and the curvature  $\kappa = -X_{yy}$  to be parameters that are determined away from the boundary. Then, the main equations are (4.3), (4.4) and (4.11) for  $\phi$ ,  $k$  and  $v$ , respectively. In the scaling detailed below, the angle  $\theta$  is of size  $\varepsilon^{3/2}$ , so that  $\cos \theta \approx 1$ ; for simplicity, this approximation will be used from the beginning. Use the relations (4.5)–(4.10) to eliminate  $\rho_{\pm}$ ,  $f_0$ ,  $w$ ,  $g$ , and  $h$ , and write the resulting equations as

$$\begin{aligned}
f_+ + f_- - v &= v\phi\kappa + \phi_t - D_E \phi_{yy}, \\
2m_{123} D_E \phi^2 - \frac{1}{2} n_{123} D_E k^2 \phi &= -2\phi(m_2 f_+ + m_3 f_-) + \frac{1}{2} k^2 (n_2 f_+ + n_3 f_-)
\end{aligned} \tag{5.1}$$

$$+ k_t - ((v/k)X_y)_y, \quad (5.2)$$

$$v - l_{123}D_E\phi k = k(l_2f_+ + l_3f_-). \quad (5.3)$$

From the steady state solution of Section 3, the governing parameters in this system are  $\varepsilon = P_E^{1/3}$ , as defined in (4.26), and  $\bar{f} = 2LF$ . The steady state solution from Section 4 shows that

$$(k, \phi, v, f_{\pm}) = (\varepsilon k', \varepsilon^2 \phi', \bar{f} v', \bar{f} f'_{\pm}).$$

As shown below, the distinguished limit is

$$\begin{aligned} \kappa &= \varepsilon^2 \kappa', \\ Y &= \varepsilon^{-1/2} Y'. \end{aligned}$$

Since  $\kappa = -X_{yy}$ , this implies that  $X = \varepsilon X'$  and also that  $\theta = O(\varepsilon^{3/2})$  as stated earlier. In addition, the dispersion relation from Proposition 3.2 shows that the time scale is  $T = \omega^{-1} = (-v\phi\ell^2)^{-1} = (-\bar{f}v'\varepsilon^2\phi'Y'^{-2})^{-1} = O(\bar{f}^{-1}\varepsilon^{-3})$ ; i.e.,

$$T = \bar{f}^{-1}\varepsilon^{-3}T'.$$

Dividing (5.1) and (5.3) by  $\bar{f}$ , and dividing (5.2) by  $D_E\varepsilon^4 = \bar{f}\varepsilon$ , we obtain

$$\begin{aligned} f'_+ + f'_- - v' &= \varepsilon^4 v' \phi' \kappa' + \varepsilon^5 \phi'_v - \phi'_{y'y'}, \\ 2m_{123}\phi'^2 - \frac{1}{2}n_{123}k'^2\phi' &= -2\varepsilon\phi'(m_2f'_+ + m_3f'_-) + \varepsilon\frac{1}{2}k'^2(n_2f'_+ + n_3f'_-) \\ &\quad + \varepsilon^3 k'_v - ((v/k)X'_{y'})_{y'}, \\ v' - l_{123}\phi'k' &= \varepsilon k'(l_2f'_+ + l_3f'_-). \end{aligned}$$

Notice that  $\cos\theta = (1 + \tan^2\theta)^{-1/2} = (1 + X_y^2)^{-1/2} = 1 + O(\varepsilon^3)$ . It follows that, to leading order (i.e., ignoring terms of size  $O(\varepsilon)$ ),  $v'$ ,  $k'$  and  $\phi'$  are given by

$$v' = f'_+ + f'_- + \phi'_{y'y'}, \quad (5.4)$$

$$k' = (l_{123}\phi')^{-1}v', \quad (5.5)$$

$$2m_{123}\phi'^2 - \frac{n_{123}}{2l_{123}^2}\phi'^{-1}v'^2 = -((l_{123}\phi')X'_{y'})_{y'}. \quad (5.6)$$

The result is more transparent if  $X' = O(\delta)$  and  $Y' = O(1/\delta)$  (i.e.,  $X = O(\delta\varepsilon)$  and  $Y = O(\delta^{-1}\varepsilon^{-1/2})$ ), in which  $\varepsilon \ll \delta \ll 1$ , so that the previous equations (5.4)–(5.6) can be linearized, but additional terms of size  $O(\varepsilon)$  from above are not required. Then  $\phi'$  and  $v'$  are given by

$$\phi' = \left( \frac{n_{123}}{4m_{123}l_{123}^2} \right)^{1/3} (f'_+ + f'_-)^{2/3} - \frac{l_{123}}{6m_{123}} X'_{y'y'} \quad (5.7)$$

$$v' = f'_+ + f'_- + \phi'_{y'y'}. \quad (5.8)$$

## 5.2 Boundary conditions and velocity with line tension and one-dimensional surface diffusion

The boundary conditions for  $\rho$  from the kinetic edge model are (cf. (4.13)–(4.14), (4.5), and (4.6))

$$\begin{aligned} -D_T \mathbf{n} \cdot \nabla \rho_+ &= D_T \rho_+ - D_E \phi, \\ D_T \mathbf{n} \cdot \nabla \rho_- &= D_T \rho_- - D_E \phi. \end{aligned}$$

From these equations, we then obtain the new boundary conditions from (5.7) and the interface velocity from (5.8), after returning to the unprimed variables, as

$$-D_T \mathbf{n} \cdot \nabla \rho_+ = D_T(\rho_+ - \rho_*) - \gamma \kappa, \quad (5.9)$$

$$D_T \mathbf{n} \cdot \nabla \rho_- = D_T(\rho_- - \rho_*) - \gamma \kappa, \quad (5.10)$$

$$v = (f_+ + f_-) + \beta \rho_{*yy} + (\gamma/D_E) \kappa_{yy} \quad (5.11)$$

where

$$\begin{aligned} \rho_* &= \left( \frac{n_{123}}{4m_{123}l_{123}^2} \right)^{1/3} \frac{D_E}{D_T} (f_+ + f_-)^{2/3}, \\ \gamma &= \frac{l_{123}}{6m_{123}} D_E, \\ \beta &= D_T/D_E, \\ \kappa &= -X_{yy}. \end{aligned}$$

The boundary conditions (5.9) and (5.10) with a reference density  $\rho_*$  and a line tension term  $\gamma\kappa$  are of the Gibbs-Thomson form, and the velocity formula (5.11) has the surface diffusion term  $(\gamma/D_E)\kappa_{yy}$ . Note however the  $\rho_*$  is a steady state value from [3], rather than the equilibrium value from [2], and the coefficient  $\gamma$  comes from a transition barrier energy rather than an equilibrium energy. The term  $\beta\rho_{*yy}$  in the velocity formula (5.11) is new. To the best of our knowledge this derivation, of the Gibbs-Thomson form in the adatom boundary conditions and the surface diffusion term in the velocity, is the first that is based on microscopic (i.e., atomistic) dynamics, rather than thermodynamics.

## 6 The infinitely fast terrace diffusion limit

Assume that the adatoms diffuse infinitely fast and attach uniformly to step edges. Then, the adatom density is  $\rho = 0$  on steps and the total flux to a step edge is

$$f = f_+ + f_- = 2FL.$$

Assume also that there is no detachment of atoms from an edge or a kink to a step. Then, the original island dynamics model with step-edge kinetics is simplified to the following model—the simplified model of one-dimensional surface diffusion and kink convection with infinitely fast terrace diffusion limit

$$\partial_t \phi - D_E \partial_s^2 \phi = 2FL - v(\phi\kappa + 1), \quad (6.1)$$

$$\partial_t k - l_1 D_E \partial_s (\phi \partial_y X) = 2m_1 D_E \phi^2 - \frac{1}{2} n_1 D_E \phi k^2 + \frac{1}{2} n_1 D_E \phi (\partial_y X)^2, \quad (6.2)$$

$$v = l_1 D_E \phi k \cos \theta, \quad (6.3)$$

where  $D_E$  is the edge diffusion constant and the coefficients  $l_1$ ,  $m_1$ , and  $n_1$  are positive constants. This system only involves a moving step edge represented by the curve  $x = X(y, t)$ , the edge-adatom density  $\phi(y, t)$ , and the kink density  $k(y, t)$ .

The steady-state solution for this system is given by

$$X_0(y, t) = v_0 t, \quad v_0 = 2FL, \quad \phi_0 = \frac{n_1}{4m_1} k_0^2, \quad k_0 = \left( \frac{4m_1}{l_1 n_1} P_E \right)^{1/3}. \quad (6.4)$$

The linearized system around this steady-state solution is given by

$$\partial_t \phi_1 - D_E \partial_y^2 \phi_1 + \partial_t X_1 - v_0 \phi_0 \partial_y^2 X_1 = 0, \quad (6.5)$$

$$\frac{1}{2} n_1 D_E k_0^2 \phi_1 - \partial_t k_1 - n_1 D_E \phi_0 k_0 k_1 + l_1 D_E \phi_0 \partial_y^2 X_1 = 0, \quad (6.6)$$

$$l_1 D_E k_0 \phi_1 + l_1 D_E \phi_0 k_1 - \partial_t X_1 = 0. \quad (6.7)$$

The existence of a nontrivial periodic solution  $(X_1, \phi_1, k_1)$  of the form (4.46) leads to the dispersion relation  $\omega = \omega(l)$  that is determined by

$$\begin{aligned} & (\omega/D_E)^3 + (l^2 + n_1 P_E/l_1 + l_1 k_0) (\omega/D_E)^2 + ((P_E n_1/l_1 + P_E^2 + l_1^2 \phi_0^2) l^2 \\ & + \frac{3}{2} n_1 P_E k_0) (\omega/D_E) + \left( \frac{3n_1}{2l_1} P_E^3 l^2 + l_1^2 \phi_0^2 l^4 \right) = 0. \end{aligned} \quad (6.8)$$

If we drop the terms  $\partial_t \phi$  and  $\partial_t k$  in (6.1) and (6.2), respectively, we obtain the corresponding steady system

$$-D_E \partial_y^2 \phi = 2FL + v(\phi \partial_y \theta - 1), \quad (6.9)$$

$$-l_1 D_E \partial_y (\phi \partial_y X) = 2m_1 D_E \phi^2 - \frac{1}{2} n_1 d\phi k^2 + \frac{1}{2} n_1 D_E \phi (\partial_y X)^2, \quad (6.10)$$

$$v = l_1 D_E \phi k \cos \theta. \quad (6.11)$$

This system has the same steady-state solution as that for the unsteady system (6.1)–(6.3), cf. (6.4). And the corresponding linearized system can be obtained by dropping the two terms

$\partial_t \phi_1$  and  $\partial_t k_1$  in (6.5)–(6.7). The dispersion relation in this case can be obtained explicitly as

$$\omega(l) = -\frac{l_1 D_E \phi_0 l^2 (2l_1 l^2 + 3n_1 P_E k_0^2)}{n_1 k_0 (2l^2 + 3l_1 k_0)}. \quad (6.12)$$

Notice that the dispersion relation for both the steady and unsteady systems of this model is of the form

$$\omega = D_E T(P_E, l)$$

with  $T$  a function of two variables.

Denote for each wavenumber  $l \geq 0$  by  $\omega_1 = \omega_1(l)$ ,  $\omega_2 = \omega_2(l)$ , and  $\omega_3 = \omega_3(l)$  the three roots of the cubic equation (6.8). Assume that  $\omega_1(l)$  is always real and that  $\omega_1(0) = 0$ . The following proposition gives the asymptotics of the dispersion relation for both small and large wavenumbers; its proof is similar to and much simpler than that for Proposition 4.2, and is therefore omitted here:

**Proposition 6.1** *For the steady system, we have*

$$\omega(l) < 0 \quad \text{for all } l > 0,$$

and

$$\begin{aligned} \omega(l) &= -v_0 \phi_0 l^2 + O(l^4) & \text{as } l \rightarrow 0, \\ \omega(l) &= -\frac{v_0 w_0}{4h_0} l^2 + O(1) & \text{as } l \rightarrow \infty. \end{aligned}$$

For the unsteady system, we have

$$\begin{aligned} \omega_1(0) &= 0, & \operatorname{Re}(\omega_1(0)) &< 0, & \operatorname{Re}(\omega_2(0)) &< 0, \\ \omega_1(l) &= -v_0 \phi_0 l^2 + O(l^4) & \text{as } l \rightarrow 0, \end{aligned}$$

and

$$\begin{aligned} \omega_1(l) &= -D_E l^2 + O(1), \\ \omega_2(l) &= r_0 + iw_0 l + O(l^{-1}), \\ \omega_3(l) &= r_0 - iw_0 l + O(l^{-1}), \end{aligned}$$

as  $l \rightarrow \infty$ , where  $i = \sqrt{-1}$  and

$$r_0 = -\frac{1}{2} D_E P_E (P_E + n_1/l_1) < 0.$$

By comparing Propositions 4.2 and 6.1, we see that the linear stability for both of the original island dynamics model with step-edge kinetics and the simplified model with the infinitely fast terrace diffusion limit are essentially the same. In particular, the dispersion relation for both of the corresponding steady systems are exactly the same.

## A Perturbation for Free Boundaries

In the analysis above, perturbation solutions are found for PDEs involving free boundaries. The method for this analysis is briefly described here. Consider a function

$$u = u(x, t, \epsilon) = u_0 + \epsilon u_1 + \dots$$

satisfying  $u = 0$  at

$$x = X(t, \epsilon) = X_0 + \epsilon X_1 + \dots,$$

where  $\epsilon$  is a parameter small in magnitude. This boundary condition is then

$$0 = u(X(t, \epsilon), t, \epsilon) = u_0(X_0(t), t) + \epsilon (u_1(X_0(t), t) + X_1 \partial_x u_0(X_0(t), t)) + \dots.$$

It follows that the equations for  $u_0$  and  $u_1$  are

$$\begin{aligned} u_0(X_0(t), t) &= 0, \\ u_1(X_0(t), t) &= -X_1 \partial_x u_0(X_0(t), t). \end{aligned}$$

## B Proof of Proposition 4.1

Set for a fixed  $P_E$  (i.e., fixed  $\varepsilon = P_E^{1/3}$ )

$$H(\xi) = \xi(2m_{123}\xi + m_{23}\varepsilon^3)(2l_{123}\xi + l_{23}\varepsilon^3)^2 - \varepsilon^6(2n_{123}\xi + n_{23}\varepsilon^3) \quad \forall \xi \in \mathbf{R}.$$

It follows from (4.21)–(4.25) that

$$H(\phi_0) = 8D_E^{-3}w_0^2(g_0 - h_0) = 0.$$

But one easily verifies that  $H(0) < 0$ ,  $H(\xi) \rightarrow +\infty$  as  $\xi \rightarrow \infty$ , and  $H''(\xi) > 0$  for all  $\xi \in \mathbf{R}$ . Hence,  $\phi_0$  is the unique positive solution of  $H(\xi) = 0$  for a fixed  $\varepsilon$ . Since  $\phi_0$  depends only on  $\varepsilon$ , we deduce from the steady-state solution (4.17)–(4.25) that  $k_0$ ,  $w_0/D_E$ ,  $g_0/D_E$ , and  $h_0/D_E$  also depend only on  $\varepsilon$ . In particular, we have by (4.21) and (4.22) that

$$k_0 = 2\varepsilon^3(2l_{123}\phi_0 + l_{23}\varepsilon^3)^{-1}.$$

Direct calculations based on the fact that  $H(\phi_0) = 0$  for each  $\varepsilon > 0$  lead to the small  $\varepsilon$  asymptotics (4.27) and (4.28), and the following large  $\varepsilon$  asymptotics

$$\phi_0 = \sigma_0 + \sigma_1\varepsilon^{-3} + O(\varepsilon^{-6}) \quad \text{as } \varepsilon \rightarrow \infty, \quad (\text{B.1})$$

$$k_0 = \frac{2}{l_{23}} - \frac{4l_{123}\sigma_0}{l_{23}^2}\varepsilon^{-3} + O(\varepsilon^{-6}) \quad \text{as } \varepsilon \rightarrow \infty, \quad (\text{B.2})$$

where  $\sigma_0$  and  $\sigma_1$  are defined in (4.29). The Eqs. (B.1) and (B.2) imply (4.30).

Set  $\xi = \phi_0$  as a function of  $\varepsilon$ . Differentiating both sides of the equation  $H(\xi) = 0$  with respect to  $\varepsilon$ , we have by a series of calculations that

$$\frac{\varepsilon}{3\xi}\xi'(\varepsilon) = \frac{n_{23}\varepsilon^9 - l_{23}^2 m_{23} \varepsilon^9 \xi + 4l_{123}(2l_{23}m_{123} + l_{123}m_{23})\varepsilon^3 \xi^3 + 16l_{123}^2 m_{123} \xi^4}{n_{23}\varepsilon^9 + 2l_{23}(l_{23}m_{123} + 2l_{123}m_{23})\varepsilon^6 \xi^2 + 8l_{123}(2l_{23}m_{123} + l_{123}m_{23})\varepsilon^3 \xi^3 + 24l_{123}^2 m_{123} \xi^4}.$$

Consequently,  $\xi'(\varepsilon) > 0$  if  $\xi \leq \sigma_0$ . Moreover, if  $\sigma_1 < 0$ , then  $\xi'(\varepsilon) > 0$  for large  $\varepsilon$ . Otherwise, if  $\sigma_1 > 0$ , then  $\xi'(\varepsilon) < 0$  for large  $\varepsilon$ . These properties, together with (4.30), imply the rest of the proposition. **Q.E.D.**

## C Dispersion Relation

Inserting the expressions (4.38)–(4.43), (4.45), and (4.46) into (4.34)–(4.37) and (4.44), we get by a series of calculations that

$$\begin{aligned} D_T(l \cosh(lL) + \sinh(lL)) \hat{s}_1 - F(1+L) \hat{X}_1 &= 0, \\ D_T(l \sinh(lL) + \cosh(lL)) \hat{r}_1 - D_E \hat{\phi}_1 &= 0, \\ 2D_T \cosh(lL) \hat{r}_1 - (2D_E + D_E l^2 + \omega) \hat{\phi}_1 + (-v_0 \phi_0 l^2 - \omega) \hat{X}_1 &= 0, \\ D_T G_r \cosh(lL) \hat{r}_1 + D_T G_s \sinh(lL) \hat{s}_1 + G_\phi \hat{\phi}_1 + (G_k - \omega) \hat{k}_1 + (G_X - w_0 l^2) \hat{X}_1 &= 0, \\ D_T V_r \cosh(lL) \hat{r}_1 + D_T V_s \sinh(lL) \hat{s}_1 + V_\phi \hat{\phi}_1 + V_k \hat{k}_1 + (V_X - \omega) \hat{X}_1 &= 0, \end{aligned}$$

where

$$\begin{aligned} G_r &= 2m_{23}\phi_0 - \frac{1}{2}n_{23}k_0^2, \\ G_s &= 2(m_2 - m_3)\phi_0 - \frac{1}{2}(n_2 - n_3)k_0^2, \\ G_\phi &= 2m_1 D_E \phi_0 + \frac{2g_0}{\phi_0} - \frac{1}{2}n_1 D_E k_0^2, \\ G_k &= -\frac{4h_0}{k_0}, \\ G_X &= 2(m_3 - m_2)FL\phi_0 - \frac{1}{2}(n_3 - n_2)k_0^2 FL, \end{aligned}$$

and

$$\begin{aligned} V_r &= l_{23}k_0, \\ V_s &= (l_2 - l_3)k_0, \\ V_\phi &= l_1 D_E k_0, \end{aligned}$$

$$\begin{aligned} V_k &= w_0, \\ V_X &= (l_3 - l_2)k_0FL. \end{aligned}$$

Thus, there is a nonzero solution  $(X_1, \rho_1, \phi_1, k_1)$  of the given form if and only if the determinant of the coefficient matrix of the above linear system for  $(\hat{r}_1, \hat{s}_1, \hat{\phi}_1, \hat{k}_1, \hat{X}_1)$  vanishes.

Notice that this determinant is zero if  $l = 0$ . Assume  $l \neq 0$ . Factoring  $D_T \cosh(lL)$  and  $D_T \sinh(lL)$  from the first and second columns of the determinant, respectively, we see that the condition that the determinant is zero becomes

$$\begin{vmatrix} 0 & l \coth(lL) + 1 & 0 & 0 & -F(1+L) \\ l \tanh(lL) + 1 & 0 & -D_E & 0 & 0 \\ 2 & 0 & -(2D_E + D_E l^2 + \omega) & 0 & -v_0 \phi_0 l^2 - 1\omega \\ G_r & G_s & G_\phi & G_k - \omega & G_X - w_0 l^2 \\ V_r & V_s & V_\phi & V_k & V_X - \omega \end{vmatrix} = 0. \quad (\text{C.1})$$

This determines the dispersion relation  $\omega = \omega(l)$  for  $l \neq 0$  for the quasi-steady system.

For the steady system, the dispersion relation  $\omega = \omega(l)$  for  $l \neq 0$  is determined similarly by

$$\begin{vmatrix} 0 & l \coth(lL) + 1 & 0 & 0 & -F(1+L) \\ l \tanh(lL) + 1 & 0 & -D_E & 0 & 0 \\ 2 & 0 & -(2D_E + D_E l^2) & 0 & -v_0 \phi_0 l^2 - \omega \\ G_r & G_s & G_\phi & G_k & G_X - w_0 l^2 \\ V_r & V_s & V_\phi & V_k & V_X - \omega \end{vmatrix} = 0. \quad (\text{C.2})$$

A simple manipulation of the above determinants using the fact that all  $v_0/D_E$ ,  $w_0/D_E$ ,  $\phi_0$ , and  $k_0$  depend only on the edge Péclet number  $P_E$  shows that these dispersion relations are of the form

$$\omega(l) = D_E S(L, P_E, l),$$

where  $S$  is a function of three variables. In particular, the dispersion relation is independent of the adatom hopping rate  $D_T$  for both the steady and quasi-steady systems. This results from the exclusion of the Ehrlich-Schwoebel effect [6, 16, 17] in our assumption.

## D Proof of Proposition 4.2

Consider first the steady system. By (C.2) we have  $\omega(l) = A(l)/B(l)$ , where

$$A(l) = \begin{vmatrix} 0 & l \coth(lL) + 1 & 0 & 0 & -F(1+L) \\ l \tanh(lL) + 1 & 0 & -D_E & 0 & 0 \\ 2 & 0 & -(2D_E + D_E l^2) & 0 & -v_0 \phi_0 l^2 \\ G_r & G_s & G_\phi & G_k & G_X - w_0 l^2 \\ V_r & V_s & V_\phi & V_k & V_X \end{vmatrix},$$

$$B(l) = \begin{vmatrix} 0 & l \coth(LL) + 1 & 0 & 0 & 0 \\ l \tanh(LL) + 1 & 0 & -D_E & 0 & 0 \\ 2a & 0 & -(2D_E + D_E l^2) & 0 & 1 \\ G_r & G_s & G_\phi & G_k & 0 \\ V_r & V_s & V_\phi & V_k & 1 \end{vmatrix}.$$

By expanding  $A(l)$  and  $B(l)$  along the first row and the last column, respectively, we get that

$$A(l) = v_0 \phi_0 q_0 l^2 + O(l^4) \quad \text{and} \quad B(l) = -q_0 + O(l^2) \quad \text{as } l \rightarrow 0,$$

where

$$q_0 = \begin{vmatrix} 1 & -D_E & 0 \\ G_r & G_\phi & G_k \\ V_r & V_\phi & V_k \end{vmatrix} = 4l_{123} D_E h_0 + 2m_{123} D_E \phi_0 w_0 + \frac{1}{2\phi_0} n_{23} k_0^2 w_0 F L > 0,$$

and that

$$A(l) = D_E w_0^2 l^6 + O(l^4) \quad \text{and} \quad B(l) = D_E G_k l^4 + O(l^2) \quad \text{as } l \rightarrow \infty.$$

These imply (4.47) and (4.48).

Consider now the quasi-steady system. Notice first that  $\omega_1(l) \rightarrow \omega_1(0) = 0$  as  $l \rightarrow 0$ . By a series of calculations, we get from (C.1) that

$$q_0 (\omega_1(l) + v_0 \phi_0 l^2) + O(l^4) + O(l^2 \omega_1(l)) + O(\omega_1(l)^2) = 0 \quad \text{as } l \rightarrow 0,$$

implying (4.49).

Observe from (C.1) that  $|\omega(l)| \rightarrow \infty$  as  $l \rightarrow \infty$ . A simple manipulation of the determinant (C.1) leads to

$$\begin{vmatrix} -(2D_E + D_E l^2 + \omega) & 0 & -v_0 \phi_0 l^2 - \omega \\ G_\phi & G_k - \omega & G_X - w_0 l^2 \\ V_\phi & V_k & V_X - \omega \end{vmatrix} + O(l|\omega|) + O(|\omega|^2/l) = 0,$$

for large  $l$ . Consequently,

$$\begin{aligned} \omega^3 + D_E l^2 \omega^2 + (2D_E + V_\phi - G_k - V_X) \omega^2 + (w_0^2 - D_E G_k - D_E V_X + V_\phi v_0 \phi_0) l^2 \omega \\ + D_E w_0^2 l^4 + O(l|\omega|) + O(|\omega|^2/l) + O(l^2) = 0 \quad \text{as } l \rightarrow \infty. \end{aligned} \quad (\text{D.1})$$

It then follows immediately that there exist constants  $c_1 > 0$  and  $c_2 > 0$  such that

$$c_1 l \leq |\omega(l)| \leq c_2 l^2 \quad \text{for large } l.$$

If  $|\omega(l)|/l^2$  is bounded below by a positive constant for large  $l$ , then (D.1) implies that

$$\omega(l) = -D_E l^2 + O(l) \quad \text{as } l \rightarrow \infty. \quad (\text{D.2})$$

If, on the other hand, up to a subsequence of  $\{l\}$ ,  $\omega(l)/l^2 \rightarrow 0$  as  $l \rightarrow \infty$ , then we must have that  $\omega(l) = O(l)$ , for otherwise, we would have a contradiction from (D.1). Therefore, the highest order terms in (D.1) are those of  $l^2\omega^2$  and  $l^4$ . Consequently, we have

$$\omega(l) = R(l) + i\sigma\omega_0 l \quad \text{as } l \rightarrow \infty, \quad (\text{D.3})$$

where  $R(l)$  is a bounded function of  $l$  and  $\sigma = 1$  or  $-1$ . If we plug the above expression back into (D.1), we then have

$$R(l) = R_0 + O(l^{-1}) \quad \text{as } l \rightarrow \infty, \quad (\text{D.4})$$

where  $R_0$  is defined by (4.51). Now, (4.50) follows from (D.2)–(D.4).

Finally, the fact that  $R_0 < 0$  for small edge Péclet number  $P_E$  follows from (4.27) and (4.28). The condition (4.52) is obviously sufficient for  $R_0 < 0$  for all  $P_E$ . The necessary condition (4.53) follows from (4.30). **Q.E.D.**

## References

- [1] G. S. Bales and A. Zangwill. Morphological instability of a terrace edge during step-flow growth. *Phy. Rev. B*, 41(9):5500–5508, 1990.
- [2] W. K. Burton, N. Cabrera, and F. C. Frank. The growth of crystals and the equilibrium of their surfaces. *Phil. Trans. Roy. Soc. London Ser. A*, 243(866):299–358, 1951.
- [3] R. E. Caflisch, W. E. M. Gyure, B. Merriman, and C. Ratsch. Kinetic model for a step edge in epitaxial growth. *Phys. Rev. E*, 1999.
- [4] R. E. Caflisch, M. Gyure, B. Merriman, S. Osher, C. Ratsch, D. Vvedensky, and J. Zink. Island dynamics and the level set method for epitaxial growth. *Applied Math Letters*, 1999.
- [5] S. Chen, M. Kang, B. Merriman, R.E. Caflisch, C. Ratsch, R. Fedkiw, M.F. Gyure, and S. Osher. Level set method for thin film epitaxial growth. *Journ. Comp. Phys.*, 167:475–500, 2001.
- [6] G. Ehrlich and F. G. Hudda. Atomic view of surface diffusion: tungsten on tungsten. *J. Chem. Phys.*, 44:1036, 1966.
- [7] R. Ghez, H. G. Cohen, and J. B. Keller. The stability of growing or evaporating crystals. *J. Appl. Phys.*, 73:3685–3693, 1993.
- [8] R. Ghez and S. S. Iyer. The kinetics of fast steps on crystal surfaces and its application to the molecular beam epitaxy of silicon. *IBM J. Res. Develop.*, 32:804–818, 1988.

- [9] M. Gyure, C. Ratsch, B. Merriman, R. Caffisch, S. Osher, J. Zinck, and D. Vvedensky. Level set method for the simulation of epitaxial phenomena. *Phys. Rev. E*, 58:R6931, 1998.
- [10] J. B. Keller, H. G. Cohen, and G. J. Merchant. The stability of rapidly growing crystals. *J. Appl. Phys.*, 73:3694–3697, 1993.
- [11] F. Liu and H. Metiu. Stability and kinetics of step motion on crystal surfaces. *Phys. Rev. E*, 49:2601–2616, 1997.
- [12] B. Merriman, R. E. Caffisch, and S. Osher. Level set methods with applications to island dynamics. In G. Makrakis I. Athanasopoulos and J.F. Rodrigues, editors, *Free Boundary Problems: Theory and Applications*, pages 51–70. 16th International Symposium on Rarefied Gas Dynamics, 1999.
- [13] W. W. Mullins and R. F. Sekerka. Morphological stability of a particle growing by diffusion or heat flow. *J. Appl. Phys.*, 34:323, 1963.
- [14] W. W. Mullins and R. F. Sekerka. Stability of a planar interface during solidification of a dilute binary alloy. *J. Appl. Phys.*, 35:444, 1964.
- [15] P. Politi, G. Grenet, A. Marty, A. Ponchet, and J. Villain. Instabilities in crystal growth by atomic or molecular beams. *Physics Reports*, 324:271–404, 2000.
- [16] R. L. Schwoebel. Step motion on crystal surfaces II. *J. Appl. Phys.*, 40:614, 1969.
- [17] R. L. Schwoebel and E. J. Shipsey. Step motion on crystal surfaces. *J. Appl. Phys.*, 37:3682, 1966.