

Lipowsky *et al.* Reply: In our Letter¹ we introduced a multicomponent order parameter (OP) for surface melting and studied its behavior within phenomenological Landau models. The *minimal* set of OP components, $\mathbf{M}(\mathbf{Q}^{\parallel})$, as required by the translational invariance parallel to the surface, is parametrized by the reciprocal-lattice vectors \mathbf{Q}^{\parallel} of the two-dimensional surface. Since the translational invariance perpendicular to the surface is broken, these OP's also depend on the distance z from the surface: $\tilde{M} = \tilde{M}(\mathbf{Q}^{\parallel}, z)$. The OP components $\hat{M}(\mathbf{Q}^{\parallel}, \mathbf{Q}^{\perp}, z)$ considered by Löwen² satisfy

$$\tilde{M}(\mathbf{Q}^{\parallel}, z) = \sum \hat{M}(\mathbf{Q}^{\parallel}, \mathbf{Q}^{\perp}, z) \exp(i\mathbf{Q}^{\perp} \cdot \mathbf{e}_z z), \quad (1)$$

where the summation runs over all reciprocal-lattice vectors \mathbf{Q}^{\perp} such that $\mathbf{Q} = \mathbf{Q}^{\parallel} + \mathbf{Q}^{\perp}$ is a reciprocal-lattice vector of the three-dimensional bulk crystal.³

Quite generally, an interface or surface acts as a planar perturbation on any density (as long as the interfacial roughness can be ignored). For systems governed by short-range forces, such a perturbation decays exponentially $\sim \exp(-z/a)$ with the distance z from the interface but the decay length a depends on the density considered. Thus, within the liquid (or melt), the densities $\hat{M}(\mathbf{Q}^{\parallel}, \mathbf{Q}^{\perp}, z)$ and $\tilde{M}(\mathbf{Q}^{\parallel}, z)$ are characterized by decay lengths $\hat{a}_L(\mathbf{Q}^{\parallel}, \mathbf{Q}^{\perp})$ and $\tilde{a}_L(\mathbf{Q}^{\parallel}) = \max_{\perp} [\hat{a}_L(\mathbf{Q}^{\parallel}, \mathbf{Q}^{\perp})]$, respectively.

We found in Ref. 1 that the length scales $\tilde{a}_L(\mathbf{Q}^{\parallel})$ satisfy a simple algebraic relation within a square-gradient Landau model for the molten layer. It turns out, however, that such a relation holds for a *whole class* of Landau models which include gradients to arbitrary order as does the density-functional (DF) approach used by Löwen.² These Landau models are defined by

$$F_L\{M\} = \int d^2x \int_0^l dz \left\{ \frac{1}{2} (\nabla M)^2 + \sum_{m \geq 2} c_m M \nabla^{2m} M + \frac{1}{2} (M/a_{L0})^2 - H_{\text{eff}} M \delta(z-l) \right\} \quad (2)$$

for the density $M(\mathbf{x}, z)$ within the molten layer which is bounded by a vapor-liquid interface at $z=0$ and by a liquid-crystal interface at $z=l$. The effective field H_{eff} at the liquid-crystal interface represents the influence of the crystalline state on the molten layer and, thus, must reflect the translational symmetry parallel to the surface:

$$H_{\text{eff}}(\mathbf{x}, z) = \sum \tilde{H}(\mathbf{Q}^{\parallel}, z) \exp(i\mathbf{Q}^{\parallel} \cdot \mathbf{x}).$$

The model as given by (2) represents a generalization of Eq. (13) in Ref. 1.

Minimizing the above functional, one obtains

$$M(\mathbf{x}, z) = \sum \tilde{M}(\mathbf{Q}^{\parallel}) \exp[i\mathbf{Q}^{\parallel} \cdot \mathbf{x} - z/\tilde{a}_L(\mathbf{Q}^{\parallel})], \quad (3)$$

with

$$\tilde{a}_L(\mathbf{Q}^{\parallel}) = \tilde{a}_L(\mathbf{0}) / \{1 + [\tilde{a}_L(\mathbf{0})\mathbf{Q}^{\parallel}]^2\}^{1/2}, \quad (4)$$

where $\tilde{a}_L(\mathbf{0})$ depends on a_{L0} and on the coefficients c_m in (2).

On the other hand, the DF approach in Ref. 2 leads to length scales $\tilde{a}_L(\mathbf{Q}^{\parallel})$ which do not satisfy (4) for small values of \mathbf{Q}^{\parallel} even though they seem to approach this relation for large \mathbf{Q}^{\parallel} . Furthermore, the scales for small \mathbf{Q}^{\parallel} are found to depend on the microscopic interparticle potential. In general, such a dependence, which cannot be obtained from phenomenological models as in (2), is indeed to be expected.

The length scales $\tilde{a}_L(\mathbf{Q}^{\parallel})$ determine the critical behavior of the surface OP's, $\tilde{M}(\mathbf{Q}^{\parallel}, z=0)$. As pointed out in Ref. 1, all surface OP's *vanish simultaneously* at the melting temperature, $T=T_*$, *but with different rates*. For systems governed by short-range forces, the critical behavior is given by (Ref. 1)

$$\tilde{M}(\mathbf{Q}^{\parallel}, z=0) \sim (T_* - T)^{\beta_1}, \quad \beta_1 = \frac{b\tilde{a}_L(\mathbf{0})}{\tilde{a}_L(\mathbf{Q}^{\parallel})}. \quad (5)$$

For the models considered in Ref. 1, the scales $\tilde{a}_L(\mathbf{Q}^{\parallel})$ satisfy the relation (4), $\tilde{a}_L(\mathbf{0}) = a_{L0}$, and $b=1$ or $b=\frac{1}{2}$. These models have, however, one important limitation since the density $\tilde{M}(\mathbf{Q}^{\parallel}=\mathbf{0}, z)$ which has the largest decay length is, in fact, *not* treated as an OP density (which would vanish at $T=T_*$) but rather as a *nonordering* density which stays finite at $T=T_*$.⁴ This limitation is not present if one considers the densities $\hat{M}(\mathbf{Q}^{\parallel}, \mathbf{Q}^{\perp}, z)$ with decay lengths $\hat{a}_L(\mathbf{Q}^{\parallel}, \mathbf{Q}^{\perp})$. Indeed, the largest decay length $\tilde{a}_L(\mathbf{0})$ can now belong to the nonordering density $\hat{M}(\mathbf{0}, \mathbf{0}, z)$ or to one of the OP densities $\tilde{M}(\mathbf{0}, \mathbf{Q}^{\perp}, z)$ with $\mathbf{Q}^{\perp} \neq \mathbf{0}$. The power-law behavior as in (5) still applies but three cases must be distinguished depending on the relative size of the two length scales $\kappa \equiv \hat{a}_L(\mathbf{Q}^{\parallel}=\mathbf{0}, \mathbf{Q}^{\perp}=\mathbf{0})$ and $\xi_0 \equiv \max_{\perp} [\hat{a}_L(\mathbf{Q}^{\parallel}=\mathbf{0}, \mathbf{Q}^{\perp})]$, where the maximum is taken over all $\mathbf{Q}^{\perp} \neq \mathbf{0}$. In terms of these scales, the coefficient b in (5) is given by $b=1$, $b=\kappa/\xi_0$, and $b=\frac{1}{2}$ for $\tilde{a}_L(\mathbf{0}) = \kappa > \xi_0$, $\tilde{a}_L(\mathbf{0}) = \xi_0 > \kappa > \xi_0/2$, and $\tilde{a}_L(\mathbf{0}) = \xi_0/2 > \kappa$, respectively.

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¹R. Lipowsky, U. Breuer, K. C. Prince, and M. P. Bonzel, Phys. Rev. Lett. **62**, 913 (1989).

²H. Löwen, preceding Comment, Phys. Rev. Lett. **64**, 2104 (1990).

³One should note, however, that the densities $\hat{M}(\mathbf{Q}^{\parallel}, \mathbf{Q}^{\perp}, z)$ are not uniquely defined by (1).

⁴R. Lipowsky, in Proceedings of the International Workshop on Magnetic Properties of Low-Dimensional Systems, San Luis Potosi, Mexico, Springer Proceedings in Physics (Springer-Verlag, New York, to be published).