

Defectlike structures and localized patterns in the cubic-quintic-septic Swift-Hohenberg equationEdgar Knobloch,¹ Hannes Uecker,^{2,*} and Daniel Wetzel²¹*Department of Physics, University of California, Berkeley, California 94720, USA*²*Institut für Mathematik, Universität Oldenburg, D26111 Oldenburg, Germany*

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We study numerically the cubic-quintic-septic Swift-Hohenberg (SH357) equation on bounded one-dimensional domains. Under appropriate conditions stripes with wave number $k \approx 1$ bifurcate supercritically from the zero state and form S-shaped branches resulting in bistability between small and large amplitude stripes. Within this bistability range we find stationary heteroclinic connections or fronts between small and large amplitude stripes, and demonstrate that the associated spatially localized defectlike structures either snake or fall on isolas. In other parameter regimes we also find heteroclinic connections to spatially homogeneous states and a multitude of dynamically stable steady states consisting of patches of small and large amplitude stripes with different wave numbers or of spatially homogeneous patches. The SH357 equation is thus extremely rich in the types of patterns it exhibits. Some of the features of the bifurcation diagrams obtained by numerical continuation can be understood using a conserved quantity, the spatial Hamiltonian of the system.

DOI: [10.1103/PhysRevE.100.012204](https://doi.org/10.1103/PhysRevE.100.012204)**I. INTRODUCTION**

Defects play an important role in both condensed matter physics and in pattern-forming systems. Topological defects arise when the pattern amplitude vanishes and organize both one- and two-dimensional patterns [1,2]. As a result their motion and creation or annihilation lead to nonlocal adjustment of the pattern wave number. Nontopological defects may involve a continuous but spatially localized transition from one wave number to another, and such transitions are usually referred to as fronts [1,2]. Fronts connecting distinct states are common in systems supporting waves as exemplified by the complex Ginzburg-Landau equation [3–5] and pattern formation in fluid dynamics [6] and chemistry [7], and are found in both wave-supporting systems and in stationary patterns. A mathematical classification of defects in one spatial dimension is provided in Ref. [8].

In this paper we are interested in properties of stationary fronts between patterns with distinct wave numbers in one spatial dimension—the simplest defect—and ask two basic questions. Do such defects come in distinct families and do they persist over time? The first question is motivated by the notion of pinning [9], whereby a front is pinned to heterogeneities on either side of the front which can be viewed as trapping the front in a particular location. Energy has to be supplied to overcome the resulting pinning potential to move the defect to a different location. The second question has to do with stability of a front. A front may lose stability because of the loss of stability of either of the far-field states or due to a localized mode at the location of the front. These instabilities are usually associated with the presence of unstable continuous and point spectra, respectively. We remark that the classical amplitude-phase description of patterns misses

the pinning effect just described since it reduces a uniform pattern to a constant amplitude state and hence treats such states as translation-invariant states. To study fronts between different wave numbers it is essential therefore to go beyond the amplitude-phase description.

The fact that defects are spatially localized structures and that such structures readily pin to heterogeneities suggests that they might come in families organized within a snaking bifurcation diagram. In the context of pattern-forming systems the term homoclinic snaking refers to a pair of intertwined branches of spatially localized patterns that oscillate back and forth across a region in parameter space (the snaking or pinning region), usually accompanied by repeated changes in stability. Prototypes for this scenario in one spatial dimension (1D) are provided by the quadratic-cubic (SH23) and cubic-quintic (SH35) Swift-Hohenberg equations. These take the form

$$\partial_t u = \lambda u - (1 + \partial_x^2)^2 u + f(u) \quad (1)$$

with $f(u) = au^2 - u^3$ and $f(u) = au^3 - u^5$, respectively, and are parametrized by the parameters $\lambda \in \mathbb{R}$ and $a > 0$. See, e.g., Refs. [10–18] for basic properties of these equations and their interpretation. In higher dimensions the situation becomes more complicated, but snaking behavior of branches of localized patterns can also be observed. Snaking of localized hexagons in SH23 and related equations is discussed in, e.g., Refs. [19,20], while snaking of localized patterns in 2D reaction-diffusion systems on nonhomogeneous backgrounds is considered in [21], and of body-centered cubes on homogeneous and nonhomogeneous backgrounds in 3D reaction-diffusion systems in [22,23]. References [24,25] provide reviews of localization and snaking in various other systems and experiments, while Refs. [26–28] give analytical results on homoclinic snaking based on beyond all orders asymptotics.

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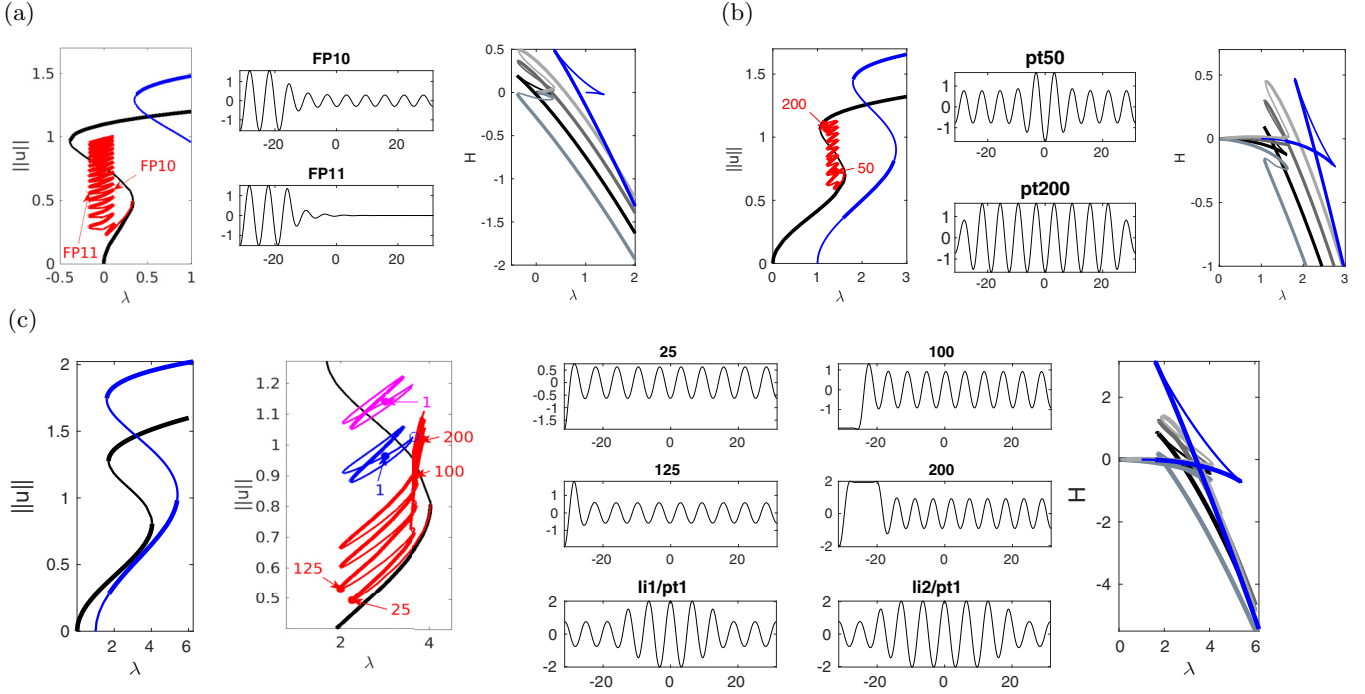


FIG. 1. Solutions of (2) on $\Omega = (-10\pi, 10\pi)$ with homogeneous Neumann BCs. (a) $a = 2$: bifurcation diagram (BD) of $u_{p,1}$ (black), $u_{\text{hom}}^{\text{large}}$ (blue) and one snaking branch (red) of fronts between $u_{p,1}^{\text{large}}$ and $u = 0$ (left folds) and $u_{p,1}^{\text{large}}$ and $u_{p,1}^{\text{small}}$ (right folds), respectively, showing bistability of $u_{p,1}^{\text{large}}$ with $u = 0$ for $\lambda < 0$, and bistability of $u_{p,1}^{\text{large}}$ and $u_{p,1}^{\text{small}}$ for $\lambda > 0$; the snake straddles $\lambda = 0$. (b) $a = 5$: “classical” snaking of heteroclinic cycles (red) between $u_{p,1}^{\text{large}}$ and $u_{p,1}^{\text{small}}$ (black). (c) $a = 9.5$: breakup of classical snaking into stacks of isolas, and a branch of solutions (red) involving $\pm u_{\text{hom}}^{\text{large}}$ (black). Sample solutions are provided in each case, followed, in the right panels, by plots of the spatial Hamiltonian $H(\cdot)$ showing self-intersections corresponding to different types of Maxwell points. Thick (thin) lines in the bifurcation diagrams and plots of H indicate linearly stable (unstable) solutions; see also Remark I.1(d) for the naming and plotting conventions in these and the following plots.

Motivated by the above considerations we focus here on 1D patterns but consider a more complex nonlinearity $f(u)$ than hitherto studied, in order to study snaking between two distinct periodic patterns. For this purpose we selected the cubic-quintic-septic (SH357) version of Eq. (1), namely,

$$\partial_t u = \lambda u - (1 + \partial_x^2)^2 u + f(u), \quad f(u) = -au^3 + bu^5 - u^7, \quad (2)$$

and study this equation on bounded domains $\Omega = (-l_x, l_x)$ with homogeneous Neumann boundary conditions (BCs) $\partial_x u|_{\partial\Omega} = \partial_x^3 u|_{\partial\Omega} = 0$. The linear stability properties of the trivial solution $u^* \equiv 0$ are independent of f and follow from the linearization $\partial_t v = -(1 + \partial_x^2)^2 v + \lambda v$. This equation has the solutions $v(x, t) = e^{ikx + \mu(k)t}$, $k \in \mathbb{R}$, where $\mu(k) = -(1 - k^2)^2 + \lambda$. Thus u^* is asymptotically stable for $\lambda < 0$ and unstable for $\lambda > 0$ with respect to periodic perturbations with wave number k near $k_c = 1$ and in 1D we expect pitchfork bifurcations to spatially periodic patterns for $\lambda \geq 0$, if permitted by the domain and BCs. In detail, if, e.g., $\Omega = (-l\pi, l\pi)$, $l \in \mathbb{N}$, then the admissible wave numbers are $k \in \frac{1}{2l}\mathbb{N}$, and for large l we have many bifurcation points for small $\lambda > 0$. The first bifurcation at $\lambda_1 = 0$ has $k_1 = 1$ and is followed by bifurcations to stripes with $k_{2,3} = 1 \pm 1/(2l)$, $k_{4,5} = 1 \pm 1/l, \dots$, corresponding to sidebands of $k = 1$.

Depending on the parameters $a, b > 0$ in Eq. (2) we find that the bifurcating branches are typically S-shaped, with stable small and stable large amplitude sections, and an unstable

intermediate amplitude section. For definiteness, we choose

$$b = 3.5 + 0.4(a - 3), \quad (3)$$

and consider a as a second free parameter, in addition to λ . We use the following abbreviations:

- (i) $u_{p,1}^{\text{small}}$ for the small amplitude part and $u_{p,1}^{\text{large}}$ for the large amplitude part of the periodic branch belonging to wave number $k = 1$;
- (ii) u_p^{small} and u_p^{large} (without specifying a wave number, but typically with k near $k_c = 1$) for general periodic solutions;
- (iii) $u_{\text{hom}}^{\text{small}}$ and $u_{\text{hom}}^{\text{large}}$ for the spatially homogeneous branch (wave number $k = 0$), bifurcating at $\lambda = 1$;
- (iv) u_*^{middle} , $*$ = p or $*$ = hom for the corresponding middle sections.

Figure 1 provides an overview of the dependence of the solution sets of Eq. (2) on a , and introduces the solution types considered in this paper. Thick lines indicate linearly stable states while thin lines correspond to linearly unstable states. For $a > 0$ and not too large the S shape of the first bifurcating branch is mild, and the subsequent branches are “neatly ordered” in the sense that only a short interval of the small amplitude periodic solutions is stable, and there is no overlap of the bistable range of the periodic solutions with wave number k near 1 with the branch of spatially homogeneous solutions (wave number $k = 0$) [Fig. 1(a) for $a = 2$]. If we increase a , then the S shape and hence the overlap of

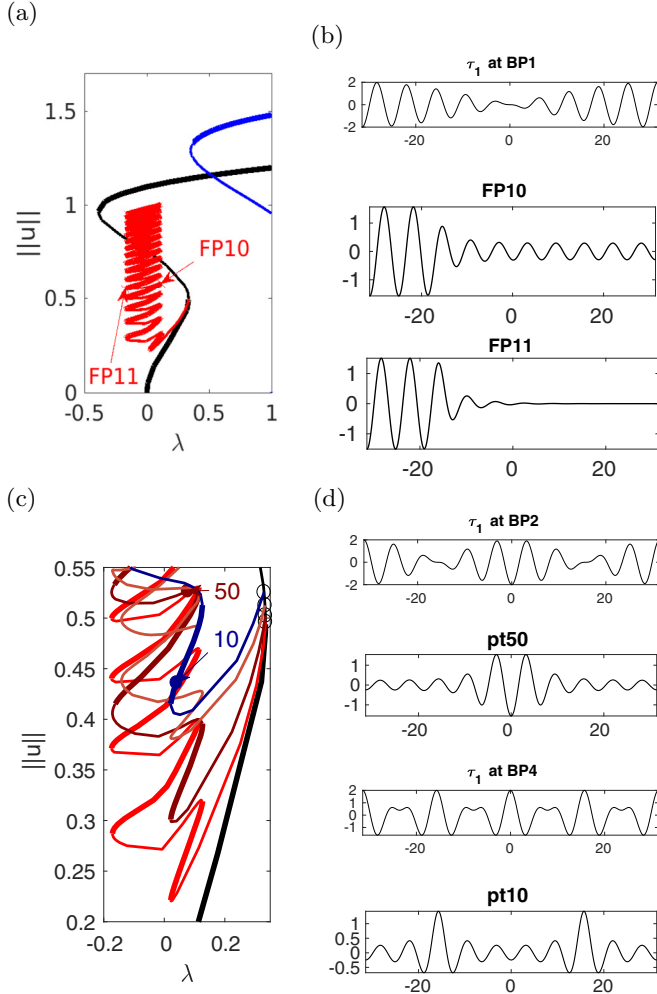


FIG. 2. $a = 2$. (a, b) Snaking fronts between u_p^{large} and u_p^{small} (for $\lambda > 0$), respectively, between u_p^{large} and $u = 0$ (for $\lambda \leq 0$, where u_p^{small} does not exist). (c) Zoom into the BD near the first fold on $u_{p,1}$. After the fold there are multiple BPs. The front solutions bifurcate at BP1 [red branch in (a)]. The first plot in (b) shows the tangent direction for this bifurcation. Branches of solutions with multiple interfaces bifurcate at subsequent BPs. For instance, a $u_p^{\text{small}} - u_p^{\text{large}} - u_p^{\text{small}}$ heteroclinic cycle bifurcates at BP2, and a heteroclinic cycle with two “pulses” at BP4; see (d) for sample plots.

the bistable ranges for solutions with different k become more pronounced [Fig. 1(b) for $a = 5$]. This overlap is of interest to us because we expect localized patterns to be generically present within such bistable (or indeed multistable) ranges.

It turns out that for the smaller a one finds “almost classical” snaking of localized states that is associated with heteroclinic cycles between u_p^{small} with wave numbers close to $k = 1$, and u_p^{large} , again with wave numbers close to $k = 1$. These states consist of a portion of u_p^{large} in a background of u_p^{small} or vice versa; localized states consisting of a portion of u_p^{large} in $u = 0$ background are also possible. For larger a , more and more branches (with wave numbers deviating from $k = 1$) enter the game, including the branch u_{hom} of spatially homogeneous solutions ($k = 0$), and the solution set of Eq. (2) becomes more and more complicated. For

instance, the snaking branches of small-to-large periodic patterns break up into isolas, and additional branches consisting of heteroclinic cycles between various distinct spatial patterns enter the picture. Figure 1 is thus intended as a preview of the subsequent results. The norm $\|u\|$ used in (a) and all similar plots is

$$\|u\| := \left(\frac{1}{|\Omega|} \int_{\Omega} u^2 dx \right)^{1/2} \quad (\text{normalized } L^2 \text{ norm}). \quad (4)$$

In summary, in this paper we numerically investigate how the set of localized patterns of Eq. (2) becomes richer and richer with increasing a , and finally “explodes” for $a \approx 9.5$ and larger. These results are presented in detail in Sec. II, while Sec. III provides a brief discussion.

Remark 1.1. (a) Equation (2) has a number of symmetries: (i) translational invariance (for $\Omega = \mathbb{R}$); (ii) odd symmetry $u \mapsto -u$; (iii) spatial reflection symmetry $x \mapsto -x$. The translational invariance (i) is broken over $\Omega = (-l_x, l_x)$ by the Neumann BCs $\partial_x u|_{\partial\Omega} = \partial_x^3 u|_{\partial\Omega} = 0$, but “periodic” solutions over $(-l_x, l_x)$ can be extended to all of \mathbb{R} by reflection at the boundaries. Thus the first “front” in Fig. 1(a) can also be seen as heteroclinic cycle between a large amplitude and a small amplitude periodic solution; (ii) implies that all nontrivial solutions are double, and we generally identify $\pm u_p^{\text{small}}$, $\pm u_p^{\text{large}}$, and $\pm u_{\text{hom}}$, respectively. (ii) and (iii) together imply that we have branches of *odd solutions* of the form $u(-x) = -u(x)$, as opposed to *even solutions* which have a maximum or minimum at $x = 0$. As a consequence, snaking branches come in odd and even families, and generally we expect ladder branches connecting these, and this *snakes-and-ladders structure* is a prerequisite for the breakup of snakes into isolas. These results are common to SH35 [29].

(b) Equation (2) is a gradient system, $\partial_t u = -\frac{\delta \mathcal{E}(u)}{\delta u}$, with respect to the energy

$$\mathcal{E}(u) = \int_{\Omega} \frac{1}{2} [(1 + \Delta)u]^2 - \frac{1}{2} \lambda u^2 - F(u) dx, \quad (5)$$

$F(u) \equiv \int_{\Omega} f(v) dv$, where either $\Omega = \mathbb{R}$, or $\Omega = (-l_x, l_x)$ with Neumann BCs $\partial_x u|_{\partial\Omega} = \partial_x^3 u|_{\partial\Omega} = 0$. In particular, local minima of \mathcal{E} are stable stationary solutions of Eq. (2), and Eq. (2) does not have time-periodic solutions (with finite energy). Moreover, the translational invariance of \mathcal{E} yields the existence of a spatially conserved quantity for steady solutions, a spatial Hamiltonian, e.g., [30, Proposition 1], here given by

$$H(u) = \partial_x u \partial_x^3 u - \frac{1}{2} (\partial_x^2 u)^2 + (\partial_x u)^2 + \frac{1}{2} (1 - \lambda) u^2 - F(u). \quad (6)$$

Hence, a necessary condition for a heteroclinic connection between, e.g., two periodic solutions u_p^{small} and u_p^{large} is that $H(u_p^{\text{small}}) = H(u_p^{\text{large}})$. For the classical SH23 and SH35 equations, this requirement provides an important wave-number selection principle that determines the wave number $k(\lambda)$ along the snaking branches. The same is true for SH357. Figure 1(e) indicates that while for small to moderate values of a there are few intersections of H for the different branches, this is no longer so for larger a where the number of possible heteroclinic cycles becomes very large.

(c) When choosing the bounded domain Ω for Eq. (2) we need to compromise between

(1) Generality: the results should be representative of the situation on large domains, ideally approximating the case $\Omega = \mathbb{R}$; this in general calls for large domains.

(2) Feasibility and clarity: the domain should be small enough to (i) avoid exceedingly expensive numerics, and (ii) allow clear plotting of results.

It turns out that the basic results can be studied and presented well on relatively small domains, for instance $\Omega = (-10\pi, 10\pi)$, which we use in most cases. In some cases, in particular for extracting wave numbers from mixed periodic solutions via Fourier transform, a larger Ω such as $\Omega = (-30\pi, 30\pi)$ is helpful. In any case, we checked that none of our results depends qualitatively on the domain size by running the same numerics on significantly larger domains, where however the results become more difficult to present graphically.

(d) For the plots of bifurcation diagrams (BDs) and sample solutions we use the following conventions. Stable parts of branches (as determined from the eigenvalues of the linearization around solutions) are plotted as thick lines, unstable parts as thinner lines. Dots labeled by an integer n correspond to solutions for which we plot profiles $u(x)$, titled “pt n ” if there is no ambiguity. Fold and branch points are indicated via FP n and BP n , respectively, and similarly in the title of the sample plots. Occasionally we give titles in the form branch/point.

(e) When $\Omega = \mathbb{R}$ the spatial dynamics picture implies that $u = 0$ is a saddle for $\lambda < 0$ with two stable and two unstable eigenvalues. Likewise, a robust connection to u_p^{small} or u_p^{large} requires these to be generalized saddles and hence that they have a three-dimensional center-stable manifold and a three-dimensional center-unstable manifold. This is a consequence of spatial reversibility and the conservation of H .

II. RESULTS

We use `pde2path` [31,32] to compute bifurcation diagrams for Eq. (2). As domain we typically choose $\Omega = (-10\pi, 10\pi)$, which is large enough to permit a multitude of patterns, cf. Remark I.1(c).

A. The case $a = 2$

We start with $a = 2$. In the plot of H as a function of λ in Fig. 1(a) we see that there are few self-intersections for the first four bifurcating branches, and in particular no overlap of their bistable ranges with the spatially homogeneous (blue) branch. This corresponds to the “easy” situation where relatively few heteroclinic connections are possible. Moreover, in this case transitions between different heteroclinic cycles and interesting codimension two points can be easily identified.

Figure 2 provides more details. In particular, we find that after the first fold on $u_{p,1}$ there are multiple BPs where branches of heteroclinic cycles with an increasing number of interfaces bifurcate. The first branch corresponds to fronts (but see also Remark I.1(a) for the interpretation of the front as a heteroclinic cycle via even parity continuation of the solution over the domain boundary), the second to a $u_p^{\text{small}} - u_p^{\text{large}} - u_p^{\text{small}}$ heteroclinic cycle, and so on. A particular feature of

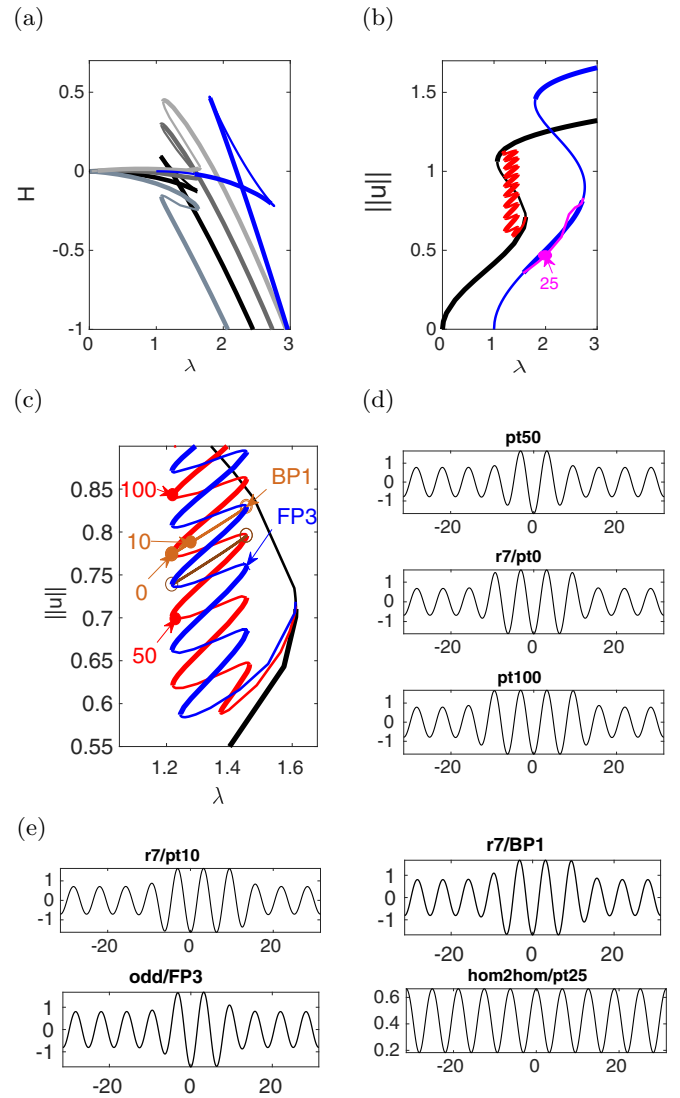


FIG. 3. $a = 5$. (a) H on the first four branches of periodic solutions (black to light gray), and the homogeneous solution u_{hom} ($k = 0$, blue). (b) $\|u\|$ for the primary periodic branch (black) and for u_{hom} (blue), together with a snake (red) of cycles between u_p^{small} and u_p^{large} , consisting of even solutions, and a pattern branch between two Turing bifurcations on u_{hom} (magenta). (c) Zoom of the snake, together with two rungs (brown) that connect the even snake and the odd snake (blue). (d) Even solutions (close to left folds) corresponding to (c). (e) Sample solutions from the light brown rung and the odd parity snake (blue branch) in (c), and from the magenta branch (hom2hom) in (b).

the small a regime is clearly visible on the first branch: for $\lambda < 0$, u_p^{large} connects to the trivial state $u = 0$, while for $\lambda > 0$ it connects to u_p^{small} . In other words, each time λ passes through $\lambda = 0$ into $\lambda > 0$ the hole in the solution fills in with small amplitude oscillations. On an infinite domain the heteroclinic cycle changes from one involving the $u = 0$ state to one involving u_p^{small} . Note that there will be parameter values such that the right folds of the snaking front branch just reach $\lambda = 0$, a situation corresponding to a codimension two bifurcation of heteroclinic cycles. There is a second codimension two transition that is relevant as well: when u_p^{small}

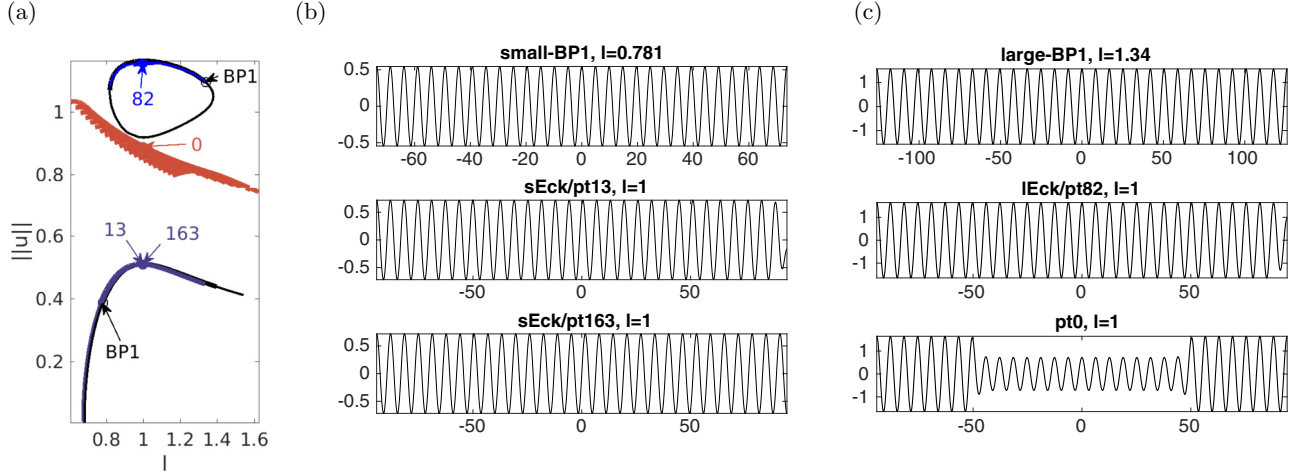


FIG. 4. Continuation of solutions of Eq. (7) in ℓ , with $a = 5$, $\lambda = 1.3$ fixed on base domain $\Omega = (-30\pi, 30\pi)$, i.e., on the effective domain (also used for solution plots) $\Omega_\ell = (-30\pi\ell, 30\pi\ell)$. Black branches correspond to u_p^{small} and u_p^{large} , with Eckhaus BPs indicated and bifurcating branches shown in blue. See Fig. 5 for zoom and details of the red snake.

changes from subcritical to supercritical the termination point of the localized solutions moves from $\lambda = 0$ to the right fold on u_p^{small} as in Fig. 2(a). An analogous transition has been observed in rotating plane Couette flow [33].

The front solutions bifurcate from BP1, the branching point nearest to the right fold of u_p^{small} ; other branching points, further away from the fold, give rise to solutions with multiple interfaces, as summarized in Figs. 2(c) and 2(d).

B. The case $a = 5$

1. Snakes and ladders of cycles between $u_{p,1}^{\text{small}}$ and $u_{p,1}^{\text{large}}$

For $a = 5$, the second (leftmost) fold on $u_{p,1}$ is at $\lambda > 0$, and the snakes bifurcating near the first fold lie entirely in the $\lambda > 0$ range. The red branch in Fig. 3(b) shows the (even) $u_p^{\text{small}} - u_p^{\text{large}} - u_p^{\text{small}}$ snake L_0 bifurcating at BP2. Looking more closely, one finds secondary bifurcation points near every fold on this branch giving rise to “rungs” [brown branches in (c)] connecting to the odd snake $[L_{\pi/2}]$, blue branch in (c). This ladder structure becomes important for understanding the breakup of snakes at larger a ; see Sec. II C. Additionally, in Fig. 3(b) we plot a branch connecting two Turing bifurcation points on u_{hom} . Note that except for $u_{p,1}^{\text{small}}$, all nontrivial branches are unstable at bifurcation but many become stable at (small but) finite amplitude.

2. Continuation in the domain size

Along the snaking branches of, e.g., $u_p^{\text{small}} - u_p^{\text{large}} - u_p^{\text{small}}$ heteroclinic cycles, the wavelengths (and hence amplitudes) of both u_p^{small} and u_p^{large} must in general change continuously as determined by the condition $H[u_p^{\text{small}}(\cdot; \lambda)] \stackrel{!}{=} H[u_p^{\text{large}}(\cdot; \lambda)]$. Here we look into this phenomenon in more detail, via continuation in the domain size scaling ℓ . Thus, we modify Eq. (2) to

$$\partial_t u = \lambda u - (1 + \ell^{-2} \partial_x^2)^2 u + f(u), \quad (7)$$

on $\Omega = (-l_x, l_x)$, such that the effective domain is $\Omega_\ell := \ell\Omega = (-\ell l_x, \ell l_x)$. Qualitative results are:

(a) In general, u_p^{large} is more rigid than u_p^{small} . This means that u_p^{large} adapts its wave number and amplitude less than u_p^{small} does.

(b) The continuation in ℓ leads to a new kind of snaking, where the small amplitude part of the heteroclinic cycles grows or shrinks via phase slips in u_p^{small} .

These results are illustrated in Figs. 4 and 5. Here, to accurately extract the wave numbers of the periodic patterns we choose a rather large base domain $\Omega = (-30\pi, 30\pi)$, although the phenomenon can be observed on significantly smaller domains, e.g., $\Omega = (-10\pi, 10\pi)$.

In Fig. 4 we show the results of continuing the *periodic* branches u_p^{large} and u_p^{small} in ℓ for both increasing and decreasing ℓ , starting from results computed earlier for $\ell = 1$ and $\lambda = 1.3$, and keeping λ fixed. For decreasing ℓ , u_p^{small} [lower black branch in Fig. 4(a)] loses stability at $\ell = \ell_0 \approx 0.78$ corresponding to an Eckhaus boundary point for u_p^{small} with wavelength ℓ_0 . The bifurcating branch sEck initiates an amplitude modulation at the right domain boundary leading to an increase in the pattern wavelength. With increasing ℓ this branch reconnects with the u_p^{small} branch near $\ell = 1.3$. In the displayed numerical continuation, this manifests itself in “branch jumping” to u_p^{small} with the corresponding ℓ , which we deliberately do not attempt to avoid here by refining the numerics. Instead, we simply continue the u_p^{small} branch so found back towards smaller ℓ , and find that this branch, like the original u_p^{small} branch, also loses stability near $\ell = 0.8$. This process can then be repeated. See (b) for sample solution profiles. We emphasize that the continuation of u_p^{small} in ℓ maintains the number of wavelengths in the domain. Thus the number of wavelengths in a solution can only change by encountering an Eckhaus point, i.e., by triggering a (dynamic) phase slip that takes a stable periodic solution with a certain number of wavelengths to a new stable periodic state with a different number of wavelengths [34].

Similar behavior takes place for u_p^{large} as well. With increasing ℓ , u_p^{large} loses stability at an Eckhaus point at $\ell = \ell_0 \approx 1.34$. Shortly thereafter the branch passes through a fold,

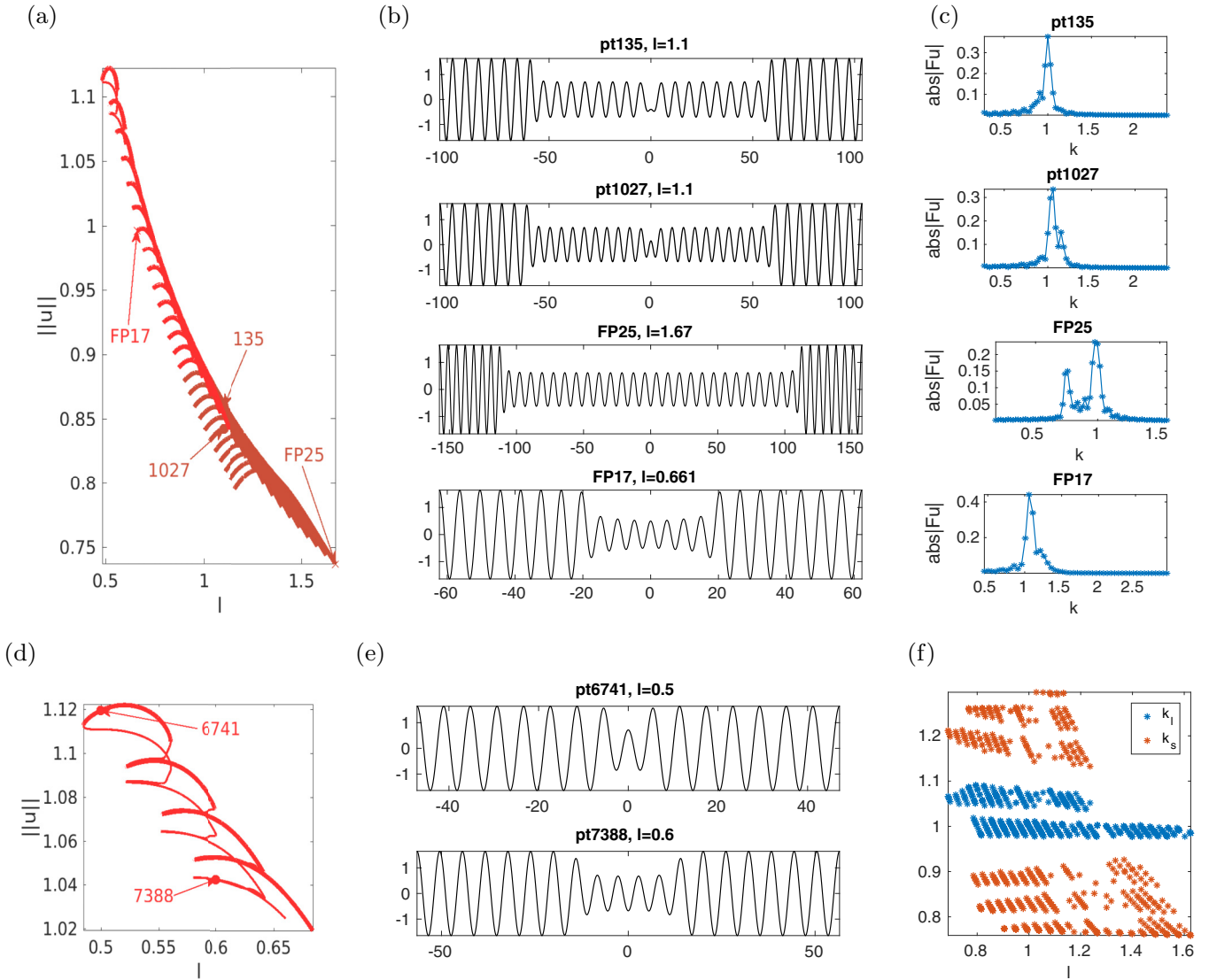


FIG. 5. (a) Details of the $u_p^{\text{large}} - u_p^{\text{small}} - u_p^{\text{large}}$ snake from Fig. 4. (b, c) Sample solutions and Fourier transforms. (d, e) Left end of the snake and sample solutions. (f) Wave numbers of u_p^{small} (red) and u_p^{large} (blue) from the snake.

and altogether we obtain a closed loop of $u_p^{\text{large}}(\cdot; \ell)$ solutions [upper black branch in Fig. 4(a)]. The bifurcation from u_p^{large} at ℓ_0 again leads to amplitude modulation at the right domain boundary [Fig. 4(c), middle], but in contrast to the u_p^{small} case, the bifurcating branch exhibits two folds near $\ell = 1$ before reconnecting to u_p^{large} at the left Eckhaus boundary in ℓ . See the first two plots in (c) for sample solutions.

Additionally, Fig. 4(a) shows a snaking branch (red) of $u_p^{\text{large}} - u_p^{\text{small}} - u_p^{\text{large}}$ heteroclinic cycles, starting at $\ell = 1$ (pt0) with a sample profile shown on the bottom right of (c). This snaking in ℓ is shown in more detail in Fig. 5. Starting at pt0 and increasing ℓ the solutions remain stable up until close to the first fold near $\ell = 1.18$. Near this fold a phase slip is initiated. This phase slip is once again associated with the onset of spatial modulation. This time it is located in the middle of the domain and manifests itself in the splitting of the central peak of the u_p^{small} portion of the profile [Fig. 5(b)]. Beyond the fold the new central peak regrows to the u_p^{small}

amplitude but the solutions are now unstable, and only recover stability at the next fold on the left. The net effect of this process is to add half a wavelength of the u_p^{small} state to the solution profile. This process then repeats adding a full wavelength of u_p^{small} in the middle of the domain after every two folds, and leading to slanted snaking with increasing ℓ . It is clear that these repeated phase slips compress the u_p^{small} portion of the solution, although the wavelength of u_p^{large} also needs to adapt, albeit only slightly. See the first three plots in Fig. 5(b) for sample profiles. This process continues indefinitely.

If, however, we start at pt0 and decrease ℓ , then we can extend the snake to the left, leading to a shrinkage of the middle section of u_p^{small} [see the last plot in Fig. 5(b) for a sample profile]. Figure 5(d) shows that this shrinkage is in all cases accompanied by significant hysteresis. Figures 5(d) and 5(e) show how the above process changes at the left end of the snake. At the leftmost fold [Fig. 5(d)], the amplitude of the last

remaining u_p^{small} peak does *not* increase to the u_p^{large} amplitude. Instead the solution grows new u_p^{small} peaks on either side of the central peak thereby initiating a parallel snake similar to that just described, but with all solutions unstable.

To see the wave-number adaption in a more quantitative way, in Fig. 5(c) we plot $|\hat{u}(k)|$ [the discrete Fourier transform of the sample solutions from Fig. 5(b)]. The larger peak, associated to u_p^{large} , stays near $k = 1$ throughout the snake, while the smaller peak, associated to u_p^{small} shifts. In Fig. 5(f) we summarize the wave numbers k_s of the small peaks (red) and k_l of the large peaks (blue) obtained from solutions in the snake. For this we only use solutions for which there is a clear peak separation in Fourier space, and thus solutions with $k_s \approx k_l$ are discarded. The plot shows that k_l adapts much less than k_s . While some scatter due to the limited resolution in Fourier space is present, bands in the wave numbers corresponding to different forward and backward transitions in the snake are clearly visible. Because of the ability of both u_p^{small} and u_p^{large} states to absorb the wavelength change generated by the phase slips in the center of the domain we conjecture that this type of double structure is more robust with respect to wavelength change than structures that are more rigid.

The above scenario is reminiscent of defect-mediated snaking [35], with two important differences. In defect-mediated snaking the solution branch appears from a fold on the branch of homogeneous states and so consists of a single branch. Moreover, the snaking is not slanted, because the phase slips do not result in the compression of a second portion of the solution. It is this compression that is in our case responsible for the slanted nature of Fig. 5(a) since it turns the problem into an effectively nonlocal one.

C. The case $a = 9.5$: Breakup of snakes into stacks of isolas and a multitude of cycles

Larger values of a result in more and more intersections among H for periodic branches with different k , and the homogeneous branch $k = 0$ [see Fig. 6(a) for $a = 9.5$], and thus many more heteroclinic cycles become possible. Up to $a = 9.3$ (say, and depending on the domain size), the basic snake-and-ladders structure from Fig. 3 stays intact, but for larger a the snakes and rungs reconnect into a stack of isolas. Moreover, $u_{\text{hom}}^{\text{large}}$ plays an increasing role in the continuation of the solutions. Figures 6(c) and 6(d) illustrate these effects. The red branch bifurcates from BP1 on $u_{p,1}$, but fails to grow a front between $u_{p,1}^{\text{large}}$ and $u_{p,1}^{\text{small}}$ as it would in “classical” snaking at lower a . Instead, it now exhibits long, nearly vertical intervals located near $\lambda = 3.5$, associated first with the growth of a segment of $-u_{\text{hom}}^{\text{large}}$ in the solution profile and then its shrinkage before another half period of u_p^{large} can be inserted. The branch bifurcating at BP2 on u_p^{small} , which at lower a generates a snaking branch of $u_p^{\text{small}} - u_p^{\text{large}} - u_p^{\text{small}}$ heteroclinic cycles, cf. Fig. 3, behaves similarly and also always involves plateaus of $\pm u_{\text{hom}}^{\text{large}}$.

This type of behavior is similar to that recently found in a Gray-Scott model studied in connection with dryland vegetation patterns where tristability between a pair of different spatially periodic states and a homogeneous state is also

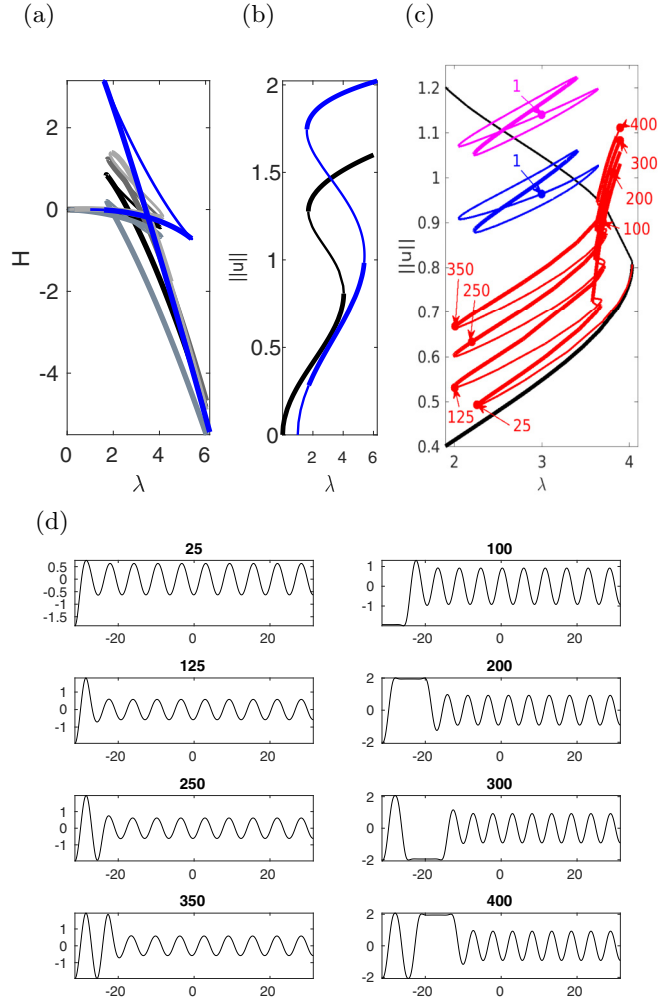


FIG. 6. $a = 9.5$, $\Omega = (-10\pi, 10\pi)$. (a) H for solution branches plotted as in Fig. 3(a); there are now multiple intersections near $\lambda = 3.5$. (b) $\|u\|$ for the $k = 1$ and $k = 0$ branches, illustrating the multistability near $\lambda = 3.5$. (c) Two isolas of cycles between $u_{p,1}^{\text{small}}$ and $u_{p,1}^{\text{large}}$ obtained from suitable initial guesses, further discussed in Fig. 7, and a branch (red) of $u_{p,1}^{\text{large}} - u_{p,1}^{\text{small}}$ cycles which passes near $\lambda = 3.5$ close to $\pm u_{\text{hom}}^{\text{large}}$. (d) Sample solutions from the red branch in (c).

present [36], suggesting that the behavior shown in Fig. 6(c) is in fact generic.

On the other hand, we can still generate “classical” heteroclinic cycles between u_p^{small} and u_p^{large} leading to, e.g., the blue and magenta isolas in Fig. 6(c). To generate starting points for these we glue together $u_{p,1}^{\text{small}}$, $u_{p,1}^{\text{large}}$ segments (using a longer middle segment for the magenta isola) and running a Newton loop to converge to a solution. Figure 7 shows details of the reorganization of the segments that make up the snakes and rungs at smaller a into the isola structure.

Figure 6(d) suggests that for $a = 9.5$ we can expect further cycles, not present at small a , where the $u_{\text{hom}}^{\text{large}}$ state plays a prominent role. Figure 8 provides some examples. The red branch in Fig. 6(a), with sample solutions in Fig. 6(b), corresponds to $u_p^{\text{large}} - u_{\text{hom}}^{\text{large}}$ heteroclinics. In states of this type it is difficult to eliminate a wavelength of the periodic

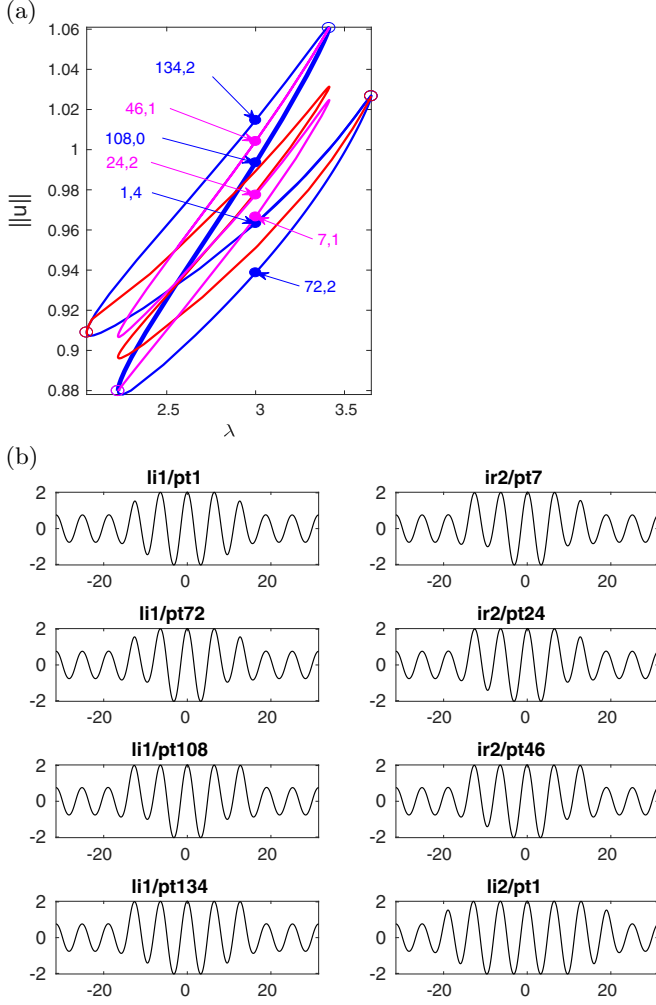


FIG. 7. (a) Zoom of the blue isola in Fig. 6(c) and associated rung structure. The labels ptr,i indicate stability properties with i specifying the number of unstable eigenvalues. (b) Sample solutions from (a), and the initial solution on $li2$ from Fig. 6(c).

state in response to parameter changes—it costs less energy to compress the state and expand the homogeneous state or vice versa. Thus phase slips will not be triggered unless the wavelength of the periodic state changes by a substantial amount. Solutions on the lower part of this branch (with $pt176$) include short loops near u_p^{small} and are unstable. Figures 6(c) and 6(d) show more heteroclinic cycles, indicating that at large a (and on sufficiently large domains) all sorts of connections are possible provided one selects λ values corresponding to intersections of H for the pertinent patterns. We used initial guesses of the form “pattern-large-to-homogeneous-small” (pl2hs, red), “pattern-small-to-homogeneous-small” (ps2hs, green), “pattern-small-to-homogeneous-large” (ps2hl, orange) to converge to the corresponding solutions and continued the resulting solutions to $\lambda \approx 3.5$, where they all form localized patterns consisting of (on this domain) up to four patches of different solutions (u_{hom}^{small} , u_{hom}^{large} , u_p^{small} , and u_p^{large}), many of which are stable, albeit in rather narrow λ intervals. These stability intervals are a consequence of what appears to be collapsed snaking of the corresponding branches in the vicinity of the Maxwell point at $\lambda \approx 3.5$. In states of this type

changes in the wavelength of the stripe portion of the solution are readily accommodated by changes in the homogeneous portion.

The reconnection of the snaking diagram into a stack of isolas as parameters are varied has been seen in SH23 [10,12] and in two-dimensional patterns may even occur as one proceeds up a snaking diagram, all other parameters remaining fixed [37,38].

III. DISCUSSION

In suitable parameter regimes, the SH357 equation allows many different heteroclinic cycles between four (recall that we identify $\pm u_p^{small}$, $\pm u_p^{large}$, $\pm u_{hom}^{small}$ and $\pm u_{hom}^{large}$, respectively) main building blocks, namely,

(1) u_p^{small} and u_p^{large} : periodic patterns of small and large amplitude, respectively, with wave numbers k near 1.

(2) u_{hom}^{small} and u_{hom}^{large} : spatially homogeneous states of small and large amplitude; these can be seen as special cases of u_p^{small} and u_p^{large} with wave number $k = 0$. However, from the point of view of heteroclinic cycles, the homogeneous states are special in the sense that segments of u_{hom}^{small} and u_{hom}^{large} can have arbitrary length.

To give some structure to our results, we made the special choice Eq. (3), i.e., $b = 3.5 + 0.4(a - 3)$. For relatively small a ($a = 2$ in Sec. II A), we have a more or less simple situation in the sense that the necessary condition,

$$H[u_1(\cdot; \lambda)] = H[u_2(\cdot; \lambda)], \quad (8)$$

for the patterns involved in a heteroclinic cycle only holds for a relatively small number of patterns. Moreover, these values of a allow for an interesting continuous transition between heteroclinic cycles between u_p^{large} and u_p^{small} , and heteroclinic cycles between u_p^{large} and $u \equiv 0$. For intermediate a ($a = 5$ in Sec. II B), we obtain “classical” snaking of heteroclinic cycles between u_p^{large} and u_p^{small} . Via continuation in the domain size we also found a new kind of slanted snaking (reminiscent of defect-mediated snaking), where, e.g., the u_p^{small} portion of the solution grows or shrinks via phase slips. Finally, for “large” a , the $u_p^{small} - u_p^{large} - u_p^{small}$ snaking breaks up into stacks of isolas. Moreover, Eq. (8) is fulfilled for a large number of different patterns, and consequently many different heteroclinic cycles become possible. In Sec. II C, we provide several examples of such states for $a = 9.5$.

In our bifurcation diagrams we indicated linearly stable (unstable) solutions by thick (thin) lines. Owing to the generally large number of different simultaneously stable solutions, periodic and localized, it would be desirable to characterize in addition the different basins of attraction. However, such basins are not yet well understood even for standard homoclinic snaking as described by SH23 or SH35. Therefore, here we confine ourselves to a few remarks: The snakes and related structures such as stacks of isolas typically have rather large basins. For instance the starting points for the computation of the isolas in Fig. 6 can also be obtained from quite rough initial guesses followed by time integration towards a stable solution. Of course, different initial guesses may lead to different but “nearby” steady states. Put differently, a localized $O(1)$ perturbation of a solution u^* in a snake typically leads to rather

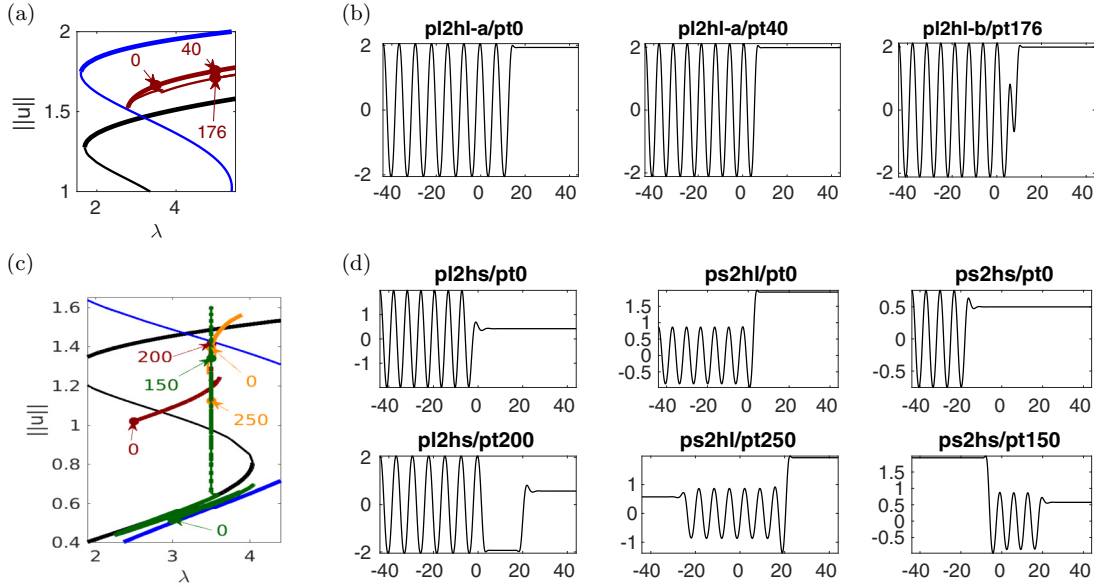


FIG. 8. Various localized patterns at $a = 9.5$, $\Omega = (-14\pi, 14\pi)$, with starting points for continuation obtained from running a Newton loop on rough initial guesses, for instance gluing together a large pattern segment and the large homogeneous solution. (a, b) Bifurcation diagram and sample plots from the red branch p12hl (“pattern-large-to-homogeneous-large”) in (a). (c) Bifurcation diagram of branches p12hs (red), ps2h1 (green), and ps2h1 (orange). (d) Sample solutions from (c).

fast relaxation either back to u^* , or to a nearby solution in the snake, with, say, one more “large” roll and one fewer “small” roll. This behavior naturally changes if we go outside the snaking region, or consider delocalized perturbations, which in particular may lead to depinning, as illustrated next.

The states satisfying Eq. (8) correspond to time-independent structures (fronts, pulses etc.) and the presence of snaking associated with these connections implies that such structures remain stationary even away from the Maxwell point, i.e., when $|\mathcal{E}[u_1(\cdot; \lambda)] - \mathcal{E}[u_2(\cdot; \lambda)]| > 0$. If, however, the energy difference between the two competing states becomes too large, then the front connecting them depins and the lower energy state invades the higher energy state in a process analogous to depinning of fronts in SH23 or SH35 [10,29]. Figure 9 shows examples of this process, focusing [Figs. 9(a)–9(d)] on competition between states that are both spatially periodic, and illustrating possibilities that do not arise in either SH23 or SH35, namely, front propagation via repeated phase slips. The fact that phase slips in SH357 can propagate is of particular interest since the Eckhaus instability that triggers them is a steady state instability. Since phase slips require a finite time for proceed to completion we may anticipate the presence of a dynamic regime in which the speed of the front is determined by the phase-slip timescale and not the energy difference alone [39]. In Fig. 9(a), with $\lambda = 1.3$ in the snaking region of Fig. 3, these phase slips adjust the wavelength of the small amplitude portion, but the front between u_p^{large} and the u_p^{small} resulting from these phase slips remains pinned. In Fig. 9(b) we have two fronts: a fast front consisting of propagating phase slips in u_p^{small} , followed by a slower front whereby the higher amplitude but lower energy state invades the smaller amplitude higher energy state. The speed of this amplitude front is not constant, however, and is strongly affected by the phase slips on the u_p^{small} portion. We conjecture that this is

a consequence of the fact that the u_p^{small} wave number has still not relaxed to its equilibrium value when the amplitude front arrives. In Fig. 9(c), corresponding to a slightly larger λ than in Fig. 9(b), the amplitude front triggers further phase slips, locally accelerating the front and resulting in a type of stick-slip motion at large times. This does not happen in case (b) even at very long times. Finally, in Fig. 9(d) the large amplitude part of the initial condition is dilated relative to its equilibrium wavelength and in this case the system relaxes to a lower energy state via phase diffusion in both u_p^{small} and u_p^{large} , instead of phase slips. Here the front remains pinned but its location adjusts accordingly.

Figure 9(e) shows a multifront at larger a . Here we obtain a $u_p^{\text{small}} - (-u_{\text{hom}}^{\text{small}}) - u_{\text{hom}}^{\text{large}}$ double front with retreating $u_{\text{hom}}^{\text{large}}$, obtained from a step-like initial condition, but as suggested by Fig. 8, all sorts of multifronts are possible via appropriate choice of initial conditions. Fronts between homogeneous states are then generically fast, while fronts involving patterned states are typically substantially slower.

Equations of the 357 form have been considered before, in context of the Ginzburg-Landau equation with real coefficients [40]. When parametrically forced by a spatially periodic function the homogeneous solutions of this equation become periodic solutions with wavelength equal to the forcing wavelength. The resulting equation thus also exhibits coexistence between different amplitude periodic states, and between periodic states and the homogeneous state. While this equation also reveals the gradual breakup of forced snaking into isolas [41] the situation is different since the wavelength of the periodic states is imposed by the forcing wavelength with the result that bistability between periodic states with different wavelengths is absent. Nevertheless, models of this type indicate that the phenomena described here may also occur in periodically forced systems exhibiting periodic states

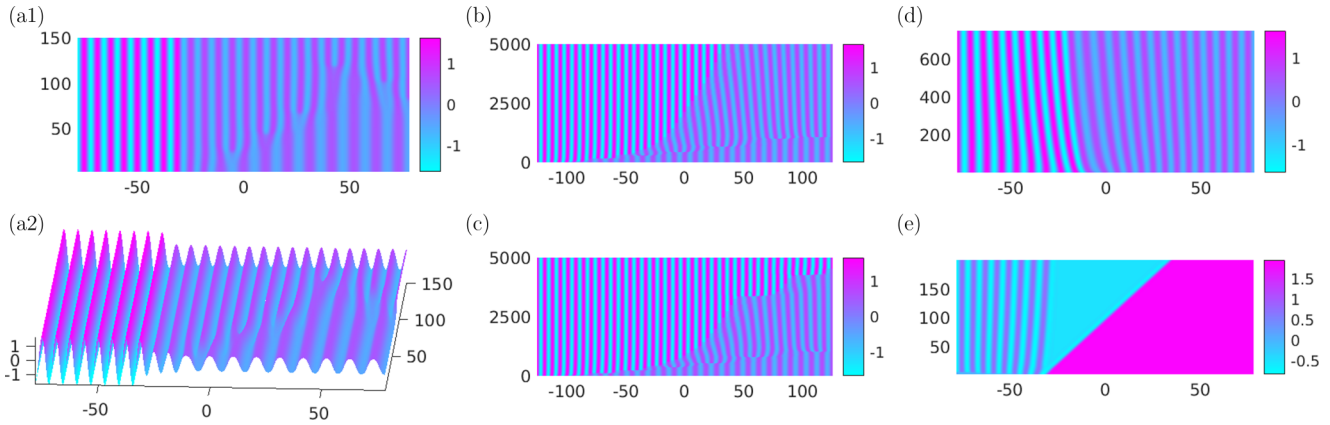


FIG. 9. Time evolution on $\Omega = (-25\pi, 25\pi)$ (a, d, e) and $\Omega = (-40\pi, 40\pi)$ (b, c) in the form of space-time diagrams. Horizontal axis shows space x , vertical axis time t , parameters as indicated. Initial conditions u_0 consist of smooth amplitude and wave-number transitions between one state on the left and a different state on the right (the precise form appears not to matter provided it is not too gradual). (a) $(a, \lambda) = (5, 1.3)$, $u_0 = 1.5 \cos(x)$ for $x < -9\pi$, $u_0(x) = 0.5 \cos(0.6x)$ for $x \geq -5\pi$; (a2) shows the same solution as (a1) as a 3D plot. (b) $(a, \lambda) = (5, 1.49)$, $u_0 = 1.5 \cos(x)$ for $x < -34\pi$, $u_0(x) = 0.5 \cos(0.6x)$ for $x \geq -28\pi$. (c) $(a, \lambda) = (5, 1.5)$, initial condition as in (b). (d) $(a, \lambda) = (5, 1.3)$, $u_0 = 1.5 \cos(0.8x)$ for $x < -4\pi$, $u_0(x) = 0.5 \cos(x)$ for $x \geq 0$. (e) $(a, \lambda) = (9.5, 3)$, $u_0 = 0$ for $x < -10\pi$, $u_0 = 2$ for $x \geq -10\pi$.

with an intrinsic wavelength. Such systems remain to be studied in detail, although preliminary studies of the spatially forced Swift-Hohenberg equations SH23 and SH35 indicate that the snaking behavior is likewise destroyed as the forcing amplitude increases [42].

In summary, the SH357 equation is an extremely rich pattern-forming system and may be seen as a one-dimensional model for studying intricate structures in systems exhibiting competition between states with distinct wavelengths. The gradient structure of this equation, and in particular the existence of the spatial invariant H , allow a more detailed understanding of the behavior of this equation than is possible for other equations exhibiting similar behavior, such as the Gray-Scott model [36], but neither property is essential for the behavior described here as is well documented for standard homoclinic snaking [25]. In particular, standard homoclinic snaking, as described by SH23 or SH35, is robust with respect

to both parameter changes and boundary conditions,¹ and for these reasons is found in more complex systems including the equations of hydrodynamics.

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¹In the presence of nonperiodic or non-Neumann boundary conditions an extended periodic state is absent and the snaking localized states turn continuously into a spatially extended state satisfying the boundary conditions [43].

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