


Non-Hermitian Band Topology and Skin Modes in Active Elastic Media

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Solids built out of active components can exhibit nonreciprocal elastic coefficients that give rise to non-Hermitian wave phenomena. Here, we investigate non-Hermitian effects present at the boundary of two-dimensional active elastic media obeying two general assumptions: their microscopic forces conserve linear momentum and arise only from static deformations. Using continuum equations, we demonstrate the existence of the non-Hermitian skin effect in which the boundary hosts an extensive number of localized modes. Furthermore, lattice models reveal non-Hermitian topological transitions mediated by exceptional rings driven by the activity level of individual bonds.

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The microscopic injection of energy into solid media via active, living, or robotic components fundamentally alters their mechanical waves [1–7]. As with optics [8,9], topoelectric circuits [10–13], and open quantum systems [14,15], the interplay between activity (gain) and dissipation (loss) can often be captured by non-Hermitian operators [16–18]. In all these contexts, a central question is the role of non-Hermiticity at the boundary of the system. Like their Hermitian counterparts, non-Hermitian systems have been shown to exhibit topological invariants that ensure localized boundary modes [19–32]. However, in some cases, the familiar bulk-boundary correspondence breaks down for non-Hermitian systems. Such systems exhibit the non-Hermitian skin effect, in which an extensive number of modes are localized to the system’s boundary [33–35].

Here we examine the non-Hermitian wave phenomena that arise from the elastic properties of a class of active solids. In the continuum, non-Hermiticity enters the linear elasticity of a solid through odd elastic moduli, which are active moduli that violate Maxwell-Betti reciprocity [7]. We show that such odd elastic moduli when combined with anisotropy can give rise to the non-Hermitian skin effect. This effect implies a dramatic localization of vibrational modes to the system’s boundary. Furthermore, we take a microscopic view of elasticity by considering 2D lattices composed of active bonds. These bonds, while active, retain two crucial features of Hookean springs: they conserve linear momentum and depend only on changes in relative distance. We uncover a non-Hermitian topological transition driven by the level of activity. This transition differs qualitatively from its Hermitian counterpart in that it is mediated by exceptional rings. Such rings arise due to geometric changes in the particle trajectories that enable the system to draw energy from nonpotential

forces. We interpret the resulting energy cycles in terms of a generalized PT symmetry.

Non-Hermitian elasticity.—We choose as our starting point elasticity theory, the continuum description of solids that captures their ability to resist shape change at large length scales [36]. Unlike conventional treatments of passive elasticity, we seek to capture at a coarse-grained level the effects induced by *nonconservative* internal forces $F_i(\mathbf{x})$ satisfying three assumptions [7]. First, the forces conserve linear momentum, and therefore can be written as the divergence of a stress $\partial_j \sigma_{ij}(\mathbf{x})$. Second, the forces only depend on the static change in shape, which is captured by gradients of the displacement field $u_{ij} = \partial_i u_j(\mathbf{x})$. Finally, we employ the phenomenological assumption that the stresses can be approximated as linearly proportional to the displacement gradients: $\sigma_{ij} = C_{ijmn} u_{mn}$. The object C_{ijmn} is the elastic tensor, and it encodes the material’s response to static deformation.

Following the approach of Ref. [7], it is useful to express the elastic tensor as the sum of two pieces:

$$C_{ijmn} = C_{ijmn}^e + C_{ijmn}^o, \quad (1)$$

where $C_{ijmn}^e = C_{mni j}^e$ is even, or symmetric, under exchange of pairs of the lower indices while $C_{ijmn}^o = -C_{mni j}^o$ is odd, or antisymmetric [7]. To understand the decomposition, consider the elastic work done (per unit volume) by a patch of material brought through a closed cycle of strain: $w = -\oint \sigma_{ij} du_{ij} = C_{ijmn}^o \oint u_{ij} du_{mn}$. If the system is passive, $w = 0$ for any cycle that begins and ends in the same state, and hence $C_{ijmn}^o = 0$. However, for an active solid, this requirement does not hold and, consequently, C_{ijmn}^o may be nonzero. We use the term “odd elastic media” to refer to this class of active systems [7].

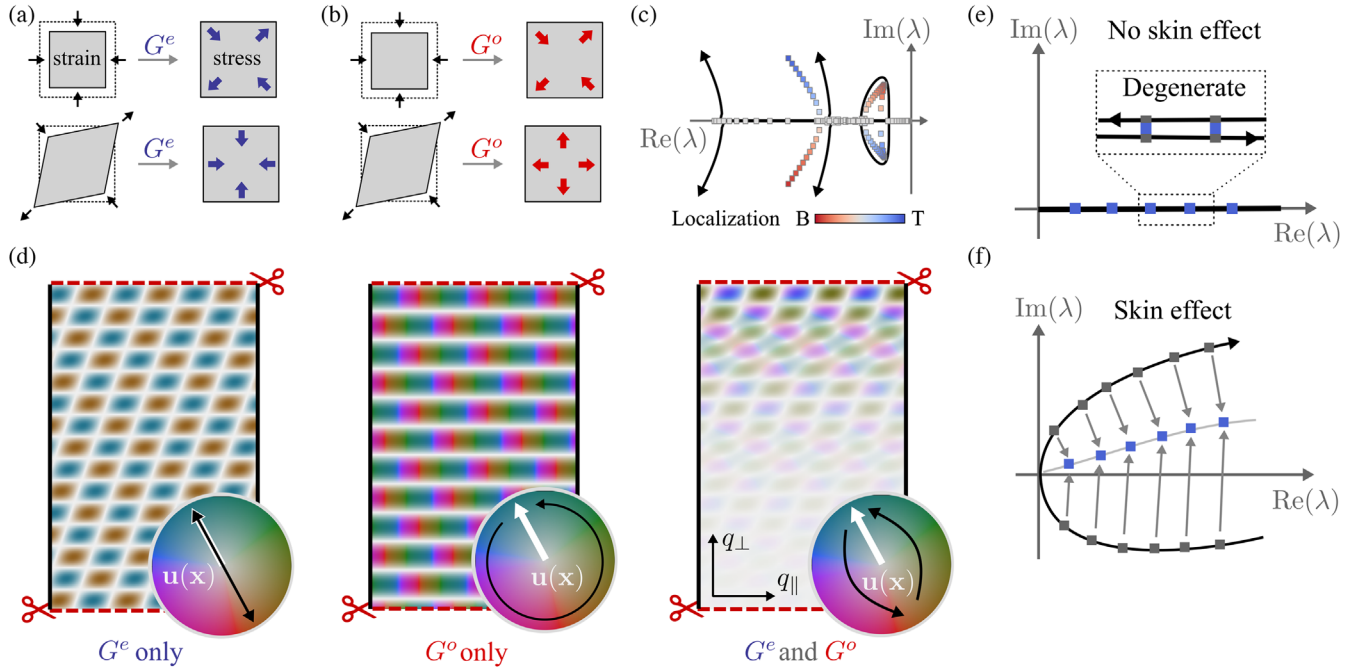


FIG. 1. Elastic non-Hermitian skin effect. (a) An anisotropic, passive elastic modulus G^e that couples dilation and shear. (b) The corresponding anisotropic, odd elastic counterpart G^o . (c) The elastic spectrum for $G^o/G^e = 1.7$. The black arrows are analytical calculations of the periodic boundary spectrum, see [41]. The colored squares denote numerical calculations of the spectrum with open boundary conditions. Blue (red) indicates localization to the top (bottom) boundary. (d) Numerically computed eigenmodes with only G^e present (left), only G^o present (middle), and both G^o and G^o (right). Hue indicates the angle and opacity indicates the magnitude of the displacement field $\mathbf{u}(\mathbf{x})$. The horizontal boundaries (red) are open and the vertical boundaries (black) are periodic. The boundary termination forms an angle $\pi/4$ with respect to the axis of anisotropy. (e) Since the spectrum of a Hermitian system lies on the real line, a generic eigenvalue is at least doubly degenerate. (f) For a non-Hermitian system, the spectrum can trace out nondegenerate arcs (black solid). In this case, the spectrum deforms (blue squares) to form degeneracies when a boundary is introduced.

Suitable experimental platforms include robotic metamaterials [2,5], solids with integrated piezoelectric components [37,38], and chiral optical matter [39–41].

Here, we focus on *anisotropic* odd-elastic media and illustrate how they generically exhibit the non-Hermitian skin effect. For concreteness, consider a minimal example of anisotropic odd elasticity represented by the following pictorial stress-strain relationship (see the Supplemental Material for standard tensor notation [41]):

$$\begin{pmatrix} \text{⊕} \\ \text{⊗} \\ \text{⊗} \\ \text{⊕} \end{pmatrix} = \begin{pmatrix} 0 & 0 & G^e + G^o \\ 0 & \mu & 0 \\ G^e - G^o & 0 & \mu \end{pmatrix} \begin{pmatrix} \text{⊖} \\ \text{⊖} \\ \text{⊖} \end{pmatrix}. \quad (2)$$

The modulus G^e (contained in C_{ijmn}^e) couples dilation (⊖) to shear stress (⊗) and shear strain (⊖) to pressure (⊕) symmetrically, see Fig. 1(a). By contrast, the modulus G^o (contained in C_{ijmn}^o) provides an antisymmetric coupling [Fig. 1(b)]. Equation (2) also includes the standard shear modulus μ to ensure mechanical stability. In the presence of these moduli, the expression for elastic forces may be written as $\mathbf{F}(\mathbf{x}) = \hat{D}\mathbf{u}(\mathbf{x})$, where \hat{D} takes the following form:

$$\hat{D} = \begin{pmatrix} \mu \nabla^2 + 2G^e \partial_x \partial_y & G^e \nabla^2 + G^o [\partial_x^2 - \partial_y^2] \\ G^e \nabla^2 - G^o [\partial_x^2 - \partial_y^2] & \mu \nabla^2 + 2G^e \partial_x \partial_y \end{pmatrix}, \quad (3)$$

where $\nabla^2 = \partial_x^2 + \partial_y^2$. In Eq. (3) we observe that the operator \hat{D} becomes non-Hermitian when G^o is nonzero, i.e., when the anisotropic moduli display an odd component arising from nonconservative elastic forces. We find that the non-Hermiticity has a dramatic effect on the nature of the bulk modes. In Fig. 1(c), we show the spectrum of \hat{D} as a function of wave number q_y for fixed q_x when both G^e and G^o are present. We find two striking features. First, the open boundary spectrum (square markers) differs dramatically from the spectrum with periodic boundaries (black lines). Second, when we examine a typical eigenmode [Fig. 1(d), right] we find that the mode is exponentially localized to the open edge. In Fig. 1(c), we color each mode by the degree of localization to the top (blue) and bottom (red) boundaries. In contrast to typical topological waves or Rayleigh waves in Hermitian systems [36], an extensive number of modes are localized to the boundary. This extensive localization of bulk modes is an elastic manifestation of the non-Hermitian skin effect.

Yet, when either $G^o = 0$ or $G^e = 0$, the skin effect disappears [Fig. 1(d), left and center, respectively]. To gain insight into its origins, we consider the notion of a generalized Brillouin zone [33–35]. Let q_{\parallel} (q_{\perp}) be the wave number parallel (perpendicular) to the boundary. For a finite system, at least two Bloch modes must have the same eigenvalue λ and wave number q_{\parallel} in order to interfere to satisfy a given boundary condition, e.g., $\mathbf{u} = 0$. If \hat{D} is Hermitian or anti-Hermitian, this condition is generically satisfied since the spectrum is confined to lie entirely along the real or imaginary line [Fig. 1(e)]. However, when both C_{ijmn}^e and C_{ijmn}^o are nonzero, the spectrum can inhabit the full complex plane. Consequently, the spectrum, plotted as a function of q_{\perp} , need not retrace itself [Fig. 1(f)]. In this case, there exist segments in which no two extended Bloch modes have the same λ and q_{\parallel} . Nonetheless, one can consider Bloch modes with complex wave numbers $\tilde{q}_{\perp} = q_{\perp} + ik(q_{\perp})$. By analytically continuing \hat{D} into the complex Brillouin zone, the eigenvalues flow together to enable interference at the boundary.

Finally, we note that the use of the two specific moduli G^e and G^o in Eq. (2) is purely illustrative. More generally, two necessary conditions must be met in order for the non-Hermitian skin effect to emerge within the continuum description of an elastic medium. First, both C_{ijmn}^e and C_{ijmn}^o must be nonzero for the spectrum to occupy the complex plane. Second, the system must be anisotropic in order for the spectrum not have a reflection symmetry over the system's boundary ($q_{\perp} \mapsto -q_{\perp}$), see the Supplemental Material [41]. For the example in Eq. (2), both C_{ijmn}^e and C_{ijmn}^o are independently anisotropic. In the Supplemental Material we examine a case in which only C_{ijmn}^e is anisotropic. Finally, we note that \hat{D} has inversion symmetry, which implies that the skin modes occur in pairs localized to each boundary [12]. This is a contrast to systems in which an external medium enables an effective violation of Newton's third law [4–6]. The continuum limit of these systems (whose interactions do not conserve linear momentum) yields $\hat{D} \propto q$, rather than the $\hat{D} \propto q^2$ dependence characteristic of elasticity [41].

Microscopic model.—A ubiquitous minimal model for elastic solids is a collection of masses connected by Hookean springs [53–66]. The Hookean spring captures two generic features of elasticity. First, the interaction conserves linear momentum, since the forces on the two participating particles are equal and opposite. Second, the force only depends on the change in bond length. Hence, the emerging mechanical response will be sensitive only to intrinsic changes in geometry. Yet, the Hookean spring has an additional feature built in: its force law follows from the gradient of a potential. Here, we retain the assumptions of length dependence and linear momentum conservation, and we study the most general 2D linear pairwise interaction when only energy conservation is lifted [7]:

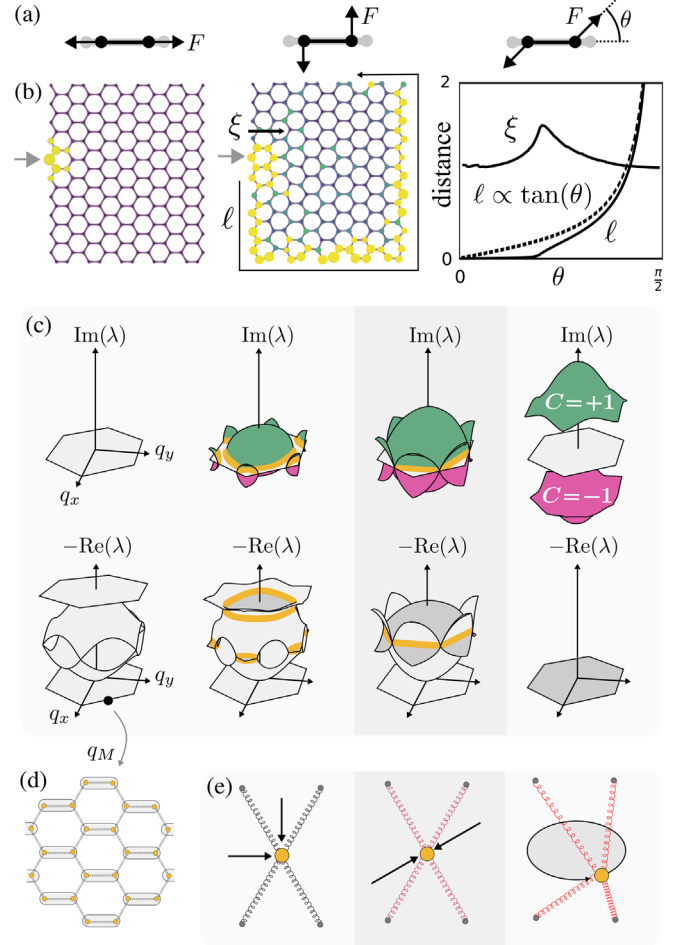


FIG. 2. Non-Hermitian topological transition and exceptional rings. (a) The generalized Hookean spring with the force \mathbf{F} oriented at an angle θ with respect to the bond vector. (b) Simulations with $\theta = 0$ (left) and $\theta = \pi/2$ (right) in which a particle at the edge is vibrated. Two lengths, the penetration depth ξ and the propagation distance ℓ , are plotted as a function of θ . (c) The spectrum plotted over the Brillouin zone for $\theta = 0, \pi/12, \pi/6, \pi/2$. Regions of positive (green) and negative (purple) Berry curvature are highlighted. The orange lines denote exceptional rings. (d) At point \mathbf{q}_M , pairs of masses move in tandem. (e) The eigenmodes (arrows) of the effective single particle system at $\theta = 0, \pi/6, \pi/2$.

$$\mathbf{F}(r) = -(k\hat{\mathbf{r}} + k^a\hat{\boldsymbol{\phi}})\delta r, \quad (4)$$

where $\hat{\mathbf{r}}$ ($\hat{\boldsymbol{\phi}}$) is a unit vector pointing along (transverse to) the bond vector, δr is the change in length of the bond, and k and k^a are spring constants [see Fig. 2(a)].

When the bond is taken on a closed cycle, the work done $W = \oint \mathbf{F} \cdot d\mathbf{r}$ is equal to k^a times the area enclosed by the path. Hence, when $k^a \neq 0$, Eq. (4) cannot be derived from a potential. In principle, Eq. (4) can be paired with any form of dynamics that governs the temporal evolution of the system. Here, for concreteness, we will interpret our results in the context of an overdamped equation of motion:

$\Gamma \partial_t \mathbf{u} = \mathbf{F}$, where \mathbf{u} is the displacement of the particle and Γ is a drag coefficient. By adjusting the angle $\theta = \arctan(k^a/k)$ between $\hat{\mathbf{r}}$ and \mathbf{F} , we can interpolate between longitudinal and transverse interactions [67–76]. In the Supplemental Material [41], we show how the equations governing the *overdamped* dynamics of an active solid with *nonconservative* bonds described by Eq. (4) are the same as those governing the *inertial* dynamics of a gyroscopic metamaterial [77–81] with *conservative* springlike interactions and weak dissipation. However, we note that deformation cycles performed with the gyroscopic media do not extract energy since the left-hand side of Eq. (4) represents torques, not forces. Similarly, in the continuum treatment of gyroscopes, the left-hand side of Eq. (2) represents angular momentum currents, not stresses.

Generalized PT symmetry and energy cycles.—For a generic network of masses connected by the bonds in Eq. (4), the linear relationship between forces $\mathbf{F}(\mathbf{x})$ and displacements $\mathbf{u}(\mathbf{x})$ can be captured by a dynamical matrix formalism $\mathbf{F}(\mathbf{x}) = \sum_{\mathbf{x}'} D(\mathbf{x}, \mathbf{x}') \mathbf{u}(\mathbf{x}')$, where \mathbf{x} and \mathbf{x}' are lattice sites and $D(\mathbf{x}, \mathbf{x}')$ is the dynamical matrix. The mere fact that the forces and displacements are real implies that $[\mathcal{K}, D(\mathbf{x}, \mathbf{x}')] = 0$, where \mathcal{K} is complex conjugation. Since \mathcal{K} is an antiunitary operator with $\mathcal{K}^2 = 1$, we say that the dynamical matrix has a generalized *PT* symmetry [17,82–84], see [41].

The *PT* symmetry has the following physical consequence: if a given eigenvalue λ of D is real, then the corresponding eigenvector $\mathbf{u}_\lambda(\mathbf{x})$ may be chosen real. Since $\mathbf{u}_\lambda(\mathbf{x})$ is real, the corresponding trajectory of each particle traces out straight lines in time [Fig. 2(e), left]. Moreover, nonreal eigenvalues come in complex conjugate pairs $\lambda_\pm = \lambda_R \pm i\lambda_I$ with eigenvectors of the form:

$$\mathbf{u}_{\lambda_\pm}(\mathbf{x}) = \mathbf{v}(\mathbf{x}) \pm i\mathbf{w}(\mathbf{x}), \quad (5)$$

where $\mathbf{v}(\mathbf{x})$ and $\mathbf{w}(\mathbf{x})$ are real vectors. Physically, a complex eigenvalue indicates energetic gain or loss. In this case, the eigenmode cycles between two states $\mathbf{v}(\mathbf{x})$ and $\mathbf{w}(\mathbf{x})$. Since the bonds are nonpotential, the cycles result in the injection (or removal) of energy [Fig. 2(e), right]. If all the eigenvalues of D are real, we say that D is *PT* unbroken, and *PT* broken otherwise [85].

Non-Hermitian topological transition.—Given a microscopic model, we can study not only the acoustic bands (accessible within the continuum theory) but also features of the optical bands. Figure 2(b) shows the response of a honeycomb lattice to vibrations applied at the boundary in two extreme cases: the passive Hookean limit $\theta = 0$, and the active transverse limit $\theta = \pi/2$. In the transverse limit, we see the emergence of a sustained, unidirectional edge wave characteristic of a Chern insulator [77,78]. Due to the translation symmetry, we may express the dynamical matrix in terms of wave number \mathbf{q} [41]:

$$D_\theta(\mathbf{q}) = \cos(\theta)D_0(\mathbf{q}) + \sin(\theta)D_{\pi/2}(\mathbf{q}). \quad (6)$$

For Hermitian systems, nontrivial topology requires breaking time-reversal symmetry (TRS): $D_\theta(-\mathbf{q}) = D_\theta^*(\mathbf{q})$ [86]. However, here $D_\theta(\mathbf{q})$ naïvely obeys TRS for all θ since the forces and displacements are real quantities. Nonetheless, band topology is still possible due to the violation of Hermiticity. At $\theta = \pi/2$, the dynamical matrix $D_{\pi/2}(\mathbf{q})$ (restricted to its optical bands, see [41]) is anti-Hermitian. Hence the relevant Hermitian Hamiltonian $H(\mathbf{q}) = iD_{\pi/2}(\mathbf{q})$ violates TRS as required. When $\theta = \pi/2$, the lattice boundary hosts a chiral edge state due to the nonvanishing Chern numbers of the optical bands.

The transition between Hermitian and anti-Hermitian limits is accompanied by two length scales ξ and ℓ [Fig. 2(b), right]. The first length scale ξ is the penetration depth into the medium, which is set by the structure of the *eigenvectors* of D_θ . Like the Haldane model [41,87,88], ξ is roughly constant as the gap opens and closes. Yet, the edge modes do not become visible until large values of θ are probed. This effect can be traced to the second length scale ℓ , which is the distance the wave propagates around the edge. This length scale is set by the *eigenvalues* of the dynamical matrix. For a given mode, $\ell \approx \tau\omega/q$, where $\tau = -1/\text{Re}(\lambda)$ is the decay rate and $\omega = -\text{Im}(\lambda)$ is the oscillation frequency. Hence, the localized edge mode becomes apparent close to the anti-Hermitian limit, i.e., θ near $\pi/2$.

Mechanical exceptional points.—Insight into the transition is gained by examining the point \mathbf{q}_M in the Brillouin zone. At this point, the 4×4 dynamical matrix $D(\mathbf{q}_M)$ can be reduced to an effective dynamical matrix that governs the motion of a single particle in a trap [Figs. 2(d) and 2(e)]:

$$D_{\text{eff}} = - \begin{pmatrix} \cos \theta & -3 \sin \theta \\ \sin \theta & 3 \cos \theta \end{pmatrix}, \quad (7)$$

with eigenvalues $\lambda_\pm = -2 \cos \theta \pm \sqrt{2 \cos(2\theta) - 1}$, see [41]. For $\theta < \pi/6$, two real modes exist that trace out straight lines [Fig. 2(e), left]; for $\theta > \pi/6$, the eigenmodes trace out cycles (right). At the transition $\theta = \pi/6$, D_{eff} permits only a single eigenvalue $\lambda_- = \lambda_+$. However, the anisotropy of the trap and the chirality of the bonds imply that no two linear eigenmodes can have the same eigenvalue unless they are parallel. Hence, the two independent modes coalesce, indicating that the dynamical matrix is defective (i.e., nondiagonalizable). Such occurrences, known as “exceptional points,” are generic features of transitions between *PT*-broken and *PT*-unbroken phases [15,30,82,89–92]. Here, the exceptional points take on a clear physical meaning: they mark the crossovers between eigenmodes with linear motion, and eigenmodes with circular motion necessary to sustain active waves.

For the honeycomb lattice, the exceptional points do not merely occur at point \mathbf{q}_M . Rather, they occur along 1D rings

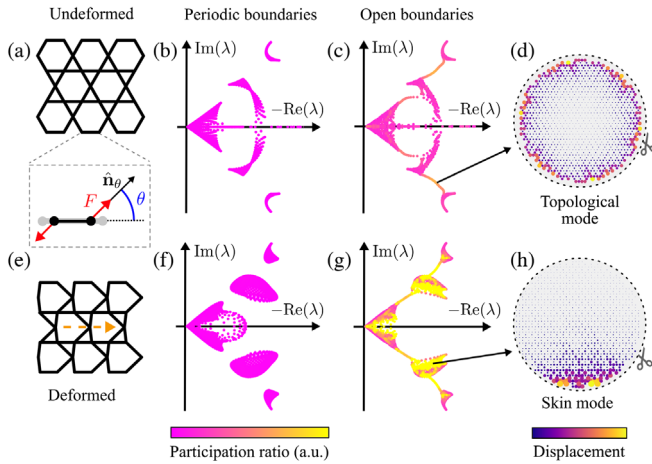


FIG. 3. Microscopic model and skin effect. (a) An undeformed kagome lattice with generalized Hookean springs. Inset: the force law displayed in Eq. (4). (b) The spectrum of a lattice with periodic boundaries. (c) The spectrum of a circle with pinned boundaries. (d) A topological edge mode with displacements visualized by size and color. (e) A deformed kagome lattice. Due to the chirality in the spring, deformation breaks reflection over horizontal axis. (f) The periodic spectrum. (g) The spectrum with open boundaries. (h) Visualization of the non-Hermitian skin mode. In the examples shown, $\theta = 70^\circ$.

depicted in orange in Fig. 2(c). In the Supplemental Material [41], we show that the inversion symmetry of the honeycomb lattice gives rise to a second manifestation of PT symmetry, given by $PT = KU$, where $U = 1 \otimes \sigma_x$ acts on $D_\theta(\mathbf{q})$. This PT symmetry applies locally at each point in the Brillouin zone. Hence contiguous regions of the Brillouin zone form PT -broken and PT -unbroken phases bounded by rings of exceptional points [3,31,93,94].

Non-Hermitian edge modes.—Finally, we note that the skin effect, previously discussed in the long-wavelength limit within continuum theory, has a counterpart in the optical bands of the lattice models. In Fig. 3, we place the active bonds on an undeformed kagome lattice illustrated in Fig. 3(a) and compute the spectrum with periodic [Fig. 3(b)] and open [Fig. 3(c)] boundaries. We color the modes by their participation ratio $\sum_{\mathbf{x}} |\mathbf{u}(\mathbf{x})|^4$, which serves as a proxy for localization [95]. For the undeformed kagome lattice, the sole difference between the periodic and open boundary spectra is the presence of a subextensive number of localized topological modes that span the band gaps. However, when we introduce a small deformation to the kagome lattice [Figs. 3(e)–3(g)], we observe a dramatic departure [96]. We find that the open boundary system not only contains gap-spanning boundary modes, but the bulk bands become highly localized and their distribution in the complex plane changes dramatically. It is instructive to note the qualitative difference between the topological and skin modes [Figs. 3(d) and 3(h)]. The localized bulk modes are confined to a single direction,

whereas the topological modes are confined to all boundaries and decay into the bulk.

Conclusions.—Our work brings to light the non-Hermitian phenomena that arise at the boundary of elastic media for which energetic sources (powered by internal activity or external fields) modify the relationship between static deformation and stress.

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