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Linking Scalar Elastodynamics and Non-Hermitian Quantum Mechanics

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(Received 8 September 2019; revised manuscript received 18 December 2019; accepted 31 January 2020; published 27 February 2020)

Recent years have seen a fascinating pollination of ideas from quantum theories to elastodynamics—a theory that phenomenologically describes the time-dependent macroscopic response of materials. Here, we open a route to transfer additional tools from non-Hermitian quantum mechanics. We begin by identifying the differences and similarities between the one-dimensional elastodynamics equation without body forces and the time-independent Schrödinger equation and finding the condition under which the two are equivalent. Subsequently, we demonstrate the application of the non-Hermitian perturbation theory to determine the response of elastic systems; the calculation of leaky modes and the energy decay rate in heterogenous solids with open boundaries using a quantum mechanics approach; and construction of degeneracies in the spectrum of these assemblies. The latter result may be of technological significance, as it introduces an approach to harness extraordinary wave phenomena associated with non-Hermitian degeneracies for practical devices, by designing them in simple elastic systems. As an example of such an application, we demonstrate how an assembly of elastic slabs that is designed with two degenerate shear states according to our scheme can be used for mass sensing with enhanced sensitivity by exploiting the unique topology near the exceptional point of degeneracy.

DOI: 10.1103/PhysRevApplied.13.024074

I. INTRODUCTION

The physics of matter at the subatomic level is described by quantum mechanics. The computational complexity associated with the theory at the macroscopic scale renders it infeasible to describe the observable mechanics of materials and hence continuum mechanics is used [1]. Despite the huge difference in the length scales for which the two theories were developed, fascinating realizations of quantum phenomena have been demonstrated using macroscopic systems in recent years [2]. Examples include the Hall effect [3,4], geometric phase [5], and negative refraction [6-10]. Special attention is given to extraordinary transport properties based on \mathcal{PT} symmetry [11–18], which corresponds to the commutativity of an operator with combined parity-time reversal operators. This concept originated from the discovery in quantum mechanics that Hamiltonians exhibiting this symmetry can have real eigenvalues, even if they are not Hermitian [19]. One of the advantages of non-Hermitian quantum mechanics (NHQM) is its quantification of the conditions for the existence of \mathcal{PT} -symmetric Hamiltonians with a real spectrum [20].

The source of these analogies originates from the connection between the governing equations in the different branches of physics. The analogy between the time-independent Schrödinger equation and the scalar elastodynamic equation without body forces that appears in part of the literature identifies the transformation

$$\hat{V}(x) - E \to \frac{\omega^2}{c^2(x)},$$

where $\hat{V}(x)$ is the potential in the quantum Hamiltonian and E is the energy, and in the elastic counterpart ω and c(x) are the wave frequency and velocity, respectively. However, this analogy is flawed, since it mixes the operator and its eigenvalues. Using a simple transformation, we here first identify the term that appears in the one-dimensional (1D) elastodynamics equation and absent from the timeindependent Schrödinger equation. In turn, this derivation allows us to determine the condition under which the two equations are equivalent. Following this analysis, we apply tools from the non-Hermitian formalism of quantum mechanics to elastodynamics, in addition to those transferred recently [21,22], as described next.

First, we show the application of the time-independent Rayleigh-Schrödinger perturbation theory of quantum mechanics to elastodynamics [23,24]. This theory provides

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the solutions of a perturbed Hamiltonian in terms of a series expansion about a Hermitian Hamiltonian, where its non-Hermitian formalism determines the radius of convergence by extending the perturbation to the complex plane. By way of example, we consider an elastic assembly composed of a PMMA slab that is perfectly bonded between two steel slabs that are fixed at the ends. We apply the aforementioned theory to calculate the shear response of an elastic assembly the properties of which are complex perturbations of the original assembly. This response is given in terms of a perturbation expansion, for which we calculate its radius of convergence [25]. Importantly, in this process we also calculate the exceptional point (EP)—the point at which the spectrum of the perturbed assembly has a non-Hermitian degeneracy, where two of its eigenmodes coalesce, together with their corresponding complex frequencies [26–28]. This occurs in our example for an assembly comprising a lossy slab with a specific viscoelastic shear modulus [29] (cf. Ref. [30], which considers a differential loss between two coupled mechanical resonators to tailor EPs).

Subsequently, we present a proof of concept of how this assembly, i.e., an elastic assembly with a non-Hermitian degeneracy in its spectrum, can be utilized for mass sensing with enhanced sensitivity. Specifically, by a combination of algebraic arguments and numerical calculations, we show that when a mass is deposited, the degenerate frequency of the elastic assembly splits into two frequencies, such that the splitting is proportional to the square root of the mass. This phenomenon is the physical manifestation of the topology near an EP in the spectrum of our elastodynamic problem. Accordingly, measurement of the frequency splitting quantifies the weight of the deposited mass, with higher mass responsivity at small masses. By contrast, standard mechanical sensors measure the shift in the mechanical resonant frequency, which scales linearly with the deposited mass and hence have inferior sensitivity at small masses [31,32]. Indeed, the square-root topology near an EP has been used in other systems for sensing [33,34], most recently by Djorwe *et al.* [35] using optomechanical cavities coupled by mechanical resonators.

The second analogy we draw is between the NHQM formalism of the particle in a box model and the previous elastic lossless assembly, when the steel slabs are semi-infinite, resulting in a non-Hermitian system. We emphasize that here the source of non-Hermiticity is the outgoing boundary conditions, not viscosity or gain. It is only in the non-Hermitian formalism of quantum mechanics that the poles of the scattering matrix are associated with metastable states, where the imaginary part of the poles provides the resonance width or the rate of decay of the resonance state [36]. Here, we obtain the physical counterparts of these quantities in the elastodynamic settings. Specifically, we show that the imaginary part of

the poles of the elastic scattering matrix equals half the decay rate of the mechanical energy in the PMMA slab, associated with leaky modes in the elastic assembly.

Finally, we develop a real perturbation theory for the non-Hermitian system using the NHQM complex scaling method [37,38]. With the framework developed in this section, we are able to derive the eigenstates of the perturbed Hamiltonian as an expansion about a non-Hermitian Hamiltonian with real parameters, such as the stiffness and length of a fourth slab in our example. Thereby, we constitute a framework to analyze and construct degeneracies by real perturbations, although its numerical study is beyond our scope here [39]. It has already been established that systems exhibit extraordinary behavior in the vicinity of non-Hermitian degeneracies, such as ultrasensitivity [40] (as we also demonstrate in the sequel), acquisition of a Berry phase [41], and asymmetric scattering properties [42,43]. Accordingly, our framework offers an approach to achieve such extraordinary wave phenomena by designing non-Hermitian degeneracies in simple systems, i.e., purely elastic, without the need for loss through viscoelastic phases or external gain as in the works mentioned earlier.

The results described above are presented in the following order. Section II provides a short summary of the elastodynamics equations and specifically their reduced form in the scalar (1D) setting. Section III identifies the similarities and differences between the scalar elastodynamic equation without body forces and the time-independent Schrödinger equation, based on a transformation that we develop. Section IV formulates the elastic counterpart of the NHQM time-independent Rayleigh-Schrödinger theory, demonstrates the calculation of the radius of convergence for our model-slabs problem, and, importantly, determines the EP in the spectrum of the slabs. Section V demonstrates how the elastic assembly that exhibits a non-Hermitian degeneracy can be utilized for mass sensing with enhanced sensitivity. Section VI details the calculation of metastable states and energy decay when the steel slabs are semi-infinite, using the analogy with the NHQM particle in a box model. The development of a real perturbation theory for non-Hermitian elastic systems is carried out in Sec. VII. A summary of our results and future outlook conclude the paper in Sec. VIII.

II. THE ELASTODYNAMICS EQUATIONS IN CONTINUUM MECHANICS

The continuum governing equations are based on the hypothesis that the interparticle forces can be replaced by the *stress*-tensor field σ ; in terms of σ , the balance of linear momentum in absence of body forces yields [44]

$$\nabla \cdot \boldsymbol{\sigma}(\mathbf{x}, t) = \rho(\mathbf{x}, t) \ddot{\mathbf{u}}(\mathbf{x}, t), \tag{1}$$

where ρ is the mass density and **u** is the displacement vector field of material points. The stress is related to the displacement field via the *constitutive* equation

$$\boldsymbol{\sigma} = \mathbf{C} \left(\mathbf{x} \right) \nabla \mathbf{u}, \tag{2}$$

where **C** is the fourth-order elasticity tensor. If the material is locally isotropic, the tensor **C** is a function of the Lamé parameters μ (**x**) and λ (**x**), and the combination of Eqs. (1) and (2) can be put in the form

$$\{\nabla [\lambda (\mathbf{x}) \nabla \cdot +2\mu (\mathbf{x}) \nabla \cdot] - \nabla \times [\mu (\mathbf{x}) \nabla \times]\} \mathbf{u} = \rho (\mathbf{x}) \ddot{\mathbf{u}}.$$
(3)

Equation (3) exposes the unique coupling in elastodynamics between the volumetric part of the vector field, proportional to $\nabla \cdot \mathbf{u}$, and its transverse or shear part, proportional to $\nabla \times \mathbf{u}$. This coupling has a significant effect on the Hermiticity of the system, discussed elsewhere [10]. When considering 1D motions, the coupling is eliminated and the problem reduces to a scalar one. Using the ansatz $u(x, t) = U(x)e^{-i\omega t}$, Eq. (3) then reduces to

$$-\frac{d}{dx}\tilde{\mu}\frac{d}{dx}U(x) = \rho(x)\omega^2 U(x), \qquad (4)$$

where $\tilde{\mu} = \mu$ when the displacements are normal to the *x* direction (termed transverse or shear waves) and $\tilde{\mu} = \lambda + 2\mu$ when the displacements are along the *x* direction (termed pressure or volumetric waves). In what follows, we focus on the former, bearing in mind that the same analysis holds for the latter, by carrying out a change of modulus.

III. SIMILARITIES AND DIFFERENCES BETWEEN THE 1D ELASTODYNAMIC EQUATION AND THE TIME-INDEPENDENT SCHRÖDINGER EQUATION

The objective of this section is to transform the equation of elastodynamics in the 1D case to a Schrödinger-type equation, in order to highlight the similarities and differences between them. To this end, we first multiply Eq. (4) by $\mu^{-1}(x)$ and define $c^2(x) = \mu(x)/\rho(x)$, to obtain

$$-\frac{1}{\mu}\frac{d}{dx}\mu\frac{d}{dx}U = \frac{\omega^2}{c^2}U.$$
 (5)

Observe that in terms of the variable $\eta(x) = \mu^{-1}(x)$, the left-hand side equals

$$-\eta \frac{d}{dx} \frac{1}{\eta} \frac{d}{dx} U = \left[\frac{1}{\eta} \frac{d\eta}{dx} \frac{d}{dx} - \frac{d^2}{dx^2} \right] U.$$
(6)

By further defining $n^2(x) = \eta(x)$, we rewrite Eq. (5) as

$$\left[2\frac{1}{n}\frac{dn}{dx}\frac{d}{dx} - \frac{d^2}{dx^2}\right]U = \frac{\omega^2}{c^2}U.$$
 (7)

We employ the transformation $U(x) = n(x)\Psi(x)$ and multiply Eq. (7) by c^2/n , to obtain

$$\left[2\frac{c^2}{n^2}\frac{dn}{dx}\frac{d}{dx} - \frac{c^2}{n}\frac{d^2}{dx^2}\right](n(x)\Psi(x)) = \omega^2\Psi(x).$$
 (8)

We rewrite the left-hand side as an operator on $\Psi(x)$, namely,

$$\left[2\frac{c^2}{n^2}\frac{dn}{dx}\left(\frac{dn}{dx}+n\frac{d}{dx}\right)-\frac{c^2}{n}\left(\frac{d^2n}{dx^2}+2\frac{dn}{dx}\frac{d}{dx}+n\frac{d^2}{dx^2}\right)\right]\Psi.$$
(9)

We denote the operator in the square brackets by \hat{H} , to rewrite Eq. (8) in the form

$$\hat{H}\Psi(x) = \omega^2 \Psi(x). \tag{10}$$

After the cancelation of the terms that depend on dn/dx, the operator \hat{H} in this eigenvalue problem for ω^2 can be separated according to $\hat{H} = \hat{T} + \hat{V}_{CM}$, where

$$\hat{T} = -c^2(x)\frac{d^2}{dx^2},\tag{11}$$

and

$$\hat{V}_{\rm CM} = 2 \left(\frac{c(x)}{n(x)} \frac{dn}{dx} \right)^2 - \frac{c^2(x)}{n(x)} \frac{d^2n}{dx^2}$$
(12)

is a local function of x and does not involve spatial derivatives, hence can be interpreted as the potential of a conservative force (CM stands for continuum mechanics). By further separating \hat{T} using the product rule according to

$$\hat{T} = \hat{T}_{\rm QM} + \hat{T}_{\rm CM}, \quad \hat{T}_{\rm QM} = -\frac{d}{dx}c^2 \left(x\right)\frac{d}{dx},$$
$$\hat{T}_{\rm CM} = \frac{d}{dx}c^2\frac{d}{dx}, \tag{13}$$

we can identify \hat{T}_{QM} with the kinetic energy operator in the Schrödinger equation of an electron with an effective mass

$$m_{\rm eff}(x) = \frac{1}{2c^2(x)},$$
 (14)

which varies when the electron traverses different semiconductors. The difference between the equations thus amounts to $\hat{T}_{\rm CM}$ —this term does not have the form of a



FIG. 1. An assembly of three elastic slabs. The Hermitian case corresponds to fixed boundaries at $x = \pm L$ and the non-Hermitian case corresponds to an infinite assembly with outgoing boundary conditions.

kinetic energy operator or a potential, as it involves one spatial derivative.

To draw the analogy with the NHQM model problem of a particle in a box with outgoing boundary conditions, we consider the prevalent case of a solid that is composed of different homogeneous slabs. The medium properties are therefore piecewise constant. For simplicity, we consider two constituents—say, material *a* with ρ_a and μ_a , which is perfectly bonded at $x = \pm l$ to two semi-infinite slabs made of a stiffer material *b* with ρ_b and $\mu_b > \mu_a$ (Fig. 1).

If we further assume that

$$\frac{\mu_a}{\rho_a} = \frac{\mu_b}{\rho_b},\tag{15}$$

then \hat{T}_{CM} vanishes; in this case—and this case only—there is an exact analogy between the 1D equation of elastodynamics and the time-independent Schrödinger equation. The corresponding potential \hat{V}_{CM} exhibits a potential well between two infinite barriers (spikes), owing to the jump discontinuities of $\mu(x)$, and hence of n(x). In the equivalent quantum-particle-in-a-box model, there are resonance phenomena and metastable states associated with complex eigenvalues that are elegantly analyzed and explained using the NHQM formalism [36]. In the sequel, we will show that this formalism establishes a powerful mechanism to study corresponding elastodynamic phenomena, even when restrictions (15) are removed and the exact analogy is broken. Before we proceed, it is useful to note that for two-dimensional (2D) elastodynamics, by contrast, an exact analogy with the time-independent Schrödinger equation always exists. To show this, it is sufficient to consider anti-plane shear waves of the form

$$u(x, y, t) = Y(y)e^{i(k_x x - \omega t)},$$
(16)

propagating in a medium that is laminated in the y direction. In each lamina, Eq. (3) reduces to the following equation for Y(y):

$$\left[\frac{d^2}{dy^2} + \frac{\omega^2}{c^2(y)}\right]Y(y) = k_x^2 Y(y).$$
 (17)

In this case, it is possible to identify k_x^2 (not ω^2) as the eigenvalue to be determined, d^2/dy^2 with the kinetic energy operator, and $\omega^2/c^2(y)$ with the potential. Physically, Eq. (17) represents the following question (cf. Ref. [45] on a similar analogy between the time-independent Schrödinger equation and Maxwell equations): given excitation frequency and mechanical properties, what would the propagation constant be in the direction perpendicular to the material modulation?

Returning to the 1D problem, we consider next the transformation $U(x) = \rho^{-1/2}(x)\Psi(x)$, which upon substitution into Eq. (4) and its multiplication by $\rho^{-1/2}(x)$, provides a different representation of the Hamiltonian, namely,

$$\hat{H} = -\frac{1}{\sqrt{\rho(x)}} \frac{d}{dx} \mu(x) \frac{d}{dx} \frac{1}{\sqrt{\rho(x)}}.$$
(18)

For real moduli, this Hamiltonian is Hermitian if it operates on functions that vanish at the boundary of the problem domain and therefore the eigenfunctions $\{\Psi(x)\}$ are orthogonal one to each other; the application of NHQM perturbation theory to such Hermitian systems in 1D elastodynamics is demonstrated first.

IV. NHQM PERTURBATION THEORY FOR ELASTODYNAMICS: THE MODEL PROBLEM OF A FINITE SLAB

In quantum mechanics, the standard time-independent Rayleigh-Schrödinger theory provides the solutions of a perturbed Hamiltonian in terms of a series expansion about a Hermitian Hamiltonian. The NHQM formalism determines the radius of convergence by extending the perturbation to the complex plane and calculating the EP—the point at which the perturbed Hamiltonian has a non-Hermitian degeneracy [27,28,36]. The process is exemplified in this section, by calculating first the eigenfrequencies and eigenmodes of a Hermitian Hamiltonian that models an elastodynamic system made of purely elastic and finite slabs; subsequently, we determine the convergence radius of the elastodynamic Rayleigh-Schrödinger expansion by calculating the EP in the perturbed non-Hermitian Hamiltonian spectrum.

Thus, we truncate the assembly at $x = \pm L$ and fix the boundaries such that the displacement field vanishes at the edges. The standard procedure to calculate the real frequencies starts with the ansatz

$$U^{(0)}(x) = \begin{cases} A\cos k_a x + B\sin k_a x, & x < |l|, \\ C_+ \sin k_b (x - L), & l < x < L, \\ C_- \sin k_b (x + L), & -L < x < -l, \end{cases}$$
(19)

where, owing to Eq. (3) and the continuity of u(x, t),

$$\omega^2 = c_i^2 k_i^2, \quad i = a, b, \tag{20}$$

(28)

and hence k_a and k_b are related via

$$\frac{k_a}{k_b} = \frac{c_b}{c_a}.$$
(21)

The continuity of the spatial parts of the displacement and stress at $x = \pm l$ takes the form

$$A\cos k_a l = C\sin k_b (l-L), \qquad (22)$$

$$-A\mu_a k_a \sin k_a l = \mu_b k_b C \cos k_b (l-L), \qquad (23)$$

from which the relation between the amplitudes A and C is determined. The resultant transcendental equation for the eigenfrequencies is

$$\tan \frac{\omega l}{c_a} = \gamma \, \tan \frac{\omega (l-L)}{c_b}, \quad (\text{odd modes}) \tag{24}$$

$$\cot \frac{\omega l}{c_a} = -\gamma \, \tan \frac{\omega (l-L)}{c_b}, \quad (\text{even modes}). \tag{25}$$

We denote the eigenfrequencies by $\{\omega_m^{(0)}\}_{m\in\mathbb{N}}$ and the corresponding transformed eigenfunctions of Eq. (18) by $\Psi_m^{(0)}(x)$. It is clear that $\{\omega_m^{(0)}\}_{m\in\mathbb{N}}$ are real and the Hamiltonian is indeed Hermitian.

Consider next another assembly, obtained by replacing the right half of the central slab by a slab whose shear stiffness differs from the shear stiffness of the original slab by α . The resultant Hamiltonian can be written as a sum of the Hamiltonian of the original medium, denoted $\hat{H}^{(0)}$, and a perturbation $\alpha \hat{H}^{(1)}$, where

$$\hat{H}^{(1)} = -\frac{1}{\sqrt{\rho_a}} \frac{d}{dx} \frac{d}{dx} \frac{1}{\sqrt{\rho_a}}$$
(26)

operates on functions over 0 < x < l. Up to a critical value of α , NHQM perturbation theory can deliver the response of the perturbed assembly, in terms of $\Psi_m^{(0)}(x)$ as the zero-order solutions [46]. Using the standard time-independent Rayleigh-Schrödinger theory, we obtain the *n*th-order correction terms $\omega^{(j)}$ and $\Psi_m^{(j)}(x)$, namely,

$$\omega_m^2(x;\alpha) = \sum_{j=0}^{\infty} \alpha^j \, \omega_m^{2(j)}, \quad \Psi_m(x;\alpha) = \sum_{j=0}^{\infty} \alpha^j \, \Psi_m^{(j)}; \quad (27)$$

the convergence of these sums is limited to values of α —including complex values—inside a circle in the complex plane; the origin of the circle is 0, and its radius is denoted by $|\alpha^{EP}|$. This radius equals the radius of the complex branch point at which two adjacent modes coalesce. Thus, a non-Hermitian degeneracy is obtained when the

conditions

and

$$\Psi_m(x;\alpha^{\rm EP}) = \Psi_{m\pm 1}(x;\alpha^{\rm EP}) \equiv \Psi_m^{\rm EP}(x)$$
(29)

are satisfied. Since, for any value of $\alpha \neq \alpha^{EP}$, the two modes $\Psi_m(x; \alpha^{EP})$ and $\Psi_{m\pm 1}(x; \alpha^{EP})$ are orthogonal to each other, at the EP $\Psi_m^{EP}(x)$ is self-orthogonal, as the two solutions coalesce [36]. To determine α^{EP} , we first represent $\hat{H}^{(0)}$ and $\hat{H}^{(1)}$ using the matrices $\mathsf{H}^{(0)}$ and $\mathsf{H}^{(1)}$, defined by

 $U_m(\alpha^{\rm EP}) = U_{m\pm 1}(\alpha^{\rm EP}) \equiv U_m^{\rm EP}$

$$\mathsf{H}^{(\alpha)}{}_{mn} = \int_{\mathscr{I}} \hat{\Psi}^{(0)}_{m} \hat{H}^{(i)} \hat{\Psi}^{(0)}_{n} \, dx, \quad i = 0, 1, \qquad (30)$$

where $\mathscr{I} = [-L, L]$ and [0, l] when i = 0 and 1, respectively, and

$$\hat{\Psi}_m^{(0)} = \Psi_m^{(0)} \left/ \int_{-L}^{L} \Psi_m^{2(0)} \, dx.$$
(31)

Note that the standard procedure to derive orthogonality relations for real functions provides

$$\left\langle \hat{\Psi}_{m}^{(0)}, \hat{\Psi}_{n}^{(0)} \right\rangle := \int_{-L}^{L} \hat{\Psi}_{m}^{(0)} \hat{\Psi}_{n}^{(0)} dx = \delta_{mn}.$$
 (32)

We are now in a position to seek the smallest α for which the matrix

$$H(\alpha) = H^{(0)} + \alpha H^{(1)}$$
(33)

has an eigenvalue multiplicity, using a modified Newton's method [47]. To proceed with numerical computations, we consider by way of example a middle slab made of PMMA, which is bonded between two steel slabs, the properties of which are

$$\rho_a = 1200 \text{ kg m}^{-3}, \quad \mu_a = 1.21 \text{ GPa}, \quad l = 1 \text{ cm},$$

 $\rho_b = 7800 \text{ kg m}^{-3}, \quad \mu_b = 78.85 \text{ GPa}, \quad L = 3 \text{ cm}.$
(34)

For simplicity, we truncate the size of $H^{(\alpha)}$ to 2 × 2 using the first odd and even modes.

The results are shown in Fig. 2 in a dimensionless form (lengths are divided by L; mass densities and shear moduli are divided by the mean value of the quantity when averaged between the two phases).

Specifically, Fig. 2(a) shows the inverse of the absolute value of the difference between the two eigenvalues of H versus Re α and Im α ; the peak at $\alpha \simeq 0.38 +$



FIG. 2. (a) The inverse of the absolute value of the difference between the eigenvalues as a function of α . Observe that $\lim_{\alpha \to \alpha^{\rm EP}} |\omega_1^2 - \omega_2^2|^{-1} = \infty.$ (b) The inverse of the inner prodbetween the first right uct left eigenvectors as a and function of α . Observe that $\lim_{\alpha\to\alpha^{\mathrm{EP}}} |\langle \hat{\Psi}_{R1}, \hat{\Psi}_{L1} \rangle$ $=\infty$. (c) The real part of the eigenvalues as a function of α . The red and blue surfaces correspond to ω_1^2 and ω_2^2 , respectively. (d) The imaginary part of the eigenvalues as a function of α (same color legend as in the previous panel). At $\alpha = \alpha^{EP}$, the two eigenvalues coalesce to a single complex value associated with ω^{EP} .

0.074*i* identifies α^{EP} and hence the radius of convergence. We note that such a value of α , i.e., with a positive imaginary part, physically corresponds to a viscoelastic slab and hence the realization of this EP does not require any gain. Figure 2(b) shows the inverse of the inner product between the first right $(\hat{\Psi}_{R1})$ and left $(\hat{\Psi}_{L1})$ eigenvectors as a function of α . Indeed, we observe that $\lim_{\alpha \to \alpha^{\text{EP}}} \left| \left\langle \hat{\Psi}_{R1}, \hat{\Psi}_{L1} \right\rangle \right|^{-1} = \infty$, confirming that this is a non-Hermitian degeneracy, since at this type of degeneracy the relevant modes are self-orthogonal [48]. Finally, we show that the spectrum in the vicinity of the EP exhibits a Riemann surface structure-the signature of non-Hermitian degeneracy—by plotting the real [Fig. 2(c)] and imaginary [Fig. 2(d)] parts of ω_1^2 (red surface) and ω_2^2 (blue surface). We denote for later use the frequency at the EP by ω^{EP} , such that

$$\omega^{\text{EP}} := \omega_1 \left(\alpha = \alpha^{\text{EP}} \right) = \omega_2 \left(\alpha = \alpha^{\text{EP}} \right).$$
(35)

V. APPLICATION TO MASS SENSING

We demonstrate next how the unique topology near the EP can be harnessed to design a mass sensor with enhanced sensitivity, based on the aforementioned assembly when tuned to operate at the EP. First, we recall that standard mechanical mass sensors are based on the shift in the mechanical resonant frequency, owing to any deposited mass [31]. For small masses, the shift is linear in the perturbation [32,49]. This linear relation can be interpreted as the first term in the Taylor series of the frequency as a function of the mass:

$$\operatorname{Re}\omega\left(m\right) - \operatorname{Re}\omega\left(0\right) = S_{T}m + \mathcal{O}\left(m^{2}\right), \qquad (36)$$

where *m* is the mass of the deposited element, Re ω (0) is the (real part of the) resonant frequency of the unperturbed system, and $S_T = \partial \text{Re} \omega (m) / \partial m$ at m = 0. By contrast, there is not a Taylor expansion of the frequency shift from an EP; however, it does admit a Puiseux series. In the case when the EP is of *two* eigenvalues and eigenmodes, we have that

$$\operatorname{Re}\omega(m) - \operatorname{Re}\omega^{\operatorname{EP}} = S_P\sqrt{m} + \mathcal{O}(m), \qquad (37)$$

with some coefficient S_P . Sensing of a device is thus quantified by the so-called mass responsivity $R = \partial \operatorname{Re} \omega / \partial m$ [32,50]. It is clear that conventional sensors have a finite R as $m \to 0$, whereas for EP-based sensors,

$$\lim_{m \to 0} \frac{\partial \operatorname{Re} \omega(m)}{\partial m} \propto \lim_{m \to 0} \frac{1}{\sqrt{m}} = \infty, \quad (38)$$

i.e., theoretically an infinite sensitivity, which in practice is limited by the resolution of the frequency measurement. This feature has been employed for sensors in different physical systems [33,34] and specifically in systems comprising optomechanical cavities coupled by mechanical resonators [35]. Here, we apply this approach to the elastodynamic system described in the previous section, emphasizing that its EP does not require any realization of gain, as that assembly comprises only elastic and viscoelastic slabs. This is carried out by calculating the eigenfrequencies when a discrete element with mass m is deposited at the center of the assembly, the Hamiltonian of which is given by Eqs. (33) and (34) with $\alpha = \alpha^{EP}$. The mass is modeled by replacing $\rho(x)$ with $\rho(x) + \delta \rho(x)$ in the Hamiltonian (18), where $\delta \rho(x) = \delta \rho_0$ over -L/50 <x < L/50 such that $m = \delta \rho_0 L/100$ [51], and calculating



FIG. 3. (a) The real part of the first (blue circles) and second (red circles) eigenfrequencies of our elastodynamic system, when operating at an EP and augmented by a deposited mass *m* (in grams). The solid lines are the functions (39), which are proportional to \sqrt{m} , and their agreement with the calculated frequencies confirms the square-root nature of the frequency splitting. By contrast, conventional mechanical sensors are linear in *m*. (b) The mass responsivity $R = \partial \text{Re } \omega / \partial m$ of the device as a function of *m*. Note the theoretical infinite responsivity in the limit of an infinitesimal mass.

the resultant eigenvalues of Eq. (33). Figure 3(a) depicts (the real part of) the first (blue circles) and second (red circles) eigenfrequencies for representative values of the deposited mass (in grams). It is shown how frequency splitting occurs owing to the added mass, in a manner that is nonlinear in the perturbation—the smaller the mass, the greater the relative change. The solid lines are the functions

$$\operatorname{Re}\omega^{\mathrm{EP}} + S_P \sqrt{m},\tag{39}$$

where $S_P = -2.02$ and 1.845 for the lower and upper curves, respectively; the matching between Eq. (39) and the calculated frequencies thereby confirms the conjectured square-root nature of the frequency dependency in the deposited mass. Again, we emphasize that the scaling is linear in conventional mechanical sensors based on the shift of the resonant frequency and therefore inferior for extremely small masses.

The enhanced sensitivity near the EP is further highlighted in Fig. 3(b), by plotting the mass responsivity as a function of *m*, using the derivative of the fitted function (39) for the higher frequency. Thereby, we show the theoretical infinite responsivity in the limit of an infinitesimal mass.

VI. A NON-HERMITIAN MODEL PROBLEM WITH OUTGOING BOUNDARY CONDITIONS

We pursue next the analogy drawn in Sec. III between the infinite purely elastic assembly and the model problem in NHQM of a particle in a box with outgoing boundary conditions. We emphasize that the non-Hermiticity of this problem stems from the outgoing boundary conditions, not viscosity, as the slabs here are purely elastic. In what follows, we present the physical interpretation of the NHQM tools for elastodynamics. Specifically, we demonstrate that the NHQM approach provides the so-called leaky eigenmodes of the system, the imaginary part of the eigenvalues of which delivers the decay rate of the elastic energy in the middle slab.

Accordingly, when the steel slabs now extend to $\pm\infty$ and the PMMA slab is released from some arbitrary initial shear deformation, we impose outgoing boundary conditions and seek solutions to U(x) in the form

$$U(x) = \begin{cases} A\cos k_a x + B\sin k_a x, & x < |l|, \\ C_+ e^{ik_b(x-l)}, & x > l, \\ C_- e^{-ik_b(x+l)}, & x < -l. \end{cases}$$
(40)

(The relation between k_a , k_b , and ω remains as in the Hermitian problem.) The continuity of the displacement and stress at $x = \pm l$ now reads

$$A\cos k_a l \pm B\sin k_a l = C_{\pm},\tag{41}$$

$$\mu_a k_a (\mp A \sin k_a l + B \cos k_a l) = \pm C_{\pm} \mu_b i k_b.$$
(42)

Manipulating these equations provides

$$i \tan k_a l = \gamma$$
 (odd modes), (43)

$$i \cot k_a l = -\gamma$$
 (even modes), (44)

where the impedance mismatch γ is

$$\gamma = \frac{\mu_a k_a}{\mu_b k_b} = \frac{\mu_a c_b}{\mu_b c_a} = \sqrt{\frac{\mu_a \rho_a}{\mu_b \rho_b}}.$$
 (45)

Equations (43) and (44) are solved by

$$k_a l = -i \operatorname{arctanh} \gamma + \frac{m\pi}{2}, \quad m \in \mathbb{Z},$$
 (46)

where odd and even m correspond to even and odd modes, respectively. Hence, there are infinitely many discrete complex solutions with a different real part and the same imaginary part; there are no bound states associated with



FIG. 4. The log of the mechanical energy stored in the PMMA slab (normalized by the energy at t = 0) as a function of t (ms), calculated using a finite-volume scheme (blue curve). The slope of its linear interpolation (red curve) matches $-2 \text{Im} \omega$.

real solutions. The obtained roots are the poles of the scattering matrix, which only in the non-Hermitian formalism of quantum mechanics delivers fundamental information on the modes, without the need to carry out wave-packet calculations. For example, in NHQM, the imaginary part of the poles provides the resonance width, or rate of decay of resonance state [36]. Here, by analogy, the imaginary part in Eq. (46) should provide information on the decay rate of the mechanical energy. This relation is demonstrated in Fig. 4, where the log of the mechanical energy stored in the PMMA slab,

$$E_a(t) = \frac{1}{2} \int_{-l}^{l} \left(\mu_a u_{,x}^2 + \rho_a u_{,t}^2 \right) \, dx, \tag{47}$$

(normalized by the energy at t = 0) is evaluated as a function of t, when calculated using the finite-volume method [52] for some (real) arbitrary initial conditions. Indeed, the slope of its linear interpolation (red curve) matches $-2\text{Im}\,\omega$, and is independent of the form of the initial conditions.

VII. THEORY FOR REAL PERTURBATIONS IN NON-HERMITIAN ELASTODYNAMICS: THE 1D MODEL PROBLEM

In this last section, we are interested in developing a perturbation theory to the latter problem, noting that the obstacle lies in the divergence of $\Psi(x)$ at $\pm\infty$. Therefore, the orthogonality relations (30) no longer hold and the components of H⁽⁰⁾ are unbounded when calculated according to Eq. (32). To overcome these obstacles, we first apply the complex scaling transformation $(x - l) \rightarrow (x - l) e^{i\theta}$ for $\Psi(x > l)$, with sufficiently large and real θ

[53]. In these rotated coordinates, the transformed function

$$\Psi_n (x > l; \theta) = e^{ik_{bn}(x-l)e^{i(\theta-\varsigma_n)}}$$

= $e^{i|k_{bn}|(x-l)\cos(\theta-\varsigma_n)-\sin(\theta-\varsigma_n)}$ (48)
 $\times e^{-|k_{bn}|(x-l)\sin(\theta-\varsigma_n)},$

with $\tan \varsigma_n = -(n\pi)^{-1} \operatorname{arctanh} \gamma$, vanishes at infinity, owing to the second decaying exponent. Similarly, $\Psi(x < -l; \theta)$ vanishes at $-\infty$ by applying the transformation $(x + l) \rightarrow (x + l) e^{i\theta}$.

To establish next an orthonormal basis set, we replace the scalar product of the Hermitian formalism with the NHQM *c-product* [36], namely,

$$(\Psi_{n},\Psi_{n}) := \int_{-\infty}^{-l} \Psi_{n}^{2}(x;\theta) \, dx + \int_{-l}^{l} \Psi_{n}^{2}(x) \, dx + \int_{l}^{\infty} \Psi_{n}^{2}(x;\theta) \, dx.$$
(49)

To show that this product indeed delivers such a set, consider first the third term in Eq. (49). Since the scaled function vanishes at infinity, the integral is zero at its upper limit and we are left with its value at x = l, such that

$$\int_{l}^{\infty} \rho_b C_{n+}^2 e^{2ik_{bn}(x-l)e^{i\theta}} e^{i\theta} dx = \frac{i\rho_b}{4k_{bn}} \left(1 \pm \cos 2k_{an}l\right),$$
(50)

where we use the fact that $C_{n+} = \cos k_{an}l$ for the even modes (with a plus sign inside the brackets) and $C_{n+} = \sin k_{an}l$ for the odd modes (minus sign). Owing to symmetry, this is also the value of the first integral in Eq. (49) and the remaining integral amounts to

$$\int_{-l}^{l} \Psi_n^2(x) \, dx = \rho_a \left(l \pm \frac{\sin 2k_{an}l}{2k_{an}} \right), \tag{51}$$

where the plus and minus signs correspond to even and odd modes, respectively. We can now redefine the basis (31) to

$$\hat{\Psi}_{n}^{(0)} = \Psi_{n}^{(0)} / (\Psi_{n}, \Psi_{n})$$
(52)

and observe that

$$(\Psi_n, \Psi_m) = \delta_{nm} \tag{53}$$

when invoking Eq. (46).

As in Sec. IV, we again replace a part of the middle slab by a third constituent, the shear stiffness of which is parametrized by real α ; now, however, we leave the location, say d, and length, say D - d, of the replacement as parameters. Using the framework developed in

this section, we can derive the eigenstates of the perturbed Hamiltonian as an expansion about a non-Hermitian Hamiltonian with real parameters. Specifically, we have that

$$H(\alpha, d, D) = H^{(0)} + \alpha H^{(1)}(d, D), \qquad (54)$$

where $H^{(0)}$ is diagonal with complex eigenvalues associated with Eq. (46) and

$$H_{nm}^{(1)} = \int_{D}^{d} \Psi_{n}(x) \frac{d^{2}}{dx^{2}} \Psi_{m}(x) dx$$
$$= k_{am}^{2} \rho_{a} \int_{d}^{D} \Psi_{n}(x) \Psi_{m}(x) dx$$
(55)

is a complex asymmetric matrix $H^{(1)}$ that depends nonlinearly on *d* and *D* (the resultant closed-form expressions are omitted here, for brevity). Notably, our developments further establish a platform for constructing degeneracies by real perturbations.

VIII. SUMMARY AND OUTLOOK

Motivated by the development of non-Hermitian quantum mechanics and the transfer of concepts from quantum theories to the macroscopic scale, we here revisit the connection between the time-independent Schrödinger equation and the 1D elastodynamics equation without body forces. Using a simple transformation, we first identify the term that appears in the elastodynamics equation and is absent from the quantum mechanics equation. This derivation allows us to determine the condition under which the two equations are equivalent.

Subsequently, we show the physical interpretation and application of different tools from non-Hermitian quantum mechanics in elastodynamics, including the timeindependent Rayleigh-Schrödinger perturbation theory to calculate the dynamic response of a finite elastic assembly; the non-Hermitian formalism of this theory to determine the perturbation-series radius of convergence and the exceptional point in the spectrum of the assembly; calculation of leaky modes and energy decay in an open elastic assembly using the poles of the scattering matrix; and the complex scaling transformation for establishing a basis from the corresponding divergent eigenfunctions.

Note that we introduce a framework to analyze and design non-Hermitian degeneracies by real perturbations. These degeneracies have great potential in applications such as ultrasensitive sensors and unidirectional energy scatterers, for which our approach offers a means of access without the need for gain or \mathcal{PT} symmetry. As a concrete application, we demonstrate how an elastic slab assembly can function as an enhanced mass sensor, when designed according to our analysis to exhibit two degenerate shear

states. Using algebraic arguments and numerical calculations, we show in Sec. V how the mass responsivity of this sensor surpasses the responsivity of conventional mechanical sensors, owing to the square-root topology near the exceptional point in its spectrum.

We expect that our application of tools from non-Hermitian quantum mechanics in elastodynamics, thereby showing their usefulness, will pave the way for further developments in more complex practical elastic systems. Examples include periodic composites and homogenization [54–59], anisotropic media [60,61], and elastodynamics in higher dimensions [62].

ACKNOWLEDGMENTS

This research was supported in parts by the I-Core: the Israeli Excellence Center "Circle of Light," the Israel Science Foundation (Grants No. 1530/15 and No. 1912/15), the United States–Israel Binational Science Foundation (Grant No. 2014358), and the Ministry of Science and Technology. We thank Ron Ziv for sharing his MATLAB code and anonymous reviewers for constructive comments that helped us improve this paper.

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