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2	High-order exceptional points in diffusive systems: robust APT symmetry
3	against perturbation and phase oscillation at APT symmetry breaking
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14	Abstract: In non-Hermitian systems with the Hamiltonians obeying parity-time (PT)
15	symmetry, exploring the counterintuitive physics induced by degeneracies known as
16	exceptional points (EPs) provides unprecedented ways to control energy flow. Recently,
17	there are growing interests in bridging wave systems and diffusive systems, where anti-
18	parity-time (APT) symmetry is demonstrated in diffusive systems. In this work, we start
19	from the thermal energy transfer in a four-channel coupling model with the background

20	flow velocities in adjacent channels opposite. A third order EP exists in this system,
21	where temperature profiles in the moving channels are static in the APT symmetric
22	phase (flow velocities below a threshold $v_{\rm EP}$ at the EP), and the profiles begin to
23	dynamically evolve in the APT broken phase (> $v_{EP}$ ). By introducing a velocity
24	perturbation into the background flow at the third order EP ( $v_{EP} \pm \Delta v$ ), we find APT
25	symmetry keeps robust with the phases of temperature profiles in adjacent channels
26	relatively static or locked. When $\Delta v$ is increased above a threshold (another EP), the
27	APT symmetry is breaking with a transition from phase locking to phase oscillation,
28	regardless of initial conditions. This work unveils rich physics in convectively coupled
29	diffusive systems and offers us new prospects for the control of complex thermal fields.
30	
31	Keywords: Anti-parity-time symmetry; Exceptional point; Phase transition; Heat
32	transfer.
33	
34	1. Introduction
35	Parity-time (PT) symmetry is attractive in quantum mechanics, since it allows for
36	real eigenvalues in the non-Hermitian Hamiltonians that are associated with observable
37	quantities in physical systems. <sup>1</sup> For a physical system, the parity operator $\hat{P}$ and time

reversal operator  $\hat{T}$  acts based on the rules of  $\hat{P}\psi(x) = \psi(-x)$  and  $\hat{T}\psi(x) = \psi^*(x)$ , 38 respectively, where \* denotes the complex conjugate.<sup>2</sup> However, in PT symmetric 39

systems, real eigenvalue is a conditional but not necessary result, since there exist phase 40

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transition points at which the PT symmetry will be spontaneously breaking and the 41 eigenvalues become complex. Such phase transition points are termed as exceptional 42 points (EPs), where eigenmodes are degenerate with both eigenvalues and eigenvectors 43 coalesced.<sup>3,4</sup> Here, EPs are completely different from diabolic points in the parameter 44 spaces of Hermitian systems at which the eigenvectors are orthogonal.<sup>5,6</sup> In the past two 45 decades, the paradigm of PT symmetry in quantum mechanics has been successfully 46 shifted into classical systems.<sup>7-13</sup> For example, in optics and acoustics, the PT symmetry 47 48 is constructed by introducing anti-symmetrically distributed gain and loss materials. New perspectives are envisioned in the PT symmetric platform, where various 49 50 counterintuitive effects are theorectically proposed and experimentally demonstrated, such as unidirectional transparency.<sup>7,8</sup> one-way cloaking.<sup>9,10</sup> mode switching.<sup>11,12</sup> EP 51 sensing,<sup>13,14</sup> coherent lasing and absorption.<sup>15,16</sup> 52

Recently, it was found that EPs also exist in anti-parity-time (APT) symmetric 53 systems, at which phase transition occurs with the eigenvalues changing from pure 54 imaginary (APT symmetric phase) into complex (APT broken phase).<sup>17,18</sup> Unlike the 55 PT symmetric Hamiltonian that satisfies the relation  $\hat{P}\hat{T}H = H\hat{P}\hat{T}$ , the APT symmetric 56 Hamiltonian follows the rule of  $\hat{P}\hat{T}H = -H\hat{P}\hat{T}$ .<sup>19,20</sup> Mathematically, the PT symmetric 57 Hamiltonian can be transformed into the APT symmetric one by simply multiplying the 58 Hamiltonian with an imaginary number *i*. Physically, for a tight-binding model, this 59 60 operation will end up with a pure virtual coupling between two tight-binding sites, which is quite challenging for the practical implementation. In optics, the virtual 61

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coupling between adjacent meta-atoms is realized in an indirect and complicated way 62 by adding a well-designed third meta-atom in-between to equivalently generate a virtual 63 coupling action.<sup>19,21</sup> It needs to be mentioned that the diffusive systems (e.g., thermal 64 systems) are inherently non-Hermitian. The most interesting is the notion that the 65 66 coupling in diffusive systems is originally imaginary. Thus, with the aid of convection and low diffusivity, we can imitate various wave-like dynamics in the framework of 67 diffusive systems.<sup>22</sup> For example, the stable temperature profile under low diffusivity, 68 which mimics a wave packet, can stop or even move in the opposite direction against 69 the background flow via convection couplings, corresponding to zero or negative group 70 71 velocity of a wave packet.

72 In this work, we explore the phase transition in a four-channel coupling thermal system, which, compared with the two-channel toy model,<sup>22</sup> provides a higher degree 73 74 of freedom to control energy flow. When the background flow velocities in neighboring channels are opposite, there exists a third order EP in the eigenspectrum, accompanied 75 with APT symmetry breaking. Adding a perturbation into the background flow velocity 76 at the high order EP ( $v_{\rm EP} \pm \Delta v$ ), the APT symmetry remains robust, where the phases 77 78 of temperature profiles in adjacent moving channels are relatively static, as manifested 79 in a locked mode. When the perturbation  $\Delta v$  surpasses a threshold (another EP), the whole system will transit into the broken APT symmetry. In this case, we observe the 80 81 effect of robust phase oscillation that is irrelevant to the initial conditions, induced by 82 the concealing dimension in the high order EP. Just as the transformation thermotics

that rapidly developed together with the transformation optics and acoustics,<sup>23-26</sup> APT
phase transition at EPs in diffusive systems, inspired from the PT phase transition in
classical wave systems, will promisingly expand the vision of counterintuitive thermal
flow regulation.

87 **2. Model and theory** 

Figures 1(a) and 1(b) show the schematics of 3D model and the corresponding 2D model, respectively. The four identical ring-shaped channels are stacked along *z*-axis, where the thickness, inner radius and outer radius of each channel are *b*,  $R_1$  and  $R_2$ , respectively. Adjacent channels are coupled through an oil layer with the thickness *d*. Rotation velocities of the background flow in the four channels are set to be  $\Omega_{1,3} = v/R_1$  and  $\Omega_{2,4} = -v/R_1$ , respectively, given that  $R_1 \approx R_2$ .



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Fig. 1 (a), (b) The schematic of 3D model and the corresponding 2D model. Here, the channel width is *b*. The thickness of oil layers is *d*. The inner radius of the ring-shaped channel is  $R_1$ . The outer radius is  $R_2$ . The velocities of background flows in adjacent channels are  $\Omega = \pm v/R_1$ . (c) The imaginary part and real part of eigenvalues *vs*. the background flow velocity. Red dots at **A** and **B** mark the EPs.

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Diffusive systems are dissipative with energy exchanges to the environment, which 101 102 can be regarded as inherently non-Hermitian systems. In stark contrast with wave systems that are governed by the real and Hermitian Hamiltonians, diffusive systems 103 104 are featured with pure imaginary Hamiltonians. Recently, great efforts have been made 105 to marry the two different scenarios, where the introduced background flow velocity in the convection process can serve as the effective group velocity for directional energy 106 107 flows, on condition that the thermal diffusivity of materials is trivial. In this case, the 108 Hamiltonians of diffusion systems become complex, with the APT symmetry possibly constructed and EPs generated. For the model displayed in Fig. 1, the convection-109 110 diffusion equations describing the temperature field evolutions take the forms of

111 
$$\frac{\partial T_1}{\partial t} = D \frac{\partial^2 T_1}{\partial x^2} + v \frac{\partial T_1}{\partial x} + \frac{h_{S1}}{\rho_r C_r}; h_{S1} = \frac{\kappa_o}{bd} (T_2 - T_1),$$

112 
$$\frac{\partial T_2}{\partial t} = D \frac{\partial^2 T_2}{\partial x^2} - v \frac{\partial T_2}{\partial x} + \frac{h_{S2}}{\rho_r C_r}; h_{s2} = \frac{\kappa_o}{bd} (T_1 + T_3 - 2T_2),$$

113 
$$\frac{\partial T_{3}}{\partial t} = D \frac{\partial^{2} T_{3}}{\partial x^{2}} + v \frac{\partial T_{3}}{\partial x} + \frac{h_{S3}}{\rho_{r} C_{r}}; h_{S3} = \frac{\kappa_{o}}{bd} (T_{2} + T_{4} - 2T_{3}),$$
  
114 
$$\frac{\partial T_{4}}{\partial t} = D \frac{\partial^{2} T_{4}}{\partial x^{2}} - v \frac{\partial T_{4}}{\partial x} + \frac{h_{S4}}{\rho_{r} C_{r}}; h_{S4} = \frac{\kappa_{o}}{bd} (T_{3} - T_{4}),$$
(1)

115 where  $T_{1,2,3,4}$  are the temperature profiles in channels 1, 2, 3, 4.  $\rho_r$  and  $C_r$  are the

mass density and heat capacity of the ring materials, respectively. In Eq. (1),  $D = \kappa_r / \rho_r C_r$  is the thermal diffusivity of the ring materials, with  $\kappa_r$  denoting the thermal conductivity. For the coupling oil layers,  $h_{s1,s2,s3,s4}$  and  $\kappa_o$  represent the coupling strengths between adjacent channels and the thermal conductivity of oil, respectively.

120 Equation (1) is based on the continuity of temperature fields on the boundaries of oil layers. For simplicity, we first consider the thermal flow in a slowly rotating ring 121 with trivial diffusivity  $D \rightarrow 0$ . The convection-diffusion equation is expressed into 122  $\frac{\partial T}{\partial t} = D \frac{\partial^2 T}{\partial x^2} + v \frac{\partial T}{\partial x}$ . Since the diffusion term is a perturbation compared to the advection 123 term, the convection-diffusion equation can be reduced into a homogeneous form  $\frac{\partial T}{\partial t}$  = 124  $v \frac{\partial T}{\partial x}$ , which has a wave form solution  $T(x, t) = Ae^{i(kx - \omega t)}$ . Substituting the wave form 125 126 solution back into the inhomogeneous convection-diffusion equation, we will end up with  $\omega = -ik^2D + kv$ . In poor thermal conducting materials, the initial temperature 127 128 field profile keeps unchanged during the rotation. After one circle, the field coincides, 129 where we define the perimeter  $L = 2\pi R_1$  as the periodic wavelength of the circulating energy packets. Straightforwardly, the wave number k in the wave form solution can 130 be defined as  $k = 2\pi/L = 1/R_1$ . Equation (1) actually takes a similar form to the 131

132 time-independent Schrödinger equation  $H\psi(x) = E\psi(x)$ , where the Hamiltonian is

133 
$$H_{0} = \begin{pmatrix} S_{0} - kv & ih & 0 & 0\\ ih & S_{0} - ih + kv & ih & 0\\ 0 & ih & S_{0} - ih - kv & ih\\ 0 & 0 & ih & S_{0} + kv \end{pmatrix},$$
 (2)

134 with  $S_0 = -i(k^2D + h)$  and  $h = \kappa_0 / \rho_r C_r b d$ . After some derivations, eigenvalues of

the Hamiltonian in Eq. (2) are solved by

136  

$$\omega_{\pm 1} = -i \left[ k^2 D + h \pm \sqrt{h^2 - (kv)^2} \right],$$

$$\omega_{\pm 2} = -i \left[ k^2 D + 2h \pm \sqrt{2h^2 - (kv)^2} \right].$$
(3)

In Eq. (3), the imaginary part of eigenvalues is mainly determined by the coupling strength *h*, which characterizes the decay rate of thermal energy. The eigenvalues will turn into complex ones, on condition that the effect of convection outweighs the coupling action. As shown by Eq. (3),  $\omega_{\pm 1}$  and  $\omega_{\pm 2}$  are complex with real and imaginary parts, when h < kv and  $\sqrt{2}h < kv$ , respectively. Eigenvectors of the Hamiltonian corresponding to  $\omega_{\pm 1}$  and  $\omega_{\pm 2}$  are derived to be

$$u_{\pm 1}(k) = \begin{pmatrix} -\frac{-h^2 + 2k^2 v^2 \pm 2ikv\sqrt{h^2 - k^2 v^2}}{h^2}, -\frac{-ikv \pm \sqrt{h^2 - k^2 v^2}}{h}, \\ -\frac{-ikv \pm \sqrt{h^2 - k^2 v^2}}{h}, 1 \end{pmatrix}^T, \\ u_{\pm 2}(k) = \begin{pmatrix} -1, \frac{h + ikv \pm \sqrt{2h^2 - k^2 v^2}}{h}, \frac{-h + ikv \pm \sqrt{2h^2 - k^2 v^2}}{h}, 1 \end{pmatrix}^T.$$

$$(4)$$

Figure 1(c) shows the relation between the eigenvalues and the flow velocity. In 144 the calculation, we set  $R_1 = 0.1$  m,  $R_2 = 0.11$  m, d = 5 mm, b = 1 mm,  $\kappa_r =$ 145 100 W/(m  $\cdot$  K),  $\kappa_o = 1$  W/(m  $\cdot$  K),  $\rho_r = \rho_o = 1000$  kg/m<sup>3</sup> and  $C_r = C_o =$ 146 1000 J/(kg · K). Thus we have  $k = 1/R_1 = 10 \text{ m}^{-1}$ ,  $h = \frac{\kappa_0}{\rho_r C_r b d} = 0.2 \text{ s}^{-1}$  and 147  $D = \kappa_r / \rho_r C_r = 10^{-4} \text{ m}^2/\text{s}$ . In Fig. 1(c), there exist two EPs in the spectrum, as 148 149 marked by the red dots A and B. From Eqs. (3) and (4), we obtain that the degenerated point A at v = h/k is a third order EP with the eigenvalues being  $\omega_{\pm 1,-2} =$ 150  $-i(k^2D+h)$  and  $\omega_{+2} = -i(k^2D+3h)$ . The corresponding eigenvectors are 151  $u_{\pm 1,-2} = (-1, i, i, 1)^T$  and  $u_{+2} = (-1, 2 + i, -2 + i, 1)^T$ , respectively. The 152 degenerated point **B** at  $v = \sqrt{2} h/k$  is a typical two order EP with the eigenvalues 153  $\omega_{\pm 1} = -i(k^2D + h) \pm h$  and  $\omega_{\pm 2} = -i(k^2D + 2h)$ . The eigenvectors are  $u_{\pm 1} =$ 154

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$$\left[-3 \pm 2\sqrt{2}, i(\mp 1 + \sqrt{2}), i(\mp 1 + \sqrt{2}), 1\right]^T$$
 and  $u_{\pm 2} = \left[-1, 1 + i\sqrt{2}, -1 + i\sqrt{2}, 1\right]^T$ .  
156 Here, it should be mentioned that the observed field evolution in diffusive systems  
157 eventually follows the minimum loss route, where the high-loss eigenfields will damp  
158 rapidly. As a result, in Fig. 1(c), the whole system would take the branch of  $\omega_{-1}$ , as  
159 marked by the brown lines. Therefore, as the flow velocity increases, the system will  
160 only experience the third order EP at  $v_{\rm EP} = h/k = 2$  cm/s, where the APT breaking  
161 phase transition occurs. Here we emphasized that the originally degenerated real parts  
162 of eigenvalues split into the upper and lower branches after the EPs, where we take the  
163 lower branch as  $\omega_{-1,-2}$  for consistency, as shown in Fig. 1(c) and the following.

164

# 165 **3. Results and discussion**

166 When the eigenvectors u satisfy  $\hat{P}\hat{T}u = \pm u$ , the whole system operates in the 167 PT/APT symmetric phase and will reach a steady state over time under the PT/APT 168 symmetry protection.<sup>22</sup> When  $\hat{P}\hat{T}u \neq \pm u$ , the system is operating in the PT/APT 169 broken phase with unstable eigenstates.

170 **3.1 Phase transition at the high order EP** 

171 From the Eq. (3), for  $v < v_{EP}$ , all eigenvalues are pure imaginary. Substituting 172  $\phi = \arcsin(v/v_{EP})$  into Eq. (4), the eigenvectors can be further simplified into

$$u_{\pm 1}(k) = (\cos 2\phi \mp i \sin 2\phi, \mp \cos \phi + i \sin \phi, \mp \cos \phi + i \sin \phi, 1)^{T},$$
  

$$u_{\pm 2}(k) = (-1, (1 \pm \sqrt{2 - \sin \phi^{2}}) + i \sin \phi, -(1 \pm \sqrt{2 - \sin \phi^{2}}) + i \sin \phi, 1)^{T}.$$
 (5)

174 Combining the forward and backward wave form solutions, we will obtain expressions

175 of eigenstates 
$$T = [T_1, T_2, T_3, T_4] = e^{-i\omega t} [u(k)e^{ikx} + u(-k)e^{-ikx}]$$
, where  $T_{1,2,3,4}$ 

# denote the steady temperature profiles in channels 1-4. All the possible eigenstates with

177 eigenfields in the four channels are derived as follows

$$T_{+1} = 2e^{-i\omega_{+1}t} [\cos(kx - 2\phi), -\cos(kx - \phi), -\cos(kx - \phi), \cos kx],$$
  

$$T_{-1} = 2e^{-i\omega_{-1}t} [\cos(kx + 2\phi), \cos(kx + \phi), \cos(kx + \phi), \cos kx],$$
  

$$T_{+2} = 2e^{-i\omega_{+2}t} \begin{bmatrix} -\cos kx, (1 + \sqrt{2 - \sin^2 \phi})\cos kx - \sin \phi \sin kx, \\ -(1 + \sqrt{2 - \sin^2 \phi})\cos kx - \sin \phi \sin kx, \cos kx \end{bmatrix},$$
  

$$T_{-2} = 2e^{-i\omega_{-2}t} \begin{bmatrix} -\cos kx, (1 - \sqrt{2 - \sin^2 \phi})\cos kx - \sin \phi \sin kx, \\ -(1 - \sqrt{2 - \sin^2 \phi})\cos kx - \sin \phi \sin kx, \cos kx \end{bmatrix}.$$
(6)

Note that the whole system ends up with the minimum loss case of  $T_{-1}$  over time, 179 showing that the steady-state temperature fields will stand still and phase differences 180 181 between  $T_1(T_4)$  and  $T_2(T_3)$  are  $\pm \phi = \pm \arcsin(v/v_{\text{FD}})$ . This claim is demonstrated 182 from the full wave simulation results by using a finite element solver COMSOL Multiphysics<sup>®</sup> 5.3, as shown in Fig. 2(a), where the initial temperature fields in all 183 channels are set by  $T_{1,2,3,4} = 293.15 + 100y$  (K). At  $v = v_{EP}$ , namely, the third order 184 transition point of eigenstates from APT symmetry to APT symmetry breaking, the 185 temperature fields in each channel keep standing still and the phase difference between 186 adjacent channels is  $\pi/2$ . To be specific, Fig. 2(b) shows the eigenfields distribution 187  $T_{1,2,3,4}$  in the four eigenstates  $T_{\pm 1,-2} \sim -2[\cos kx, \sin kx, \sin kx, -\cos kx]$  and  $T_{\pm 2} \sim -2[\cos kx, \sin kx, \sin kx, -\cos kx]$ 188  $2[\cos kx, -\sqrt{5}\cos(kx+\theta-\pi/2), \sqrt{5}\cos(kx-\theta-\pi/2), -\cos kx] \quad (\theta = \arctan 1/2).$ 189 For  $v > v_{EP}$ , the whole system transits into the APT breaking phase, where the 190 evolution of temperature profiles becomes very complicated. Intuitively, the convection 191 192 effect outperforms thermal coupling, making the system impossible to reach a steady 193 state. Substituting  $\psi = \operatorname{arccosh}(v/v_{\rm EP})$  into Eq. (4), we will derive the eigenstates

194 as

$$195 \quad T_{+1} = e^{\operatorname{Im}\omega_{+1}t} \begin{bmatrix} -e^{-2\psi}\cos(kx - \operatorname{Re}(\omega_{+1})t) - e^{2\psi}\cos(kx + \operatorname{Re}(\omega_{+1})t), \\ e^{-\psi}\cos(kx - \operatorname{Re}(\omega_{+1})t) + e^{\psi}\cos(kx + \operatorname{Re}(\omega_{+1})t), \\ e^{-\psi}\cos(kx - \operatorname{Re}(\omega_{+1})t) + e^{\psi}\cos(kx + \operatorname{Re}(\omega_{+1})t), \\ cos(kx - \operatorname{Re}(\omega_{+1})t) + cos(kx + \operatorname{Re}(\omega_{+1})t), \\ cos(kx - \operatorname{Re}(\omega_{-1})t) - e^{-2\psi}\cos(kx + \operatorname{Re}(\omega_{-1})t), \\ e^{\psi}\cos(kx - \operatorname{Re}(\omega_{-1})t) + e^{-\psi}\cos(kx + \operatorname{Re}(\omega_{-1})t), \\ e^{\psi}\cos(kx - \operatorname{Re}(\omega_{-1})t) + e^{-\psi}\cos(kx + \operatorname{Re}(\omega_{-1})t), \\ e^{\psi}\cos(kx - \operatorname{Re}(\omega_{-1})t) + cos(kx + \operatorname{Re}(\omega_{-1})t), \\ cos(kx - \operatorname{Re}(\omega_{-1})t) + cos(kx + \operatorname{Re}(\omega_{-1})t), \\ cos(kx - \operatorname{Re}(\omega_{-1})t) + cos(kx + \operatorname{Re}(\omega_{-1})t), \\ cos(kx - \operatorname{Re}(\omega_{-1})t) - cos(kx + \operatorname{Re}(\omega_{+2})t), \\ cos(kx - \operatorname{Re}(\omega_{+2})t) - cos(kx + \operatorname{Re}(\omega_{+2})t), \\ -\cos(kx - \theta_{1} - \operatorname{Re}(\omega_{+2})t) - cos(kx - \theta_{1} + \operatorname{Re}(\omega_{+2})t), \\ cos(kx - \operatorname{Re}(\omega_{+2})t) + cos(kx + \operatorname{Re}(\omega_{+2})t), \\ cos(kx - \operatorname{Re}(\omega_{+2})t) - cos(kx + \operatorname{Re}(\omega_{+2})t), \\ cos(kx - \operatorname{Re}(\omega_{+2})t) - cos(kx + \operatorname{Re}(\omega_{+2})t), \\ cos(kx - \operatorname{Re}(\omega_{+2})t) - cos(kx + \operatorname{Re}(\omega_{+2})t), \\ cos(kx - \operatorname{Re}(\omega_{-2})t) - cos(kx + \operatorname{Re}(\omega_{-2})t), \\ cos(kx - \operatorname{Re}(\omega_{-2})t) - cos(kx - \operatorname{Re}(\omega_{-2})t), \\ -cos(kx -$$

$$L \cos(kx - \operatorname{Re}(\omega_{-2})t) + \cos(kx + \operatorname{Re}(\omega_{-2})t) \qquad J$$
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$$\left(\tan\theta_2 = \frac{\cosh\psi}{1 - \sqrt{2 - \cosh^2\psi}}\right).$$
(7)

In inspection of Eq. (7), the amplitudes of forward and backward wave form  
components in the eigenstate 
$$T_{-1}$$
 are  $A_{-1,1} = -e^{2\psi}$ ,  $B_{-1,1} = -e^{-2\psi}$ ,  $A_{-1,2} = e^{\psi}$ ,  
 $B_{-1,2} = e^{-\psi}$ ,  $A_{-1,3} = e^{\psi}$ ,  $B_{-1,3} = e^{-\psi}$  and  $A_{-1,4} = B_{-1,4} = 1$ . Obviously, the  
temperature fields  $T_{1,2,3,4}$  are unstable over time evolution. Here we utilize the local  
maximum to trace the position (or phase) of temperature fields in each channel. From  
 $\partial T_j / \partial x = 0$  and Eq. (7), we will obtain the relation  $\tan kx_j = \tan[\operatorname{Re}(\omega)t](A_{-1,j} - B_{-1,j})/(A_{-1,j} + B_{-1,j})$ . Define the phase difference between channels 1 and 2 as  
 $\varphi_1 - \varphi_2 = k(x_1 - x_2)$ . Considering  $\tan k(x_1 - x_2) = \frac{\tan kx_1 - \tan kx_2}{1 + \tan kx_1 \cdot \tan kx_2}$ , we have  
 $\varphi_1 - \varphi_2 = \arctan \frac{S_1 \tan[\operatorname{Re}(\omega)t]}{1 + S_2 \tan^2[\operatorname{Re}(\omega)t]}$ , (8)  
where  $S_1 = \frac{A_{-1,1} - B_{-1,1}}{A_{-1,2} + B_{-1,2}}$ ,  $S_2 = \frac{A_{-1,1} - B_{-1,1}}{A_{-1,2} + B_{-1,2}}$ . Figures 2(c) and 2(d)

211 show the theoretical calculation and numerical simulation of the phase difference  $\varphi_1$  –

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212  $\varphi_2$  over time evolution, respectively. The results unequivocally reveal that the phase

- 213 difference  $\varphi_1 \varphi_2$  keeps increasing and will never reach a steady state on condition
- that APT symmetry is breaking.



Fig. 2 APT symmetry breaking at the high order EP in the four-channel diffusive system. (a) The time evolution of phase differences  $\varphi_1 - \varphi_2$  and  $\varphi_4 - \varphi_3$  vs. the flow velocity v, where we set  $v_n = v_{EP} \sin\left(\frac{n\pi}{18}\right), (n = 1, 2, ..., 9)$ . (b) The eigenstates  $T_{\pm 1}$  and  $T_{\pm 2}$  at the EP. In (a) and (b),  $T_{1,2,3,4}$  and  $\varphi_{1,2,3,4}$  denote the steady temperature profiles and the related phases in channels 1-4. (c), (d) Theory and simulation results of time evolution of  $\varphi_1 - \varphi_2$  at  $v = 2\sqrt{2}$  cm/s, when the APT symmetry is broken.

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#### **3.2 Perturbation at the high order EP**

In this section, we explore the case that a perturbation is introduced to the high order EP. Here we introduce a flow velocity modulation  $\pm \Delta v$  into the four-channel system, as schematically shown in Fig. 3(a). In this case, the Hamiltonian containing the perturbation is rewritten as follows

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$$H_{1} = \begin{pmatrix} S_{0} - (kv_{\rm EP} - \Delta v) & ih & 0 & 0\\ ih & S_{0} - ih + (kv_{\rm EP} + \Delta v) & ih & 0\\ 0 & ih & S_{0} - ih - (kv_{\rm EP} + \Delta v) & ih\\ 0 & 0 & ih & S_{0} + (kv_{\rm EP} - \Delta v) \end{pmatrix}.$$
230 (9)

The imaginary part and the real part of the Hamiltonian are shown in Figs. 3(b) 231 and 3(c). The result shows that as the modulation strength  $\Delta v$  increases from 0 cm/s, 232 233 there exist three EPs. One is the third order EP A' at  $\Delta v = 0$  cm/s, while the typical two order EPs **B**' and **C**' locate at  $\Delta v = 0.57$  cm/s and  $\Delta v = 1.6$  cm/s. As we 234 235 mentioned before, the diffusive system is always following the lowest dissipation state over time. Therefore, the eigenvalue of the Hamiltonian takes the brown branch in Figs. 236 3(b) and 3(c). The results show that the APT symmetry is robust against the weak 237 perturbation at the high order EP. When the perturbation strength is above the threshold 238 239  $\Delta v = 1.6$  cm/s, the whole system will transit into APT symmetry breaking across the EP C' and the temperature fields become unstable. Figure 3(d) displays the time 240 evolution of phase difference  $\varphi_1 - \varphi_2$  at different modulation strengths, where the 241 242 APT is symmetric. In Fig. 3(d), we find that the phases of steady-state temperature profiles in adjacent circulating channels are relatively static (t > 60 s), as manifested 243

in a locked mode with a nearly constant phase difference. To be specific, the phase differences of three locked modes (in degrees) are 55° for  $\Delta v = 0.5$  cm/s, 41.2° for

246  $\Delta v = 1 \text{ cm/s}$ , and 25.6° for  $\Delta v = 1.5 \text{ cm/s}$ , respectively.



Fig. 3 Perturbation at the high order EP. (a) The schematic of flow velocity modulation in the four channels. (b), (c) The imaginary part and real part of eigenvalues *vs.* the flow velocity modulation  $\Delta v$ , with the brown lines denoting the scenario of the lowest energy dissipation. Red dots at **A'**, **B'**, and **C'** mark the EPs. (d) The phase difference  $\varphi_1 - \varphi_2$  between channels 1 and 2 over time evolution in APT symmetry, where  $\Delta v = 0.5$  cm/s, 1.0 cm/s, 1.5 cm/s.

As aforementioned, the temperature fields in diffusive systems are unstable in the APT broken phase. Here we show that by introducing flow velocity modulation at the

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high order EP, the phase difference between adjacent channels will oscillate over time
instead of continuously diverging at APT breaking. In addition, the phase oscillation at
APT breaking is independent with the initial condition. In Figs. 4(a)-4(c), we present a
numerical demonstration of the phase oscillation effect at APT breaking with three
different initial conditions. Specifically, in Fig. 4(a), the temperature profile at $t = 0$ s

261	is $T_{1,3} = 293.15 + 100x$ (K) and $T_{2,4} = 293.15 + 100y$ (K), with $\varphi_1 - \varphi_2 =$
262	$-90^{\circ}$ . In Fig. 4(b), the temperature profile at $t = 0$ s is $T_{1,2,3,4} = 293.15 + 1000$
263	100y (K), with $\varphi_1 - \varphi_2 = 0^\circ$ . In Fig. 4(c), the temperature profile at $t = 0$ s is
264	293.15 - 100x (K) and $T_{2,4} = 293.15 + 100y$ (K), with $\varphi_1 - \varphi_2 = 90^\circ$ . Here
265	the flow velocity modulation strength $\Delta v = 4$ cm/s. The results reveal that for the
266	three cases, phase oscillation occurs at $t > 40$ s, with the oscillation center angle
267	$\varphi_{\text{center}} = 45^{\circ}$ and the time cycle $T_{\text{period}} = 32$ s. In Figs. 4(a) - 4(c), the insets show
268	the temperature profiles in channels 1 and 2 at different times when the phase difference
269	$\varphi_1 - \varphi_2$ takes maximum or minimum values, vividly displaying the relative periodic
270	oscillation of phase difference over time. Figure 4(d) shows that both the oscillation
271	center angle and the time cycle decrease as the flow velocity modulation strength
272	increases. When the flow modulation strength $\Delta v$ is much larger than the EP velocity
273	$v_{\rm EP}$ , for example $\Delta v = 40$ cm/s, the flows in channels 1 and 2 can be regarded as
274	almost synchronous circulation. Therefore, the phase oscillation is weak with the center
275	angle close to 0 ( $\varphi_{center} = 1.4^\circ$ ), while the time cycle of the phase jittering $T_{period} =$
276	21.6 s.



Fig. 4 Periodic phase oscillation in APT symmetry breaking. (a), (b) and (c) Phase oscillation of  $\varphi_1 - \varphi_2$  at  $\Delta v = 4$  cm/s with the initial condition being  $\varphi_1 - \varphi_2 = -\pi/2$ , 0 and  $\pi/2$  at t =0 s, respectively. (d) The oscillation center angles/time cycles *vs*. the flow velocity modulation  $\Delta v$ .

At last, we would like to briefly discuss the cases of more channels coupled, odd-282 number channels coupled and the suggestions for experiment. When more channels are 283 coupled, we can basically obtain four-order EPs and beyond. In this case, the technical 284 difficulty for experimental observation is increasing. Higher ordered EPs are intuitively 285 associated with very complicated behaviors in phase evolution. However, the properties 286 protected by APT symmetry are expected to be unchanged. For example, the phase 287 288 differences of temperature profiles between adjacent channels for the steady states are locked in the APT symmetric case. Phase oscillation at APT symmetry breaking is also 289

supposed to occur when the higher ordered EP is perturbed with velocity modulation. 290 In this work, we focus on the model of even-number coupled channels with inside flows 291 having equal-but-opposite velocities. It will be interesting if an additional channel is 292 introduced to break the symmetry of the whole system (*i.e.*, odd-number channels). In 293 294 light of the Hamiltonian analysis, there will exist an isolated branch with no conjugate 295 pair in the eigenvalue spectrum, which does not degenerate with other paired branches. When the eigenmodes on the isolated branch have the minimum loss, the system will 296 follow this single branch and do not experience EP-induced phase transition. For the 297 experimental demonstration, we can use nylon rings ( $\kappa_r < 2 W/(m \cdot K)$ ) and grease 298  $(\kappa_0 \approx 0.3 \text{ W}/(\text{m} \cdot \text{K}))$  as the channels and coupling layers, respectively. The flow 299 300 velocity is implemented by rotating the rings with motors, where the rotation speed can be accurately controlled. The initial temperature fields are added to the rings through a 301 302 copper plate with the ends immersing into the hot and cold water baths. Meanwhile, the 303 copper plate is closely contacting with the rings.

304

#### 305 4. Conclusions

In summary, we comprehensively investigate the phase evolution in a four-channel coupling-chain diffusive model. We show that a third order EP is generated in the eigenspectrum of the Hamiltonian for the four-channel toy model with the background flow velocities in adjacent channels being opposite. The high order EP is featured with APT phase transition. At the APT symmetry, we theoretically derive and numerically

verify the phase differences of the steady-state temperature profiles between adjacent 311 channels, where the flow velocity is below the threshold  $v_{\rm EP}$ . The divergence of phase 312 difference is also verified in the unstable diffusive system at APT symmetry breaking. 313 314 More interesting is to introduce a perturbation into the background flow velocity at the 315 high order EP. We find that the APT symmetry is robust against weak perturbation, 316 where the system operates in a locked mode with the phase difference of moving temperature profiles in adjacent channels relatively unchanged. As the modulation is 317 318 strengthened to surpass a threshold (another EP), the four-channel system will transit into the broken APT symmetry. In APT breaking, the phenomenon of robust phase 319 oscillation is observed, which is irrelevant to the initial conditions. This work marries 320 321 the two scenarios of diffusive systems and high order EP physics, which paves the way 322 of counterintuitive thermal flow regulation via phase transition in coupling-chain 323 diffusive systems.

324

## 325 **Conflict of interest**

326 The authors declare no conflict of interest.

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371