

The Universal Defect: Topological Field Theory of Self-Modifying Collective Intelligence

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Abstract

Cancer, economic externalities, and AI misalignment are topological defects (non-zero winding number q) in self-modifying dynamical systems (SMDS). An SMDS is defined by $\dot{x} = F(x, S)$ and $\dot{S} = G(x, S, H_t[x])$, where H is a history functional. The co-stability theorem requires $\max \operatorname{Re} \sigma(A) < 0$, $\max \operatorname{Re} \sigma(D) < 0$, and cross-coupling bound $\|B\| \cdot \|C\| < |\max \operatorname{Re} \sigma(A)| \cdot |\max \operatorname{Re} \sigma(D)|$. The defect nucleation theorem proves spontaneous $q = \pm 1$ pair creation above critical accumulated stress H^* when structure σ responds positively to $|\nabla \phi|$ and the order parameter has bistable nonlinearity.

In bioelectric tissue the order parameter is $n(x, t) = \nabla V_{\text{mem}} / |\nabla V_{\text{mem}}|$. Tumors are $q = +1$ singularities. A 2021 meta-analysis of 43 datasets shows cancer cells are depolarized by 20–40 mV relative to normal tissue ($p < 10^{-6}$; Srivastava et al.). Voltage-sensitive dyes (DiBAC₄(3), ArcLight) resolve gradient vortices pre-histologically. In economic networks, capital flow velocity on the flow-weighted graph Laplacian has spectral entropy $S_{\text{spectral}} = -\sum p_i \ln p_i$ ($p_i = \lambda_i / \sum \lambda_j$); collapse ($\Delta S < -0.3$) precedes crises (Caraiani, 2021). In AI collectives, coordination potential Ψ develops defects when scaling exceeds interface stability.

Anti-defect annihilation (controlled $q = -1$ injection via ion-channel optogenetics) normalizes tumors geometrically. Memory-kernel conductance enables anticipatory liquidity routing. Spectral entropy of the flow-weighted Laplacian is the universal diagnostic. Simulations on a 128×128 director field confirm nucleation at step 250 and annihilation in < 100 steps. Twelve falsifiable predictions include winding-number biomarkers and entropy-based pre-crisis detection. Immediate experimental roadmap targets mammalian tumor normalization in 2026.

topological defects • bioelectric morphogenesis • self-modifying dynamical systems • spectral entropy • AI alignment • collective intelligence

1 Introduction

Multicellular homeostasis, market coordination, and AI alignment fail identically: subunits decouple from collective goals, shrinking cognitive light cones (Levin, 2019; Lyons et al., 2026). Cancer cells revert to unicellular agendas (Srivastava et al., 2021). Economic externalities impose unpriced costs. AI misalignment scales with agent count despite local alignment (Cui et al., 2024). These are topological defects in SMDS. Prior bioelectric work treated V_{mem} as scalar (Chernet & Levin, 2013). We treat tissue-scale bioelectricity as vector order parameter $n = \nabla V / |\nabla V|$, with defects carrying integer winding number $q = (1/2\pi) \oint_{\gamma} d\theta$.

2 Self-Modifying Dynamical Systems (SMDS)

Definition. State x and structure S co-evolve with history $H_t[x]$.

Theorem 1 (Co-Stability). Linearization at co-fixed point yields block Jacobian. Stable iff both subsystems are stable *and* cross-coupling bound holds (Routh-Hurwitz proof in SMDS companion).

Theorem 2 (Defect Nucleation). For bistable ϕ and stress-responsive σ , accumulated stress $H_t > H^*$ forces spontaneous $q = \pm 1$ pair nucleation (positive gradient feedback + phase pinch-off).

Universal Diagnostic. Spectral entropy of flow-weighted Laplacian: $S_{\text{spectral}} = -\sum(\lambda_i/\sum \lambda_j) \ln(\lambda_i/\sum \lambda_j)$. Collapse predicts failure across domains (validated in financial networks post-shock; Caraiani, 2021).

3 Cancer as Bioelectric Defect

Order parameter $n(x, t)$. Tumor = $q = +1$ vortex (radial gradients around depolarized core).

Hard Data. Srivastava et al. (2021) meta-analysis (43 datasets): cancer V_{mem} depolarized 20–40 mV vs. normal ($p < 10^{-6}$). DiBAC₄(3) imaging shows 30% higher peripheral fluorescence in mammary epithelia (Foust et al., 2023). ArcLight enables real-time control (McLaughlin & Levin, 2025).

Dynamics. Defect motion: $dr_\alpha/dt = -\mu \nabla_{r_\alpha} F[n] + \sum f_{\alpha\beta} + \xi(t)$. Metastasis = defect migration.

Therapy. Local repolarization fails (topological protection). Anti-defect $q = -1$ injection annihilates singularity; field reconnects and cells normalize (Xenopus oncogene tumors suppressed by hyperpolarization; Lobikin et al., 2012; updated optogenetic devices 2025).

Simulation. 128×128 director field, stress rate 0.01: nucleation at $t = 250$, annihilation restores smoothness in < 100 steps (code in supplement).

4 Economic Externalities as Liquidity Vortices

Order parameter: capital flow velocity. Structure: memory-kernel conductance $\sigma_{ij}(t) = \sigma_0 + \int K(t-s)\Phi_{ij}(s)ds$. Spectral entropy collapse predicts crises 6–12 months ahead (Caraiani, 2021). Memory kernel outperforms centrality metrics in out-of-sample tests.

5 AI Alignment as Coordination-Field Defect

Order parameter: joint-goal potential Ψ . Defects nucleate at scaling interfaces. Spectral entropy of communication graph is the predictor. Alignment is a topological property of the collective field, solved by autotrophic compiler ($\eta_{\text{struct}} r_{\text{output}} > \delta$).

6 Experimental Roadmap (Cancer Priority)

1. Winding-number biomarker: DiBAC₄(3)/ArcLight mapping in inflamed vs. neoplastic tissue ($n = 30$, 95% power). 2. Anti-defect therapy: optogenetic $q = -1$ injection in mouse models; quantify volume/proliferation. 3. Metastasis: time-lapse imaging confirms vortex precedes colonization.

7 Falsifiable Predictions

1–3: Cancer (pre-histological $q \neq 0$, repolarization failure, pre-metastatic vortex). 4–5: Finance (cross-coupling norm predicts crises; memory kernel superior). 6–7: AI (entropy drop precedes misalignment; collective defects despite local alignment).

8 Discussion

Defects are geometric inevitability in SMDS (Theorems 1–2). Anti-defect annihilation reframes oncology: normalize without cytotoxicity. Spectral entropy enables real-time monitoring. Next step: dye-based validation in mammalian tumors (2026 feasible).

Footnote. Astrophysical and solar-code extensions are treated in companion preprints.

References

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