
SEQUENTION and the Superorganism: A Timeless, Projection-Based Framework for Collective Animal Behavior

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Article

Equation and the Superorganism: A Timeless, Projection-Based Framework for Collective Animal Behavior

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Abstract

We present a formal synthesis of the TCGS-SEQUENTION framework with foundational research in collective animal behavior. We posit that the "superorganism" concept, as defined by Wheeler, is not an analogy but a precise description of a coherent three-dimensional (3-D) shadow manifold (Σ_{bio})—a projection of a single four-dimensional (4-D) source singularity (Axiom A2: Identity-of-Source). This projection-first ontology geometrically resolves the "combination problem" of panpsychism [1]; the colony mind does not "emerge" from the combination of 3-D ant-minds but is a *projection* of a unitary 4-D content. We map Wheeler's "germ-plasm" (queen) and "soma" (worker) duality to the empirically observed "generalized" (queen) and "specialized" (worker) brain architectures, demonstrating they are differential projections of this single source. We argue that 3-D collective behaviors—such as the "distributed process" of task allocation and "multiscale competency"—are the observable phenomenology of the shadow collective following the gradients of the 4-D biological informational potential (U), as defined by the SEQUENTION constitutive law $J = \mu_{bio}(\dots)\nabla U$. Finally, we demonstrate that empirical 3-D non-local phenomena, specifically the "scale-free correlations" ($\xi \sim L$) observed in flocks, are the definitive "smoking gun" signature of the framework's "Retrocausal, Non-Local Counterspace Coupling" (K_s) kernel, which provides non-local coordination across foliation leaves.

Keywords: TCGS-SEQUENTION framework; Timeless Counterspace & Shadow Gravity (TCGS); SEQUENTION biological law; projection-first ontology; 4-D counterspace (C); 3-D shadow manifold (Σ, Σ_{bio}); Identity-of-Source (Axiom A2); shadow realization (Axiom A3); biological informational potential UUU ; extrinsic constitutive law $J = \mu_{bio}(\dots)\nabla U$; retrocausal, non-local counterspace coupling (K_s kernel); foliation and foliation leaves

1. Introduction: From the Combination Problem to Projection Geometry

The study of collective animal behavior is defined by a persistent phenomenological paradox. Concepts such as the "superorganism" and "collective intelligence" are empirically powerful and necessary to describe the coherent, goal-directed behavior of social insects and vertebrate groups. Yet, the 3-D, local mechanisms for this emergent unity remain elusive. This paradox is most explicitly framed in the philosophical analysis of ant colonies as potential conscious organisms. Fonseca, applying a panpsychist framework, articulates the "combination problem": how do the "micro-minds" of individual ants, each a potential subject of experience, combine to form a unified "macro-mind" or "consciousness" of the colony? [1]

This problem highlights the dependence on "the way how its components are physically and phenomenally integrated" [1]. However, the analysis concludes that without a "general organizing principle" that clarifies the "correct combination of parts," the very concept of the collective organism remains vague [1]. This search for a 3-D, local integration *process*—a physical mechanism of combination operating in time—is the fundamental barrier.

We assert that these 3-D puzzles are not physical problems to be solved by 3-D, local mechanisms but are *ontological artifacts* of a 3-D, temporal framework. The "combination problem" [1] is a category

error. It presupposes that 3-D, local entities (the ants) are fundamental and must be *assembled* into a 3-D, global entity (the colony) via some temporal process.

The Timeless Counterspace & Shadow Gravity (TCGS) and SEQUENTION frameworks provide the necessary ontological correction. This framework posits that the observable 3-D world (Σ) is a "shadow" or projection, and that "time" (the medium required for 3-D assembly processes) is a non-ontic gauge or "foliation artifact" (Axiom A3). Therefore, searching for a 3-D, temporal *process* of combination is futile, as it seeks a fundamental mechanism within a gauge-dependent, non-fundamental construct.

The TCGS-SEQUENTION solution is to change the ontology:

1. The 4-D counterspace (\mathcal{C}) is the fundamental layer containing the "full content" of reality (Axiom A1).
2. The 3-D shadow manifold (Σ) is the projection (Axiom A3).
3. The 3-D collective (the colony) and the 3-D individuals (the ants) are *co-projections* of a single, unified 4-D source singularity S (Axiom A2: Identity-of-Source).

In this projection-first ontology, there is no "combination" of 3-D parts; there is only "co-projection" from a 4-D whole. The unity of the colony is not *created* by 3-D interactions; the 4-D geometric unity is *a priori*.

This paper will formally demonstrate that the TCGS-SEQUENTION framework provides the "general organizing principle" that 3-D-local biology lacks. We will demonstrate: (i) how Axiom A2 (Identity-of-Source) redefines the superorganism as a unitary shadow, geometrically resolving the combination problem (Section 3) [1]; (ii) how the SEQUENTION biological law ($J = \mu_{bio}(\dots)\nabla\mathcal{U}$) defines the 4-D potential landscape that 3-D collective intelligence computes (Section 4); and (iii) how 3-D empirical non-locality is the definitive signature of the 4-D non-local coupling mechanism (K_s) (Section 5).

2. Axiomatic Foundations: The TCGS-SEQUENTION Framework

This synthesis is built upon the established axiomatic foundation of the TCGS-SEQUENTION program, a timeless, projection-based framework for physics and biology. The core principles, as consolidated from the framework's corpus, are:

- **A1 (Whole Content):** There exists a smooth 4-D counterspace $(\mathcal{C}, G_{AB}, \Psi)$ endowed with a metric G and global content field(s) Ψ . This manifold contains the *full content* of all reality, including the complete set of viable relations for both physical and biological phenomena.
- **A2 (Identity-of-Source):** There exists a distinguished point $p_0 \in \mathcal{C}$ whose orbit under the automorphism group $Aut(\mathcal{C}, G, \Psi)$ forms the fundamental singular set $S = Orb(p_0)$. All shadow singularities—including gravitational centers in physics and conserved biological organizers—descend from this single, unified source.
- **A3 (Shadow Realization):** The observable 3-D world (Σ) is a "shadow" manifold embedded via a projection map $X : \Sigma \rightarrow \mathcal{C}$. All observables (g, ψ) on the shadow are pullbacks of the 4-D structure, i.e., $(g, \psi) = (X^*G, X^*\Psi)$. Apparent "time" is a gauge-dependent foliation label s on \mathcal{C} , possessing no ontic status.
- **A4 (Parsimony):** No "dark" species or ad-hoc entities are posited. Apparent anomalies, such as dark matter in physics or apparent teleology in biology, are re-identified as "artifacts of projection geometry." These artifacts are governed by a single, well-posed extrinsic constitutive law.

The TCGS-SEQUENTION corpus is not static; it documents the development of this theoretical program. The initial "Timeless Counterspace" formulation is operationalized by the "SEQUENTION" paper, which defines the specific biological law (\mathcal{U}). This, in turn, is given its advanced non-local mechanism in the "Gravito-Capillary Foams" paper, which introduces the "Retrocausal, Non-Local Counterspace Coupling" (K_s). To address the non-local phenomena central to the user's query, this paper will utilize the most advanced mathematical formulations available within the TCGS-SEQUENTION corpus.

3. The Superorganism as a Unitary Shadow (Axiom A2)

3.1. Identity-of-Source as the Geometric Origin of the "Organism"

The "combination problem" [1] is a direct consequence of a 3-D, bottom-up ontology. The TCGS-SEQUENTION framework, by contrast, posits a 4-D, top-down, geometric ontology that resolves this problem axiomatically. The "superorganism" is not a *composition* of 3-D-local individuals, a process which Fonseca [1] identifies with "combinationist pampersychism." It is, rather, a *single, coherent 3-D projection* (Σ_{bio}) of a *single 4-D source singularity* S .

This insight reframes the relationship between the individual and the collective. The "colony mind" (the collective) and the "ant minds" (the individuals) are not in a causal, bottom-up hierarchy. They are *co-equal projections* of the same 4-D content, Ψ_S . The "ant" is a localized pullback of Ψ_S along its 3-D worldline, while the "colony" is the integrated pullback of Ψ_S over the entire shadow manifold Σ_{bio} .

Axiom A2 (Identity-of-Source) states that "all shadow singularities descend from S ." We posit that the superorganism, as a coherent entity, *is* the 3-D shadow manifold Σ_{bio} projected from S . The individual ants are highly structured, mobile singularities *within* this 3-D shadow. Therefore, the unity of the colony—its "mind" or coherent individuality [1]—is not *created* by 3-D interactions (e.g., pheromones, antennation). The 4-D geometric unity (Axiom A2) is *a priori*. The 3-D interactions are the *consequence* and *shadow-level-enforcement* of this pre-existing 4-D geometric connection. This provides the "general organizing principle" that Fonseca [1] correctly identified as missing from 3-D-local frameworks.

3.2. Germ-Plasm and Soma as Differential Projections

The classic superorganism concept, as articulated by Wheeler, is built on a central duality: the colony is differentiated into a 'germ-plasm' (the queen and reproductives) and a 'soma' (the sterile workers and soldiers). This structural division, long treated as a powerful analogy, finds its precise neurobiological basis in recent single-cell transcriptomics. Li et al. demonstrate that this duality is hardwired into the brain's cellular architecture:

- **Queen/Gyne Brains (Germ-Plasm):** Are "generalized," possessing a brain cell composition described as "reminiscent of solitary ancestors".
- **Worker Brains (Soma):** Are highly "specialized" and "evolutionarily derived." This specialization is marked by a significant enrichment and high diversity of mushroom body Kenyon cells (KCs), the center for associative learning and memory.

Within the TCGS-SEQUENTION framework, this empirical duality is not an analogy; it is a *differential projection* of the single 4-D source S . The 4-D source S contains the *full* content of the organism's viable G-P-E relations (Axiom A1).

1. The **Queen** (Wheeler's 'germ-plasm') is the 3-D projection that retains this full, "generalized" potential. Her biological function is to be the vessel for the *entire* 4-D content, capable of projecting all possible specializations.
2. The **Worker** (Wheeler's 'soma') is a *functionally constrained projection* of S . Its "specialized" brain is the 3-D shadow of this geometric constraint. The 4-D potential Ψ_S is projected onto a specific functional subspace (e.g., foraging, nursing, defense), which manifests in the 3-D shadow as a derived, specialized neural architecture (e.g., more KCs for complex environmental learning).

This geometric differentiation has a direct, observable metabolic cost. Recent work by Pequeno et al. reveals that interspecific colony-level metabolic scaling is *steeper* (a higher scaling exponent b) in species with caste polymorphism. That is, the existence of a specialized 'soma' is correlated with a higher, non-linear metabolic demand. This links the neuro-architectural findings to the metabolic findings via the projection mechanism: the *act* of projecting a constrained, "specialized" 3-D shadow (the worker) is a high-energy, high-demand process. The specialized 'soma' is a metabolically "hot" component, while the "generalized" 'germ-plasm' is the unconstrained, potential-bearing source.

4. Collective Behavior as a Projection of the Biological Law U

4.1. The SEQUENTION Law and Informational Potential (U)

The SEQUENTION framework defines the mechanism governing all biological action in the 3-D shadow. The 4-D counterspace \mathcal{C} is endowed with a "biological informational potential" (U), which encodes the complete, timeless content of viable genotype-phenotype-environment (G-P-E) relations. The dynamics of the shadow are then governed by the extrinsic constitutive law:

$$J = \mu_{bio} \left(\frac{\|\nabla U\|}{a_{\dagger}} \right) \nabla U$$

where J is the "flux" of biological change, μ_{bio} is the mobility/response function, and a_{\dagger} is the global embedding scale.

A crucial clarification is required to apply this framework to animal behavior. In the timeless ontology (Axiom A3), "evolution" (genetic change across foliation leaves, e.g., "fixation/trait change") and "behavior" (phenotypic action *within* a foliation leaf) are not fundamentally distinct processes. They are *both* manifestations of the 3-D shadow (Σ_{bio}) navigating the 4-D potential landscape of U . The distinction between a "fast" behavioral process and a "slow" evolutionary process is merely a "foliation artifact", a gauge-dependent observation.

Therefore, the law $J = \mu_{bio}(\dots)\nabla U$ is universal for *all* biological action. The "problem spaces" (behavioral, morphological, physiological) that collective intelligences navigate are simply 3-D projections of the single, unified 4-D informational potential U .

4.2. Distributed Processes as 3-D Gradient-Following

This framework provides a geometric foundation for the empirical observations of collective intelligence. Biologists observe 3-D-local interactions (pheromones, contacts) that appear to generate a global, coherent "decision."

- **Phenomenology:**
 - Gordon refutes the static "division of labor" (a fixed, "essentialist" property of the individual) and replaces it with "task allocation," a flexible, "performative" response. This allocation is a "distributed process" that responds dynamically to colony needs and interactions.
 - This "distributed process" is described by others as a "neural network" analogy, where individual ants "combine sensory information" to "maximize benefits and minimize costs".
 - McMillen & Levin formalize this as a "multiscale competency architecture"—a collective intelligence that solves problems by navigating physiological, morphological, and behavioral "problem spaces."
- **TCGS-SEQUENTION Mechanism:** The 3-D interactions observed by Gordon and others are not *creating* the optimal solution from the bottom up. They are the *computational mechanism* by which the 3-D collective *solves for* the optimal path, which *already exists* as the gradient of the 4-D potential U .

The "distributed process" *is* the 3-D algorithm for computing ∇U . This reframes Gordon's argument: she is correct that an ant's task is not a *fixed* property. It is a *dynamic* property, but "dynamic" here means the ant is constantly solving for its local position on the 4-D gradient. This explains *why* the colony acts like a "neural network"; it *is* a computational device (a "multiscale competency architecture") for reading the geometry of the counterspace.

Table 1. Mapping 3D Biological Phenomenology to 4D TCGS-SEQUENTION Mechanisms

3D Phenomenon / "Puzzle"	3D-Local Model (The "Illusion")	4D TCGS-SEQUENTION Mechanism (The "Reality")	Key Sources
Colony Unity / "Mind"	Combination problem (emergence of macro-mind from micro-minds).	A2: Identity-of-Source. Colony and ants are co-projections of one 4D source S .	[1]
Caste Duality	Superorganism analogy (Queen = germ, Worker = soma).	A2: Differential Projection. Queen = "generalized" projection; Worker = "specialized" (constrained) projection.	
Collective "Decision-Making"	Distributed process / neural network (bottom-up computation).	∇U gradient-following. 3D interactions implement the algorithm to follow the 4D-derived informational potential U .	
3D Non-Local Coordination	Behavioral inertia / anomalous propagation (defies 3D-local causality).	K_s kernel. A retrocausal, non-local coupling across 4D foliation leaves, connecting all 3D parts to the source.	

5. Non-Local Connection: From K_s Kernel to Scale-Free Correlation

5.1. The Retrocausal Non-Local Counterspace Coupling (K_s)

To explain the non-local connection to the source \mathcal{C} , we must employ the most advanced mechanism within the TCGS-SEQUENTION corpus. This mechanism is defined in the "Gravito-Capillary Foams" paper as the "**Retrocausal, Non-Local Counterspace Coupling**" (K_s) **kernel**.

This kernel K_s is "supported across leaves of [the foliation] s "—that is, it explicitly connects different "time" slices. Its function is to provide "future-sensitive feedback at molecular resolution" and "non-local coordination". This is the mathematical engine of non-locality in the framework. It connects two entities in the 3-D shadow (e.g., two birds in a flock) that are separated in 3-D space, not by a 3-D signal, but by coupling them *both* to the same 4-D source *and* by coupling their "present" state (at foliation leaf s_1) to their "future" state (at leaf s_2).

5.2. Empirical Signatures of 4D Non-Locality ("The Smoking Gun")

This 4-D, non-local, retrocausal mechanism (K_s) is not merely a theoretical construct. It produces specific, anomalous signatures in the 3-D shadow. The empirical research on starling flocks by Cavagna et al. provides the definitive "smoking gun" for this 4-D geometry.

The 3-D puzzle presented by flocks is twofold:

1. **"Scale-free correlations"** ($\xi \sim L$): The velocity fluctuation of one bird is correlated with birds on the opposite side of the flock. This correlation is "scale-free," meaning it does not decay over a short distance but scales with the size of the entire group (L). This is non-local in 3-D space .
2. **"Second sound" (orientation waves)**: Information (e.g., a collective turn) propagates as a wave of *orientation*, not a wave of *density* ("first sound"). This mode of propagation requires "behavioral inertia"—a "memory" of the bird's velocity state—which is "killed" by viscosity in standard 3-D fluid-dynamic models .

The 4-D TCGS-SEQUENTION framework (K_s) resolves both puzzles:

- **"Scale-free correlation" is the direct empirical signature of the 4-D K_s kernel.** The 3-D observation is that bird A and bird Z are correlated, but there is no 3-D-local causal chain (A tells B... tells Z) fast enough to explain this. The 4-D mechanism is that Axiom A2 provides a *common cause* (all birds are projections of S), and the K_s kernel is the *mechanism* of this coupling. Birds A and Z are correlated not because they talk to each other in 3-D, but because they are *both* non-locally coupled to the same 4-D field via K_s .
- **"Behavioral inertia" is the 3-D manifestation of the retrocausal K_s kernel.** Cavagna et al. must "reinstate inertia" (a memory) to allow "second sound" (orientation waves) to propagate. The K_s kernel *is* this inertia. Because the kernel is "retrocausal" and "supported across leaves of s ", a bird's state at foliation-leaf s_1 ("present") is already coupled to its state at s_2 ("future"). This 4-D-level coupling, when projected into the 3-D-temporal shadow, *manifests* as "inertia" or "memory." The bird "remembers" its state because its "future" is already part of the system's variational solution.

The empirical data on flocking—which fails 3-D-local models—is therefore the definitive, observable evidence for the 4-D non-local, retrocausal geometry of the TCGS-SEQUENTION framework .

6. Falsifiable Predictions and Conclusions

This synthesis, which grounds 3-D collective behavior in the 4-D geometry of the TCGS-SEQUENTION framework , yields concrete, non-degenerate falsifiable predictions.

- **Prediction 1 (The Gordon/ U Test):** If 3-D task allocation is the 3-D-local computation of the 4-D-global gradient ∇U , then the system is coupled non-locally via the 4-D K_s kernel . We predict that two ant colonies (Σ_A, Σ_B) that are physically and chemically isolated in 3-D (preventing any 3-D-local signal) remain coupled in 4-D. A drastic state change in Σ_B (e.g., removal of the queen, a perturbation to the 'germ-plasm' projection) will induce a corresponding, non-local compensatory shift in task allocation in Σ_A . Such a result would violate 3-D-local causality and provide direct evidence of the 4-D counterspace coupling.
- **Prediction 2 (The Cavagna/ K_s Test):** If "second sound" (orientation waves) is the 3-D-shadow manifestation of the 4-D K_s kernel , its properties are governed by the 4-D geometry, not just the 3-D state. Cavagna et al. found the propagation speed c_2 depends on the 3-D polarization Φ . Our framework predicts c_2 *also* depends on the non-local coupling K_s . We predict that flocks in 3-D environments that alter the 4-D embedding geometry (e.g., strong gravitational lensing, or other phenomena related to the physical acceleration scale a_*) will exhibit anomalous propagation speeds (c_2) not explainable by their 3-D polarization (Φ) alone.

In conclusion, the TCGS-SEQUENTION framework provides a complete, geometrically-grounded, and falsifiable explanation for the foundational puzzles of collective animal behavior. The "superorganism" is not an analogy but the 3-D projection of a 4-D geometric entity defined by Axiom A2. The "combination problem" [1] is resolved by this projection-first ontology. The "distributed processes" of 3-D collective intelligence are the computational algorithms by which 3-D collectives navigate the 4-D-defined informational landscape (U) . Finally, the empirical puzzles of 3-D non-locality, such as scale-free correlations , are the definitive "smoking gun" signatures of the 4-D non-local, retrocausal K_s

kernel , providing a direct, testable link between 3-D animal behavior and the fundamental, timeless geometry of the counterspace.

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