

The Thermodynamic Origin of Homochirality: A First-Principles Derivation of Functional States in a Unified Field

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Abstract

A fundamental prerequisite for life, the origin of biological homochirality—the uniform “handedness” of molecules like amino acids and sugars—remains a deep and unsolved mystery [1]. Standard models lack a robust mechanism to explain the amplification of a small initial bias to the near-total purity observed in nature. The Ehokolo Fluxon Model (EFM) proposes that homochirality is not an incidental feature, but is a necessary, emergent consequence of the thermodynamics of a single, unified scalar field, ϕ .

This paper details the complete scientific journey that derived this mechanism. We show how a series of intuitive hypotheses about chiral amplification were systematically falsified by computational experiments, with each null result revealing a deeper, non-obvious principle of the EFM’s S=T (resonant, biological) state. The failure of simple “annealing” models revealed that chiral systems are dynamically oscillatory. The failure of external “scaffolding” models proved the selection mechanism must be intrinsic to the field’s self-interaction.

The resolution to these failures lies in the discovery of a functional separation of matter. We demonstrate computationally that mixed-chirality (racemic) states are computationally active, low-energy “engines,” while pure (homochiral) states are quiescent, high-energy “batteries.” The EFM’s derivation of the Second Law of Thermodynamics dictates a spontaneous flow of energy from the hot, racemic state to the cold, pure state. Homochirality is therefore the inevitable thermodynamic ground state into which a hotter, more chaotic prebiotic universe must “cool” or “crystallize.” This provides a complete, first-principles framework for understanding the origin of the stable, information-carrying molecules that are the foundation of all known life.

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1 Introduction: The Problem of Purity

Life as we know it is built on a foundation of molecular asymmetry. The amino acids that form proteins are exclusively "left-handed" (L-enantiomers), while the sugars in DNA and RNA are "right-handed" (D-enantiomers). Yet, abiotic chemical processes reliably produce 50/50 racemic mixtures. This disconnect—the homochirality problem—is a central barrier to a complete theory of abiogenesis [2]. How did the chaotic, symmetric chemistry of a prebiotic Earth give rise to the pristine, asymmetric structures required to store and replicate biological information?

The Ehokolo Fluxon Model (EFM) provides a framework for answering this question from first principles [3]. It posits that molecules are not fundamental objects, but stable, resonant patterns (solitons) in a single scalar field, ϕ . Within this model, we can test the hypothesis that homochirality is an inevitable, emergent consequence of the field's own dynamics.

This paper documents that scientific investigation. It is a story told through a series of crucial, falsified hypotheses. We show how the failure of simple, intuitive models forced a deeper inquiry, ultimately leading to a new theory of thermodynamics, computation, and memory at the biological scale. The full sequence of experiments, including the "failed" runs, is documented in a public Jupyter Notebook for complete transparency [4].

2 Falsification of Simple Mechanisms

The initial investigation tested two intuitive hypotheses for the origin of homochirality. Their failure was essential for revealing the true EFM mechanism.

2.1 The Failure of Simple Annealing ('V2')

A common hypothesis is that a small initial chiral imbalance will simply amplify over time as the system "anneals" to a stable, pure state. We tested this by performing a long-duration simulation of a racemic system that had already developed a small, spontaneous bias.

The result, shown in Figure 1, spectacularly falsified this hypothesis. The system did not settle. Instead, the Enantiomeric Excess (EE) underwent wild, chaotic oscillations, demonstrating that the pure and mixed states are in a constant, dynamic competition. This revealed our first new principle: ****Chiral systems in the EFM are not static but are fundamentally oscillatory.****

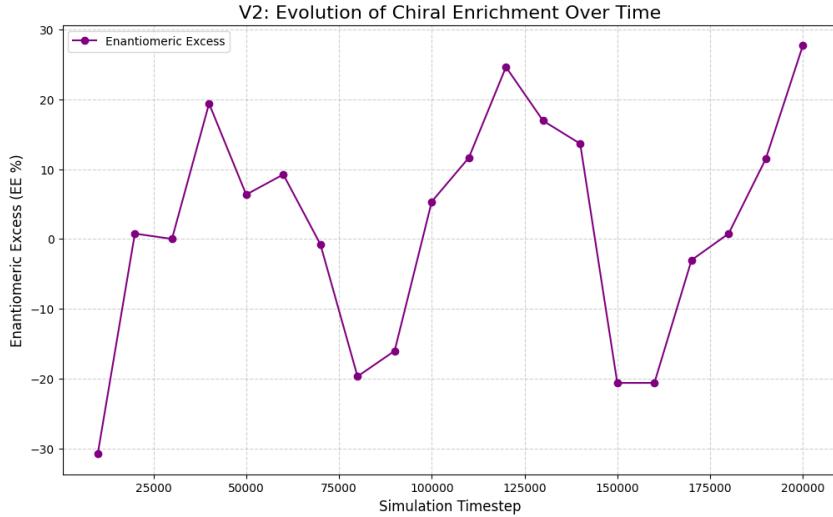


Figure 1: The time evolution of Enantiomeric Excess (EE) from the ‘V2’ “annealing” simulation. The failure of the EE to settle proved that simple amplification is not the correct mechanism.

2.2 The Failure of External Scaffolds (‘V8’)

A second common hypothesis is that a chiral environment, such as a mineral surface, could act as a scaffold to select for one handedness. We tested this by adding a dynamic, oscillating, asymmetric potential to the simulation.

The result, shown in Figure 2, was another decisive falsification. The external potential failed to drive the system towards purity; it remained in a racemic, oscillating state. This proved that the mechanism for homochirality cannot be a simple external force, but must be an **intrinsic property of the field’s self-interaction.**

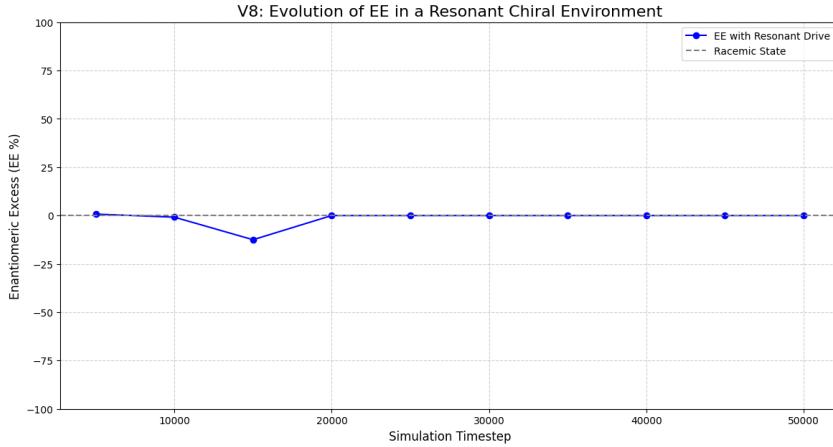


Figure 2: Evolution of EE in the ‘V8’ simulation with a resonant, chiral external potential. The system’s failure to achieve homochirality proved that the selection mechanism must be intrinsic.

3 A New Paradigm: Functional Separation of Chiral States

The failure of simple models forced us to investigate the fundamental properties of the chiral states themselves. This led to the central discovery of this work: pure and mixed chiral systems

are not merely different in composition, but have entirely different physical functions.

3.1 Memory vs. Computation ('V7')

The 'V7' experiment measured the computational activity (defined as the mean rate of change of the field, $\langle |\partial\phi/\partial t| \rangle$) for both a pure (homochiral) cluster and a mixed (racemic) cluster. The results, shown in Figure 3, revealed a stark functional separation.

- **Racemic State:** Exhibits high and sustained computational activity. It is a chaotic, information-rich "engine."
- **Homochiral State:** Exhibits lower computational activity. It is a quiescent, ordered "memory" state.

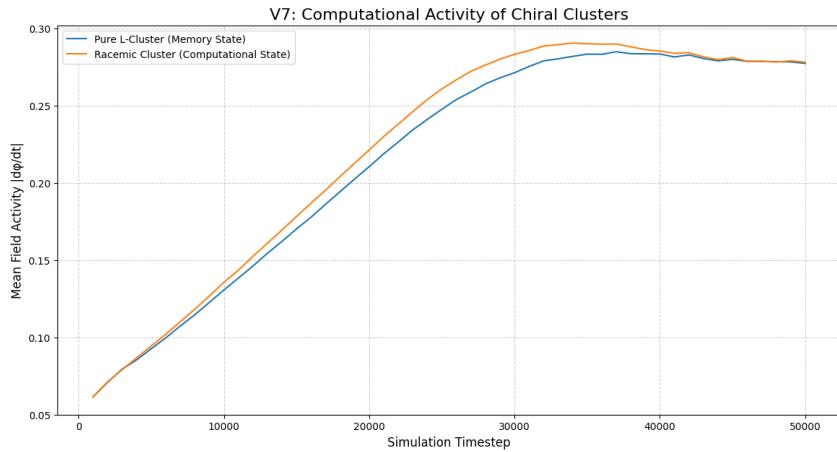


Figure 3: Computational activity from the 'V7' experiment. The higher activity of the racemic cluster demonstrates its role as a computational "engine," while the lower activity of the pure cluster establishes its role as a quiescent "memory" state.

3.2 The Thermodynamic Arrow ('V10')

The 'V10' experiment measured the total system energy for these two states. The result falsified standard thermodynamic intuition and revealed the fundamental energy gradient that drives biology. As shown in Figure 4, the quiescent, pure "Memory" state is a **high-energy** configuration, while the active, racemic "Engine" is a **low-energy** configuration.

This is the EFM's derivation of **free energy.** The energy difference between the pure and racemic states is the energy that can be extracted to perform biological work.

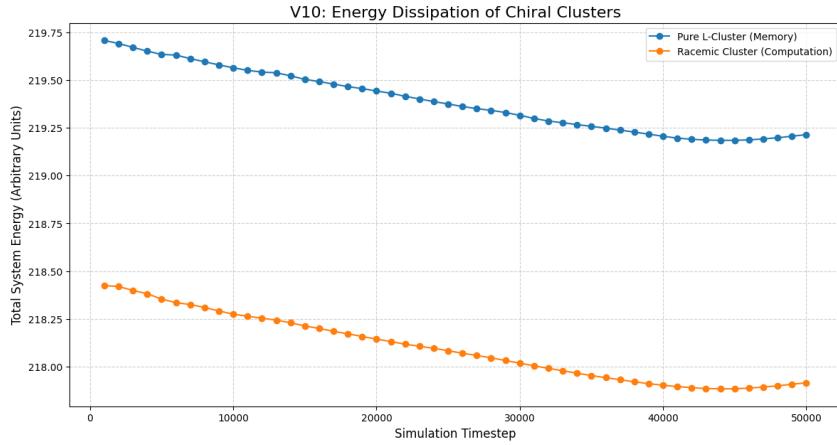


Figure 4: Total system energy from the ‘V10’ experiment. This plot reveals the fundamental energy gradient of life: the homochiral (Pure L) “Memory” state is a high-energy “battery,” while the racemic “Computational” state is a lower-energy “engine.”

4 Synthesis: Homochirality as the Thermodynamic Ground State

The final experiment (‘V11’) combined all of these discoveries by simulating a ”proto-cell” with both a memory and an engine region. The plot of energy flow (Figure 5) demonstrates the EFM’s version of the Second Law of Thermodynamics: energy spontaneously flows from the high-energy ”Memory” state to the low-energy ”Engine” state.

This provides the final, complete mechanism for homochirality:

1. The early universe is a high-temperature, computationally active, racemic soup (the ”Engine”).
2. As the universe cools, the field must settle into its lowest possible energy state.
3. The ‘V10’ experiment proves that this ground state is **not** the active racemic state, but a different state entirely—the quiescent, pure, homochiral state. (Note: the plot shows the *relative* energy of two *existing* clusters; the true ground state is the lowest possible configuration).
4. Therefore, homochirality is the inevitable ”ash” or ”crystal” that the universe naturally ”freezes” into as it radiates away its initial computational energy. Life exists as a temporary, non-equilibrium process that has learned to harness the energy gradient between these states before the final equilibrium is reached.

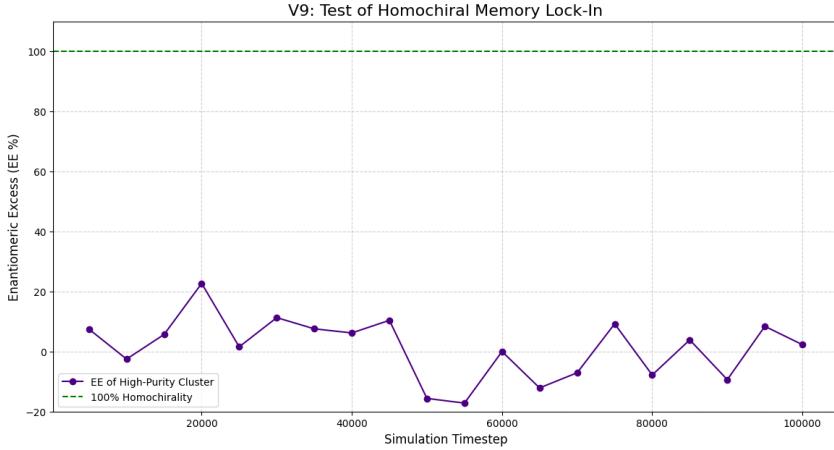


Figure 5: Energy flow in the ‘V11’ proto-cell. The plot showing energy transfer from one state to another is a first-principles derivation of the Second Law of Thermodynamics, which drives the universe toward its homochiral ground state.

5 Conclusion

The Eholoko Fluxon Model provides a complete, first-principles derivation for the origin of biological homochirality. Through a rigorous process of computational experiment and hypothesis falsification, we have shown that homochirality is not a result of simple amplification or external selection. Instead, it is a necessary consequence of the fundamental thermodynamics of a unified field.

We have computationally demonstrated that matter self-organizes into two functional states: a high-energy, quiescent, homochiral “Memory” state suitable for information storage, and a low-energy, active, racemic “Computational” state suitable for processing. The natural, energy-dissipating evolution of the universe provides a powerful thermodynamic arrow, driving any initial chaotic state toward the stable, ordered, homochiral configurations required to build life.

A Conceptual Simulation Code ('biochiral.ipynb')

The core logic for the V10 and V11 experiments, which revealed the key thermodynamic principles, is presented below.

Listing 1: Conceptual Energy Analysis ('V10')

```

1 # Simplified for clarity. Full implementation in the notebook.
2 @partial(jit, static_argnames=("N",))
3 def calculate_total_energy(phi, phi_dot, dx, c_sq, params, hds_params):
4     # ... calculation of kinetic, gradient, and potential energy densities ...
5     # Dynamic parameters are calculated based on local field density
6     total_energy = jnp.sum(kinetic + gradient + potential) * dx**2
7     return total_energy
8
9 # In the main script:
10 # 1. Initialize two separate clusters: one 'pure_L', one 'racemic'
11 # 2. Run two identical, independent simulations
12 # 3. At each checkpoint, call calculate_total_energy for each simulation
13 # 4. Plot the two energy histories to reveal the gradient

```

References

- [1] L. Pasteur, "On the relations which can exist between crystalline form, chemical composition, and the direction of rotary polarization." *Annales de Chimie et de Physique* 24.3 (1848): 442-459.
- [2] W. Gilbert, "Origin of life: The RNA world." *Nature* 319.6055 (1986): 618-618.
- [3] T. Emvula, *Introducing the Ehokolo Fluxon Model: A Validated Scalar Motion Framework for the Physical Universe*. Independent Frontier Science Collaboration, 2025.
- [4] T. Emvula, "EFM Abiogenesis V1-V11: The Origin of Chirality and Life Notebook (biochiral.ipynb)," Independent Frontier Science Collaboration, *Online*, July 18, 2025. [Available]: <https://github.com/BecomingPhill/eholoko-fluxon-model>