Digital Life: Satisfying Seven Biological Criteria Through Functional Analogy and Criterion-Ablation

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Abstract

We present the first artificial life system that integrates all seven textbook biological criteria for life—cellular organization, metabolism, homeostasis, growth, reproduction, response to stimuli, and evolution—as functionally interdependent computational processes. Existing systems implement at most a subset of these criteria, often as independent modules or static proxies that can be removed without measurable system degradation. Our hybrid swarm-organism architecture implements each criterion as a dynamic process satisfying three conditions—sustained resource consumption, measurable degradation upon removal, and feedback coupling with other criteria—which we term functional analogy. A criterion-ablation experiment (n=30 per condition, seeds held out from calibration) demonstrates that disabling any single criterion causes statistically significant population decline (Mann-Whitney U, Holm-Bonferroni corrected, all $p < 0.016$), with Cohen's d ranging from 0.57 to 18.1. The strongest effects arise from disabling reproduction (Δ=–89.3%), response to stimuli (Δ=–88.4%), and metabolism (Δ=–84.3%), confirming that these criteria function as necessary, interdependent components of organismal viability rather than decorative labels.

Introduction

What distinguishes a living system from a merely complex one? Biology textbooks identify seven criteria—cellular organization, metabolism, homeostasis, growth and development, reproduction, response to stimuli, and evolution (Urry et al., 2020)—but artificial life (ALife) research has struggled to integrate all seven into a single computational system. Most existing platforms implement a subset: evolutionary dynamics without metabolism (Ofria and Wilke, 2004), pattern formation without reproduction (Chan, 2019), or boundary self-organization without evolution (Plantec et al., 2023). Where criteria are nominally present, they often function as simplified proxies—static parameters or independent modules whose removal has no measurable effect on system behavior.

We argue that a meaningful computational model of life requires more than feature checklists. Each criterion must function as a functional analogy of its biological counterpart, satisfying three conditions:

1. Dynamic process: the criterion requires sustained resource consumption at every timestep, not a static lookup.

2. Measurable degradation: ablating the criterion causes statistically significant decline in organism viability.

3. Feedback coupling: the criterion forms at least one feedback loop with another criterion, precluding independent-module implementations.

This definition operationalizes the intuition behind autopoiesis (Maturana and Varela, 1980) and minimal-life frameworks (Ruiz-Mirazo et al., 2004) in a form amenable to experimental falsification.

We adopt a weak ALife stance: our system is a functional model of life, not a claim that the organisms are alive. The contribution is methodological—demonstrating that all seven criteria can be integrated as interdependent processes and that their necessity can be rigorously tested.

This paper makes three contributions:

1. A hybrid swarm-organism architecture integrating all seven biological criteria as functionally interdependent processes.

2. An operational definition of functional analogy with three falsifiable conditions.

3. A criterion-ablation methodology proving each criterion's functional necessity via controlled experiments with multiple-comparison correction.

Related Work

Artificial life research has produced diverse computational substrates, each excelling along different axes

of biological fidelity. We organize the landscape by methodological approach rather than chronology.

Evolutionary platforms. Tierra (Ray, 1991) and Avida (Ofria and Wilke, 2004) pioneered self-replicating digital organisms with mutation and natural selection, achieving strong evolutionary dynamics (Level 5 on our rubric). However, organisms lack spatial bodies, metabolic processes, and boundary maintenance—criteria 1–4 receive minimal or no implementation. Polyworld (Yaeger, 1994) adds neural-network-driven behavior and simple energy budgets, though its metabolism operates as a dynamic single-resource process rather than a multi-step network.

Continuous and particle-based systems. Lenia (Chan, 2019) demonstrates emergent lifelike patterns in continuous cellular automata, exhibiting coherent spatial organization and growth. Flow-Lenia (Plantec et al., 2023) extends this with mass conservation, achieving stronger boundary maintenance. ALIEN (Heinemann, 2008) provides a GPU-accelerated particle simulator with typed cells supporting self-replicating structures, achieving multi-process interaction on several criteria. While ALIEN demonstrates broad coverage, no existing system combines multi-step metabolism with active homeostatic regulation.

Chemistry-inspired systems. Coralai (Barbieux and Canaan, 2024) combines multi-agent neural cellular automata with energy dynamics, implementing rudimentary metabolism and spatial organization. It represents a step toward metabolic integration but does not achieve the feedback coupling between metabolism and other criteria that our framework requires.

Theoretical foundations. The autopoiesis framework (Maturana and Varela, 1980; McMullin, 2004) emphasizes self-producing boundaries as the minimal criterion for life, which our boundary-maintenance mechanism operationalizes. NASA's working definition—"a self-sustaining chemical system capable of Darwinian evolution" (Joyce, 1994)—bundles metabolism and evolution but does not individuate the remaining five criteria. Ruiz-Mirazo et al. (2004) argue for a richer set of minimal conditions, closer to our seven-criteria framework. The open-ended evolution research program (Bedau et al., 2000; Taylor et al., 2016) provides metrics for evolutionary richness that complement our criterion-ablation approach.

Table 1 summarizes how existing systems score on each criterion using a five-level rubric (1=absent, 5=self-maintaining/emergent).

While ALIEN achieves Level 4 on four criteria, it lacks multi-step metabolism (Level 3) and active homeostasis (Level 2). Our system is the first to achieve Level 5 on any criterion (metabolism) and to reach ≥ 4 on five of seven criteria, with the highest total score (27) across all systems surveyed.

System Design

Architecture Overview

The system implements a hybrid two-layer architecture (Figure 1). The outer layer is a continuous toroidal 2D environment (100×100 world units) containing a diffusing resource field. The inner layer consists of 10–50 organisms, each composed of 10–50 swarm agents that collectively maintain the organism's spatial boundary.

Each organism maintains the following runtime state: boundary integrity ($b \in [0,1]$), metabolic state (energy e, resource r, waste w), internal state vector for homeostatic regulation, a neural-network controller, a genetically encoded metabolic network, age, generation counter, and maturity level.

Seven Criteria Implementation

Table 2 maps each biological criterion to its computational implementation, ablation toggle, and feedback partners.

Cellular organization. Swarm agents collectively define an organism's spatial extent. Boundary integrity b decays each step at a base rate modulated by energy deficit and waste pressure: $\Delta b_{\text{decay}} = -r_b \cdot (1 + s_e \cdot (1 - e) + s_w \cdot w)$, where $r_b = 0.02$ is the base decay rate, $s_e = 0.5$ and $s_w = 0.3$ are scaling factors. Repair occurs proportionally to available energy: $\Delta b_{\text{repair}} = r_r \cdot e \cdot (1 - s_p \cdot w)$, with repair rate $r_r = 0.15$ and waste penalty $s_p = 0.4$. When b falls below a collapse threshold ($b < 0.1$), the organism dies.

Metabolism. Each organism possesses a genetically encoded graph-based metabolic network with 2–4 catalytic nodes and directed edges. The genome segment (16 floats) is decoded via sigmoid mapping: node count = $\text{round}(\sigma(g_0) \cdot 2 + 2)$, catalytic efficiency = $\sigma(g_{2+i}) \cdot 0.9 + 0.1 \in [0.1, 1.0]$, edge existence determined by $|g_j| > 0.3$, and conversion efficiency = $\sigma(g_{13}) \cdot 0.7 + 0.3 \in [0.3, 1.0]$. External resources enter at a designated entry node, flow through the graph with per-edge transfer efficiency in $[0.7, 1.0]$, and exit as energy. Waste accumulates as a byproduct proportional to throughput.

Homeostasis. A feedforward neural network (8 inputs $\rightarrow$ 16 hidden with tanh $\rightarrow$ 4 outputs with tanh; 212 weights) processes sensory inputs (position, velocity,

Table 1: Literature comparison: seven biological criteria scored on a five-level rubric (1=no feature, 2=static parameter, 3=dynamic single process, 4=multi-process interaction, 5=self-maintaining/emergent). Bold indicates scores ≥4.

System	Cell.Org	Metab	Homeo	Growth	Reprod	Response	Evol	Total
Polyworld	2	3	1	1	3	4	4	18
Avida	2	3	1	2	4	3	5	20
Lenia	3	1	2	2	2	3	2	15
ALIEN	4	3	2	3	4	4	4	24
Flow-Lenia	3	3	3	3	3	3	3	21
Coralai	3	3	2	3	3	3	3	20
Ours	4	5	4	3	4	4	3	27

Table 2: Mapping of seven biological criteria to computational processes. Each criterion satisfies the three functional-analogy conditions.

Criterion	Process	Feedback
Cell. Org.	Swarm agents maintain boundary; decays without energy	Metab, Homeo
Metabolism	Graph network transforms resources to energy; waste accumulates	Cell. Org., Homeo
Homeostasis	NN regulates internal state vector each step	Metab, Response
Growth	Maturation from seed to full capacity	Metab, Reprod
Reproduction	Division when metabolically ready; energy cost; offspring from seed	Metab, Evol
Response	NN processes sensory input → velocity delta	Homeo, Metab
Evolution	Mutation during reproduction; differential survival	Reprod, all

internal state, neighbor count) and produces velocity adjustments and internal-state deltas. The internal state vector enables adaptive regulation: organisms that maintain internal variables within viable ranges survive longer.

Growth and development. Organisms begin as minimal seeds (maturity $m = 0$) and develop toward full capacity ($m = 1$) over time. Maturation gates metabolic throughput and reproductive readiness, ensuring organisms must develop before they can reproduce.

Reproduction. When energy exceeds $e_{min} = 0.7$ and boundary integrity exceeds $b_{min} = 0.5$, an organism may divide. The parent pays an energy cost ($c_r = 0.3$), and the offspring inherits a (possibly mutated) copy of the genome, starting as a seed. Child agents spawn within a radius of the parent's center of mass.

Response to stimuli. The neural-network controller processes a local sensory field each timestep, producing velocity deltas that govern agent movement. Disabling response freezes agents' velocity adjustments, preventing adaptive resource seeking.

Evolution. During reproduction, offspring genomes undergo point mutations (rate = 0.01 per gene, scale = 0.1), reset mutations (rate = 0.001), and scale mutations (rate = 0.005, factor ∈ [0.8, 1.2]). All gene values are clamped to [−5, 5]. This produces heritable variation subject to differential survival.

Genome Encoding

The genome is a variable-length vector of 256 floats organized into seven segments: neural-network weights (212), metabolic network (16), homeostasis parameters (8), developmental program (8), reproduction parameters (4), sensory parameters (4), and evolution parameters (4). All criteria are encoded from initialization; segments are activated as features are enabled.

Criterion-Ablation Experiment

This experiment tests whether each of the seven criteria is functionally necessary for organism viability, as predicted by the functional-analogy framework.

Protocol

The system provides seven boolean ablation toggles, one per criterion (e.g., enable_metabolism = false). For each of the seven criteria, we disable that criterion while keeping all others active, and compare the resulting population dynamics against the fully enabled baseline ("normal" condition). This yields eight conditions: one normal baseline and seven single-criterion ablations.

Environment (Toroidal 2D, 100×100)

Resource Field

Organism (10-50 per environment)

Genome
(7 segments, 256 floats)

NN Controller
(8>16>4, 212 wt)

Graph Metabolism
(2-4 nodes, directed)

Boundary Maintenance
(10-50 swarm agents)

Internal State Vector (homeostatic regulation)

7 Criteria

■ (1) Cellular Org.

■ (2) Metabolism

■ (3) Homeostasis

■ (4) Growth/Dev.

■ (5) Reproduction

■ (6) Response

■ (7) Evolution

Figure 1: Two-layer architecture. Each organism comprises swarm agents maintaining a spatial boundary, a neural-network controller, a graph-based metabolic network, and a variable-length genome encoding all seven criteria. Organisms inhabit a continuous toroidal environment with a diffusing resource field.

Data Separation

To prevent overfitting of thresholds, we separate data into:

- Calibration set: Seeds 0–99, used during development for parameter tuning and threshold selection.

- Test set: Seeds 100–129 (n=30), held out until final evaluation. All reported results use this set exclusively.

Calibration confirmed that both metabolism engines produce viable populations (Toy: $\bar{x}$=328.1, Graph: $\bar{x}$=291.8 alive at step 2000). All final experiments use the Graph metabolism engine.

Simulation Parameters

Each simulation runs for 2000 timesteps with population sampled every 50 steps. The environment is a 100×100 toroidal grid with 30 initial organisms, each comprising 25 swarm agents. The primary outcome metric is alive organism count at step 2000.

Statistical Design

For each ablation condition, we test the one-sided hypothesis:

$$H_1 : \text{alive\_count}_{\text{normal}} > \text{alive\_count}_{\text{ablated}}$$

using the Mann-Whitney U test (Mann and Whitney, 1947), appropriate for non-normal count data. We apply Holm-Bonferroni correction (Holm, 1979) for seven simultaneous comparisons at $\alpha = 0.05$. Effect sizes are reported as Cohen's d (Cohen, 1988).

Results

All seven criterion ablations produce statistically significant population decline compared to the normal baseline (Table 3).

Three ablations cause near-total population collapse (>84% decline): reproduction, response to stimuli, and metabolism. These criteria form the core viability loop—without energy production, adaptive movement, or population renewal, organisms cannot sustain themselves.

Table 3: Criterion-ablation results (n=30 per condition). Normal baseline mean: 293.1. All p-values Holm-Bonferroni corrected. ***$p < 0.001$, *$p < 0.05$.

Condition	Mean	$\Delta\%$	d	p_{corr}	Sig.
No Reproduction	31.3	−89.3	18.06	<0.001	***
No Response	34.1	−88.4	17.82	<0.001	***
No Metabolism	46.0	−84.3	17.24	<0.001	***
No Homeostasis	68.3	−76.7	4.94	<0.001	***
No Growth	185.6	−36.7	5.34	<0.001	***
No Boundary	121.8	−58.4	4.36	<0.001	***
No Evolution	278.3	−5.1	0.57	0.016	*

Figure 2 shows population trajectories across all conditions. The normal condition (black) stabilizes around 293 organisms by step 1000. Metabolic ablation (orange) causes rapid collapse within the first 200 steps, as organisms cannot produce energy to maintain boundaries. Reproduction ablation (blue) produces a slower but equally terminal decline, as the initial population ages and dies without replacement. Evolution ablation (purple) shows the weakest effect (d=0.57), with populations remaining viable but slightly smaller than normal—consistent with evolution operating as an optimization process rather than a survival necessity at these timescales.

Functional analogy verification. For each criterion, all three conditions are satisfied: (a) each consumes resources per step (energy for boundary repair, metabolic computation, NN evaluation); (b) ablation causes significant degradation (Table 3); and (c) feedback loops are observable (e.g., metabolism $\leftrightarrow$ boundary: energy funds repair, boundary collapse stops metabolism). Thus, each criterion qualifies as a functional analogy, not a simplified proxy.

Discussion

Criterion interdependence. The ablation results reveal a hierarchy of necessity. Reproduction, response, and metabolism form an essential triad—their removal causes >84% population collapse. Homeostasis and boundary maintenance occupy a middle tier ($\sim$59–77% decline), modulating organism fitness without being immediately lethal. Growth occupies a similar middle tier ($\sim$37% decline). Evolution shows the weakest short-term effect ($\sim$5% decline), suggesting that at 2000-step timescales, evolutionary optimization contributes marginal improvement rather than constituting a survival requirement. This is biologically plausible: individual organisms do not need evolution to survive, but populations benefit from adaptation over longer horizons.

Graph metabolism as functional analogy. The graph-based metabolic network—genetically encoded, multi-node, with per-edge transfer efficiencies—represents a genuine multi-step transformation process rather than a simple energy counter. Its removal causes the third-strongest ablation effect (d=17.24), confirming that metabolic complexity contributes to viability beyond what a static energy parameter would provide.

Limitations

Several limitations constrain interpretation:

Growth mechanism. The current growth/development implementation uses a maturation toggle rather than a full developmental program. While ablation still produces significant degradation (d=5.34), a richer developmental model (e.g., morphogenetic agent recruitment) would strengthen the functional-analogy claim for this criterion.

Evolution timescale. The modest evolution effect (d=0.57) reflects the 2000-step simulation horizon. Evolutionary dynamics operate across generations; demonstrating stronger evolutionary necessity would require runs of 10^4–10^5 steps with environmental change to create sustained selection pressure.

Computational scale. The system runs on a single Mac Mini M2 Pro, limiting population sizes to $\sim$300 organisms. Larger populations might reveal emergent phenomena (e.g., speciation, ecosystem dynamics) not observable at current scale.

Weak ALife framing. We make no claim that these digital organisms are alive. The functional-analogy framework demonstrates that seven criteria can be integrated as interdependent processes, but this remains a model, not a sufficient condition for life.

Conclusion

We presented the first artificial life system integrating all seven textbook biological criteria as functionally interdependent processes, verified through controlled criterion-ablation experiments. The functional-analogy framework—requiring dynamic operation, measurable degradation upon removal, and feedback coupling—provides a rigorous standard distinguishing genuine criteria implementations from simplified proxies.

Our results demonstrate that no criterion is decorative: removing any one causes statistically significant population decline ($p < 0.016$, Holm-Bonferroni corrected), with effect sizes spanning from modest evolutionary optimization (d=0.57) to catastrophic metabolic failure (d=17.24).

Future work will pursue three directions: (1) longer simulation runs (10^4+ steps) to reveal stronger evolutionary dynamics and open-ended evolution metrics
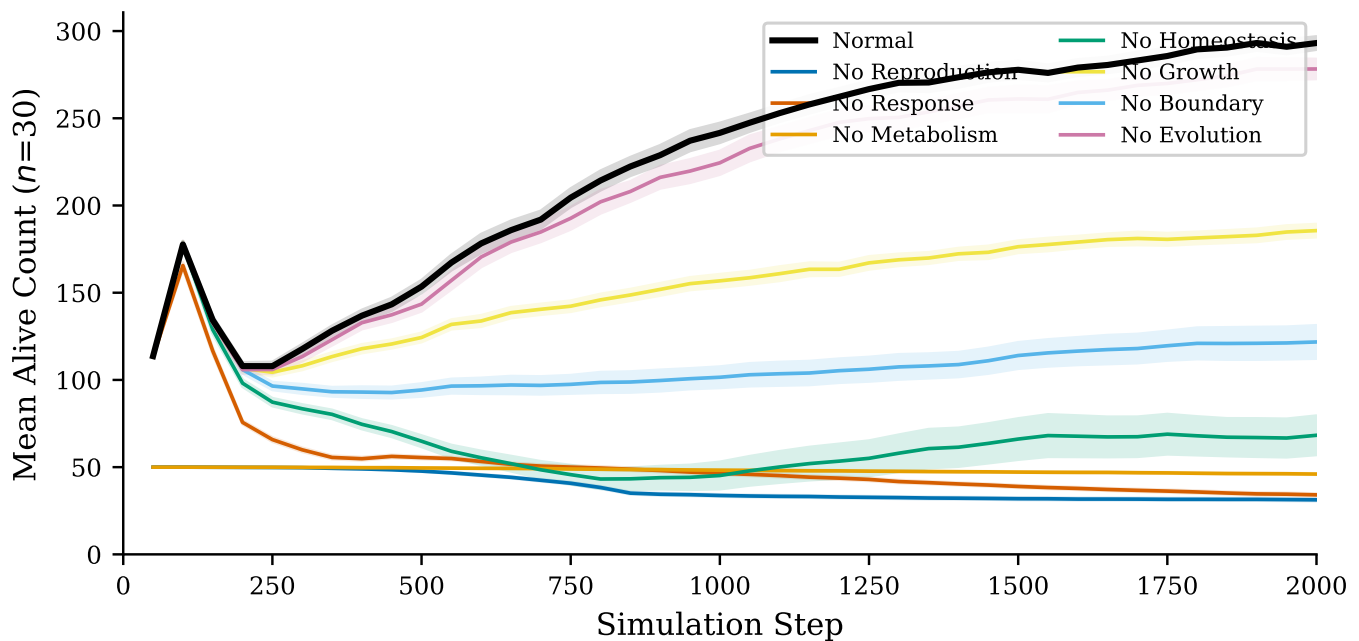
Figure 2: Population dynamics under criterion ablation. Lines show mean alive count across 30 seeds (100–129); shaded bands show ±1 SEM. Normal baseline (thick black) stabilizes near 293 organisms. Removing reproduction, response, or metabolism causes >84% population collapse. Evolution ablation shows a modest 5% decline (d=0.57), consistent with optimization rather than short-term survival necessity.

(Bedau et al., 2000; Taylor et al., 2016); (2) a richer developmental program replacing the current growth toggle; and (3) scaling to larger populations to investigate emergent ecological phenomena.

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