

Integrated Human Dynamics

A concept note and research agenda for multi-layer modeling of repair, regeneration, and whole-body integration

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Positioning note: this document does not claim a completed unified theory. It proposes a research framing, a layered vocabulary, and near-term experimental entry points for a longer-term program.

Abstract

Human biology is better approached as a distributed, adaptive, multi-layer control system than as a static assembly of replaceable parts. For that reason, local repair, enhancement, or redesign can create downstream incompatibilities across neural, circulatory, immune, endocrine, and mechanical layers even when a local intervention appears successful. This concept note introduces Integrated Human Dynamics as a provisional upper-level framework for describing, measuring, and eventually controlling those cross-layer interactions. The aim is not to present a completed grand theory, but to define why such a theory is needed, how it may be organized, and which experimental entry points are tractable today. The central proposition is that repair and redesign should be modeled as re-synchronization across layers rather than isolated modification of a single module. Within this framing, neuroregeneration becomes a near-term test bed because exploration, selection, reinforcement, pruning, and stabilization can be observed and manipulated. The note places the author's prior stigmergic neuroregeneration hypothesis as a local, experimentally accessible sub-framework inside a broader agenda that also includes developmental biology, mechanobiology, immune compatibility, circulatory integration, and digital-twin-style safety modeling.

Keywords: systems biology; regenerative medicine; neuroregeneration; mechanobiology; immune compatibility; digital twin; whole-body integration; concept note

Why this note

Human repair is often discussed at the level of tissues, molecules, or individual organs. But if the body is better understood as a distributed adaptive system, then local repair, enhancement, or redesign can propagate across neural, circulatory, immune, endocrine, and mechanical layers.

A framework is therefore needed that can describe not only local intervention, but also cross-layer coupling, failure propagation, recovery, and safe control. This note records that framing as a working concept rather than a final doctrine.

Working definition

Integrated Human Dynamics is a working concept for treating genetic, cellular, tissue, organ, circuit, systemic, and environmental interactions as a multi-layer control problem with delays, feedback, and safety constraints.

As descriptive science	Describe the body as a coupled hierarchy rather than a stack of isolated parts.
As predictive science	Estimate where local success will create downstream incompatibility, instability, or incomplete recovery.
As an engineering program	Design repair, reconstruction, and limited augmentation under closed-loop control and explicit safety envelopes.

Central propositions

1. Repair and redesign should be modeled as cross-layer re-synchronization rather than isolated modification of a single module.
2. Local success is insufficient if neural, circulatory, immune, and mechanical coherence is not restored.
3. The first practical entry point is therapeutic reconstruction, not radical morphological redesign.
4. Neuroregeneration offers an early test bed because exploration, reinforcement, pruning, and stabilization can be measured and perturbed.
5. Future augmentation, if ever pursued, must remain downstream of safety modeling, ethical boundaries, and demonstrated whole-body compatibility.

Layered architecture

The framework can be organized as six major layers plus a cross-cutting bidirectional feedback logic.

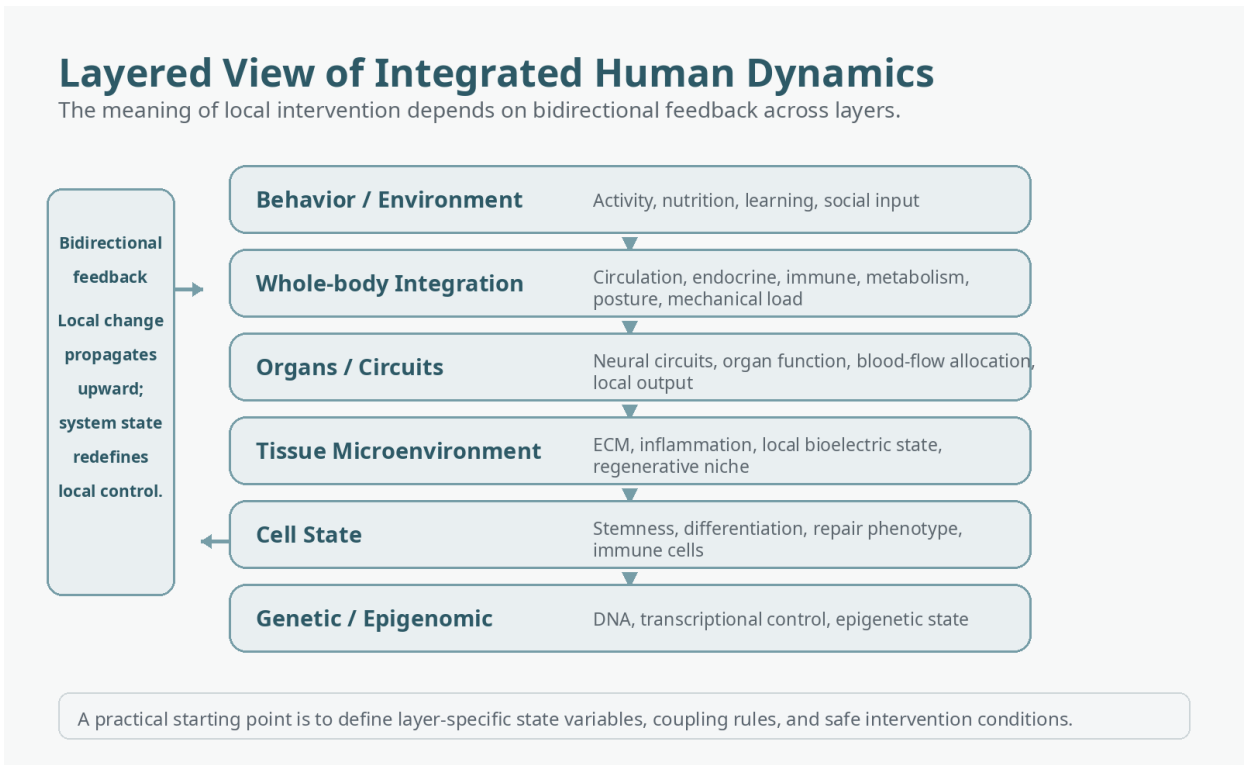


Figure 1. Conceptual layer structure of Integrated Human Dynamics.

5. Layer table

Layer	Example state variables	Primary question
Genetic / Epigenomic	DNA, transcriptional control, epigenetic state	What local changes are stable, reversible, or developmentally constrained?
Cell State	Stemness, differentiation, repair phenotype, immune cells	Which cell states can be shifted, retained, and safely coordinated?
Tissue Microenvironment	ECM, inflammation, local bioelectric state, regenerative niche	Which local conditions permit repair without dysplasia or collapse?
Organs / Circuits	Neural circuits, organ function, blood-flow allocation, local output	How does local repair become usable function?
Whole-body Integration	Circulation, endocrine, immune, metabolism, posture, mechanical load	How is local success coupled to systemic stability?
Behavior / Environment	Activity, nutrition, learning, social input	Which external inputs stabilize or destabilize outcomes?

Connection to the prior neuroregeneration framework

The author's prior manuscript, *Stigmergic Neuroregeneration*, proposed a local grammar of trace writing, distributed exploration, selective reinforcement, pruning, and consolidation for adaptive nerve repair.

In the present framing, that work is not displaced. It is repositioned as a local and experimentally accessible subsystem inside a broader program concerned with organ/circuit integration, whole-body compatibility, and safe intervention.

Local sub-frame: write traces -> explore -> reinforce -> prune -> consolidate. Upper frame: connect local success to organ/circuit function, whole-body stability, and safety control.

Earliest falsifiable program

Minimal hypothesis	Minimal test	What would weaken it
Local regenerative success without organ/circuit integration plateaus or misfires.	Compare tissue-local readouts against functional circuit readouts in nerve-muscle repair models.	If tissue-local success fully predicts recovery without higher-level variables.
Activity-gated or feedback-shaped reinforcement outperforms static guidance in heterogeneous repair settings.	Compare fixed cue scaffolds with adaptive reinforcement in PNS repair models.	If static guidance repeatedly wins across heterogeneous lesions.
Mechanical and immune boundary conditions improve failure prediction beyond purely local regenerative models.	Add immune and mechanical variables to closed-loop predictions of repair outcome.	If extra variables add complexity without predictive gain.

Staged roadmap

Phase	Focus	Expected output
Phase 0	Vocabulary and boundaries	Operational definitions, exclusions, and a first variable map.
Phase 1	Neuroregeneration entry model	A closed-loop local repair model with explicit failure criteria.
Phase 2	Nerve-muscle-vessel-immune coupling	Boundary models that connect local success to organ integration.
Phase 3	Development/regeneration bridge	Links between pattern formation, regenerative niches, and reconstruction.
Phase 4	Digital twin and safety envelope	Prediction of stability, failure points, and allowable intervention windows.
Phase 5	Therapeutic reconstruction first	Only after safety, cautiously test limited structural optimization or augmentation.

What this note does not claim

1. It does not claim a completed unified theory of the human body.
2. It does not claim a validated route to radical body redesign.
3. It does not argue for bypassing ethics, safety, immune compatibility, or long-term monitoring.
4. It does not assume that local regenerative success automatically scales to whole-body redesign.

Public positioning for Zenodo

This document is best released on Zenodo as a concept note, working hypothesis, and research agenda. Versioned updates should refine state variables, readouts, failure criteria, and ethical boundaries rather than pretend final closure.

The public value of v0.1 lies in clarifying scope and research entry points. Its success condition is not immediate completeness, but whether it helps organize more precise future work.

Declarations

Manuscript type	Concept note / research agenda
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Ethics	Not applicable. This note reports no new studies involving human participants or animals.
Data and code	No new datasets or code were generated for this note.
Use of generative AI	Generative AI tools were used for language support and editorial assistance. Final review and responsibility remain with the author.

Next-version priorities

1. Operationalize the vocabulary: coherence, safety envelope, cross-layer synchronization, and redesign.
2. Translate the local neuroregeneration sub-frame into measurable variables, readouts, and failure criteria.
3. Formalize coupling rules across nerve, muscle, vessel, immune, and mechanical boundaries.
4. Separate therapeutic reconstruction, limited augmentation, and radical morphology in both safety language and ethics language.