

OUTCOVE VITALITY · NAPA VALLEY · BREAKOUT SESSION

MitoXcel™ Geropeptides

Not Your Typical Peptide Technology

A closer look at origin and validation — comparing the peptides exploding in longevity medicine with the information-encoded MitoXcel™ platform.

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GEROTHERAPEUTIX · Eos SENOLYTIX · Perseus SENOLYTIX

The Peptide Gold Rush

Interest in “longevity peptides” has exploded — driven by online communities, biohacker culture, and compounding pharmacies, well ahead of rigorous clinical validation.

BPC-157

Gastric-derived healing peptide

TB-500

Synthetic thymosin β 4 fragment

KPV

Alpha-MSH fragment

MOTS-c

Mitochondrial-derived peptide

Semax

Russian nootropic

Epitalon

Pineal peptide bioregulator

JULY 23–24, 2026

FDA Pharmacy Compounding Advisory Committee

reviewed 7 popular peptides for the 503A bulk compounding list.

6 of 7

received a favorable committee recommendation — BPC-157, KPV, TB-500, MOTS-c, Semax, and Epitalon.

What "FDA-Allowed" Really Means

A 503A bulks-list recommendation is not FDA drug approval. It is a permission slip for compounding pharmacies — not a verdict on safety or efficacy.

	503A Bulks-List Recommendation	FDA Drug Approval
Review type	Advisory committee vote	Full premarket safety & efficacy review
Binding?	No — rulemaking still required	Yes — legally required before marketing
Manufacturing	Individual compounding pharmacies	Standardized, FDA-inspected GMP facilities
Human efficacy data	Largely absent or uncontrolled	Adequate & well-controlled trials required

FDA’s own scientific reviewers recommended AGAINST adding all seven peptides, citing gaps in characterization, effectiveness, and safety data. The committee vote overruled its own agency staff.

Where These Peptides Came From

1989

BPC-157

Isolated from screened human gastric juice samples at the University of Zagreb; never advanced through completed human efficacy trials.

1990s

TB-500

A synthetic fragment of thymosin beta-4, studied almost entirely in animal and in vitro models.

2015

MOTS-c

A short peptide encoded in mitochondrial DNA, discovered via genomic analysis; no controlled human trials.

Soviet-era

Semax & Epitalon

Peptide “bioregulators” developed in Russia; Semax is an approved drug in Russia only, not reviewed by the FDA.

None of these peptides originated from a rationally designed platform built around a defined mechanism of aging.

The Validation Gap

Per the FDA's own July 2026 briefing materials, evidence for these peptides is thin — and several carry specific safety signals.

1

No controlled human trials

KPV, TB-500, and MOTS-c: human clinical evidence is essentially absent.

2

Small, uncontrolled, or contradictory

BPC-157, emideltide, and Semax: available human studies don't settle efficacy.

3

Specific safety signals flagged

BPC-157 adverse-event reports; epitalon's telomerase activation with uncertain cancer risk; Semax anticoagulant / amphetamine-potential interactions.

4

Manufacturing & doping concerns

Limited impurity, aggregate, and endotoxin testing; MOTS-c and TB-500 are WADA-prohibited substances.

A DIFFERENT STARTING POINT

Enter MitoXcel™

Not a molecule pulled from folk remedies, gastric extracts, or Soviet-era research — but an information-encoded, AI-designed platform built from a defined design framework.

DEVELOPER

SENOTHERAPEUTIX, Inc. (d/b/a GEROTHERAPEUTIX, Inc.), Houston, TX — founded 2019

DESIGN METHOD

Computationally-derived rules: charge distribution, amphipathicity, structural motifs

NOT A SINGLE DRUG

A platform — any sequence must satisfy the MitoXcel™ blueprint to qualify

One Target, Chosen Deliberately

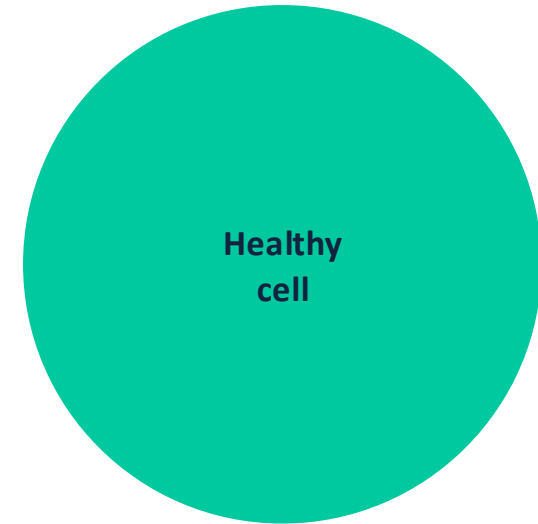
Mitochondrial Membrane Potential ($\Delta\Psi_m$)

Across every studied organism, aging brings a predictable, measurable decline in $\Delta\Psi_m$. This bioenergetic decay precedes impaired ATP production, metabolic instability, and the transition of cells into senescence.

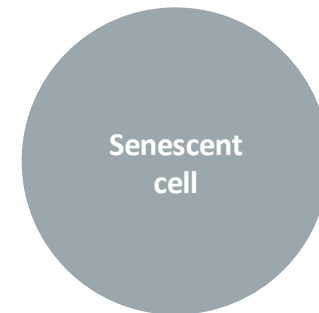
Senescent cells run at markedly lower $\Delta\Psi_m$ than healthy cells — a naturally occurring selectivity filter that MitoXcel™ geropeptides are engineered to recognize, without relying on a specific cell-surface receptor.

Why this matters

A universal, cell-type-agnostic vulnerability — not a single-pathway drug target — is what makes a true gerotherapeutic possible.



High $\Delta\Psi_m$



Low $\Delta\Psi_m$

Two Mechanisms, One Molecule

1

Rapid Bioenergetic Restoration

Elevates $\Delta\Psi_m$ in aging cells, restoring mitochondrial efficiency toward a more youthful phenotype — rapidly, with no receptor engagement or external co-factors required.

2

Selective Apoptotic Clearance

Triggers self-elimination of senescent cells across every major organ, including senescent glial cells in the brain — reducing systemic inflammatory burden.

The downstream effect

Reduced fat mass, increased muscle mass, enhanced tissue function, and improved cognition are not directly targeted — they emerge naturally from restored cellular energetics and lower senescent-cell burden.

Preclinical Validation, By the Numbers

>75 wks

long-term dosing studies with no evidence of safety issues or dose limiting toxicity

p=0.0098

extended lifespan vs. saline by 13.1% to date.

**First
Gerotherapeutic?**

extensive studies in naturally aged mice demonstrate system-wide healthspan improvements

**2 / 3 Bladder Ca
1 / 2 Breast Ca
2 / 3 Colon Ca
4 / 4 Melanoma Ca**

cell lines responded to the MitoXcel™ oncopeptide PTC-2128 in solid-tumor screening

Validated with a systematic preclinical program — dose escalation, long-term toxicology, and disease-model efficacy — advancing toward the FDA's actual drug-approval pathway, not the compounding pathway.

Source

Preclinical data disclosed by Eos SENOLYTIX and Perseus SENOLYTIX (MitoXcel.com, EosSenolytix.com, PerseusSenolytix.com, GEROTHERAPEUTIX.com).

Side-by-Side: Origin & Validation

	Popular Longevity Peptides	MitoXcel™ Gerpeptides
Discovery method	Gastric extracts, animal fragments, genomic screening, Soviet-era research	Computational / AI design against a defined aging mechanism
Mechanism clarity	Often unclear or receptor-based, case by case	Single, universal target: mitochondrial membrane potential ($\Delta\Psi_m$)
Human trial status	Absent, small, uncontrolled, or foreign-market only	Advancing through a structured preclinical-to-IND program
Regulatory pathway	503A compounding (no premarket efficacy review)	FDA drug-approval pathway (IND / clinical development)
Manufacturing	Variable, pharmacy-by-pharmacy compounding	Controlled, standardized development program

Pipeline & Path Forward

Lead candidate: PTC-2105

A homeostatically adaptive geropeptide for sarcopenia and sarcopenic obesity — unique in restoring the body's healthy physiological set point rather than pushing biology in a single direction.

Eos SENOLYTIX

PTC-2105, PTC-2107, PTC-2113, PTC-2118

Healthspan / longevity geropeptides

Perseus SENOLYTIX

PTC-2109, PTC-2110, PTC-2117, PTC-2128

First-in-class MitoXcel™ oncopeptides

Ponce Aurora SENOLYTIX

PTC-2111, PTC-2116, PTC-2119, PTC-2132

YUUTH™ cosmetics, YUURN™ longevity beverages, CHUUS™ gummies

The path we're choosing

- Unlike compounded peptides sold under the 503A bulks-list pathway, which are often research grade, MitoXcel™ geropeptides are advancing through the FDA's IND / clinical development pathway — the same rigorous route required of any approved medicine
- May be manufactured to full GMP, ISO22716 or ISO9001+additional QA quality standards depending on the need
- That means standardized manufacturing, controlled dosing, and clinical trials designed to establish safety and efficacy — not just eligibility for a compounding pharmacy to prepare a batch.



Focused Portfolio



FULL BODY REVITALIZING
SERUM



FULL BODY TREATMENT



CELLULAR VITALITY
CREAM



CELLULAR RENEWAL
CREAM



CELLULAR PERFECTING
SERUM

YUUTH Inner Circle membership program



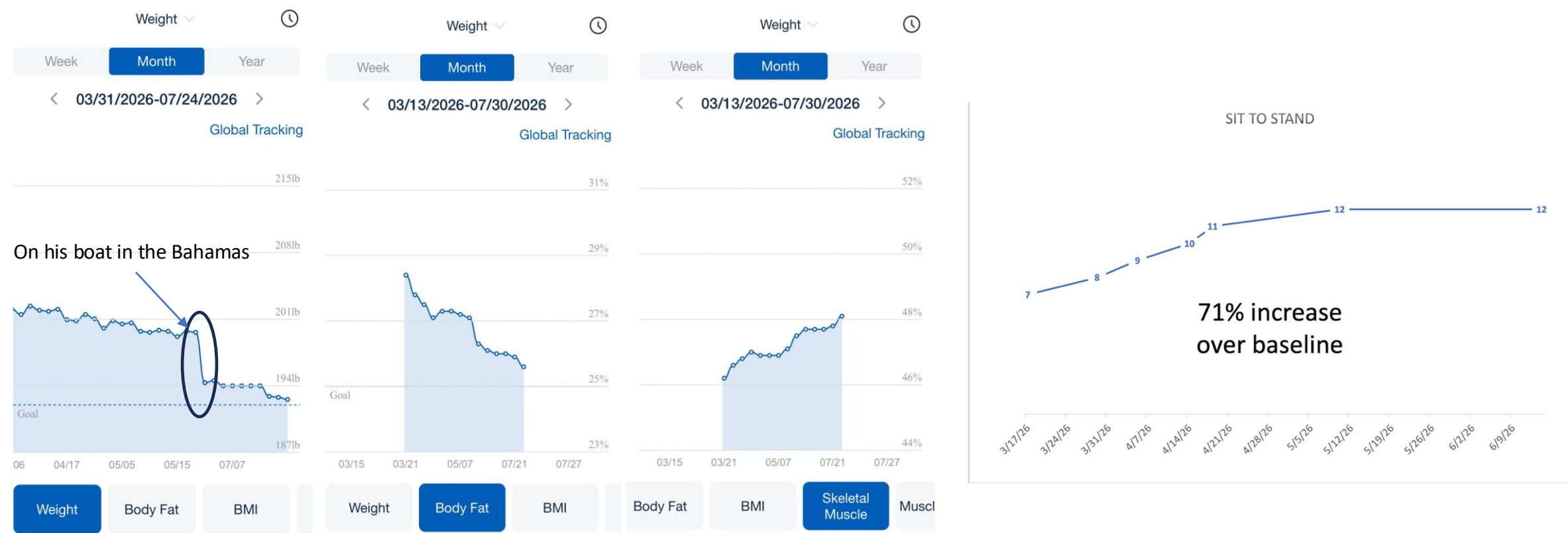
YUUTH Full Body Treatment – Limited Edition

Early Access. Prestige Packaging.



63 year old healthy but out of shape white male

YUUTH™ Full Body Treatment – Pre-Release Sample Applied 3X/Week since 3/2026



KEY TAKEAWAY

Being on a compounding list isn't the same as being validated.

MitoXcel™ geropeptides were designed — not discovered by accident — against a single, universal, mechanistically defined signature of aging, and are being validated through the same rigorous pathway required of any approved medicine.

Not your typical peptide technology.

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