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PRIMEIA · THE FERTILITY & PREGNANCY INSTITUTE · OVALAIRE · SUPERBABY

A LETTER TO THE MAN WHO LOVES HER

Why her thyroid is the most consequential gland in the body that may carry your child.

A scientific brief, prepared for the husband, partner, fiancé, or co-parent.



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EXECUTIVE SUMMARY

In one page

What this letter is. A scientific brief, prepared for the man who loves a woman whose thyroid, hormones, and full neuroendocrine axis are about to be addressed with the deepest intelligence available to her. A translation, not a pressure letter. Designed to help you understand what she may not be able to say without exhaustion, tears, or the careful flatness women use when they have stopped expecting to be believed.

What she is asking. Not for you to fix her. Not for you to become an endocrinologist. To stand beside her while she addresses the gland that often has to be relieved before her ovaries, her eggs, her uterus, her gut, and her nervous system can come back online — inside one twelve-week container, co-led by Dr. Amie Hornaman and me.

What this transformation is. Total Thyroid & Hormone Fix with Thyroid Burnout Recode. A twelve-week, two-layer transformation. The Clinical Layer reads her labs and prescribes when clinically indicated. The Psychogenomic Layer reads her life and rewrites the survival pattern that burned the thyroid out in the first place. Both are necessary. Neither is sufficient alone.

What it is not. Not a diagnosis. Not a treatment. Not a promise. Not generic thyroid medication. Not another bottle of selenium. Not a scattered protocol stitched together across five practitioners.

This is Primemester-level work concentrated specifically on the thyroid and the HPA-axis that governs both her thyroid and her eggs — because the thyroid is so critical for fertility, pregnancy, fetal programming, and offspring outcomes that it earns its own dedicated container.



THE SCIENTIST WRITING TO YOU

Dr. Cleopatra Kamperveen, Ph.D.

Fertility & Pregnancy Scientist · World's First Psychogenomics Expert

30+ years teaching the science of fertility · since 1996

\$3M in NIH and NSF research funding awarded

2,700+ scholarly citations · Google Scholar

47.90 NIH Weighted Relative Citation Ratio

Academic and scientific credentials. Retired Tenured Professor, University of Southern California — first woman of color on the tenure track at the USC Davis School. Tenure across three USC departments. h-index 20. Founder, The Fertility & Pregnancy Institute, Global Fertility Inc, Third Horizon Research Group, ProgeniTech, and Superbaby Nutraceuticals™. Creator of the Primemester Protocol®, the Reproductive Readiness Index, the Taboo Terrain Assessment, the Burnout Personality framework, and the Couple-as-Fertility-Organ model.

Research disciplines. Reproductive health · intergenerational health · psychogenomics · epigenetics · social epidemiology · maternal-child-family health · lifespan development · developmental origins of health and disease (DOHaD).

And my co-lead. Dr. Amie Hornaman is the thyroid-literate physician most fertility clinics wish they could send their patients to. She prescribes T3-containing thyroid medication, bioidentical hormones, peptides, and GLP-1 receptor agonists when clinically indicated. She reads the full lab panel — not the single number. She knows when to treat and when to leave biology alone. I read your wife's life. Dr. Amie reads her labs. Together, that is the complete reading.

Why this matters. I am writing to you because the woman you love wants to address her thyroid and her hormones with the same rigor she would want for any other consequential decision in your life together. She may have asked you to read this because she wants your support. What follows is what I would say if we were sitting across a desk together.



The opening, stated plainly

Dear Sir,

The woman you love wants to address her thyroid and her hormones with seriousness, intelligence, and heart. She may have asked you to read this because she wants your support.

She may be afraid you will think it is too much money. She may be afraid you will think she is overthinking. She may be afraid you will say, "Your last labs said you were fine." She may be afraid you will only trust her primary care doctor, only trust the endocrinologist, or only trust time. She may be afraid that asking for one more investment — of money, attention, belief, or hope — will feel like too much.

So I want to explain, plainly and respectfully, what Total Thyroid & Hormone Fix with Thyroid Burnout Recode is, why it exists, why it is different from a standard thyroid workup or a fertility clinic's thyroid panel, and why supporting her in addressing this gland may be one of the most consequential decisions you make together — before, during, or after conception.

This is not a pressure letter. It is a translation.

She is not asking you to fix her. She is not asking you to guarantee an outcome. She is asking you to stand beside her while she addresses the gland that often has to be relieved before her ovaries, her eggs, her uterus, and her nervous system can come back online.



The bottom line

Total Thyroid & Hormone Fix with Thyroid Burnout Recode is a twelve-week, two-layer transformation co-led by Dr. Amie Hornaman and me.

It is for women trying naturally. It is for women preparing for IVF. It is for women already pregnant who need first-trimester thyroid targets met inside the closing window. It is for women who have been told their thyroid is fine when their body is telling them otherwise. It is for women with TSH at 3.8 that no clinician has flagged for forty-one months. It is for women whose free T3 is low and whose reverse T3 is high while their TSH "looks normal." It is for women with TPO antibodies one unit below the diagnostic threshold, walking into a fourth IVF cycle no one has explained.

It is for women in their thirties and forties who have been on oral contraceptives for fifteen years and now wonder why their testosterone is depleted, their libido is gone, and their cycles never quite normalized. It is for women whose insulin is high-normal, whose ferritin is below seventy, whose vitamin D is below thirty, whose homocysteine is climbing — all markers their primary care doctor called "fine."

It does not replace medical care. It does not diagnose. It does not treat. It does not promise pregnancy.

It does something different. The Clinical Layer — led by Dr. Amie Hornaman — orders the full panel, reviews her labs in context, and prescribes thyroid medication, bioidentical hormones, and other clinically indicated medications when her labs and symptoms call for them. The Psychogenomic Layer — led by me — addresses the survival pattern that burned her thyroid out in the first place, so the cofactors and the medication can finally land at the cellular level instead of being intercepted by the chronic cortisol biochemistry of a body that has been running on urgency for years or decades.

Dr. Amie reads her labs. I read her life. Both are necessary to bring her thyroid, her eggs, and her full HPTAO axis back from burnout so they have the power to create new life.



Why the thyroid deserves its own container

The thyroid is a small, butterfly-shaped gland at the front of her neck. It weighs less than an ounce. It is small enough that you could miss it with your hand if you were not looking for it.

It is also one of the most consequential glands in her body — for her own metabolism and energy, for her ovaries and eggs, for her capacity to conceive, for her capacity to stay pregnant, for the neurocognitive development of the child she may carry, and for the lifelong health trajectory of that child after birth.

The thyroid does not work in isolation. It sits at the center of an axis we call HPTAO — Hypothalamic-Pituitary-Thyroid-Adrenal-Ovarian. Five organs that share the same hypothalamus, the same pituitary, and the same survival-versus-thriving logic. The HPA axis — Hypothalamic-Pituitary-Adrenal — governs both her thyroid and her ovaries. When her HPA axis is in chronic survival mode, her thyroid is suppressed and her ovaries are suppressed in the same biochemical breath.

This is why a woman with a "normal" TSH can still be cold, exhausted, anovulatory, anxious, hair-shedding, and unable to conceive. The HPA axis is overriding the HPT axis, which is overriding the HPG axis. Three suppressions, one underlying signal.

This program is Primemester-level work concentrated specifically on the thyroid and the HPA-axis that governs both the thyroid and the eggs. The Primemester Protocol — my comprehensive 120-day preconception program — addresses the full reproductive system across every layer of biology and biography. It includes thyroid and HPA-axis work as part of its full scope, alongside dozens of other interlocking systems. Total Thyroid Fix takes the same depth of science and rigor and concentrates it specifically on the thyroid and the HPA-axis — because the thyroid is so critical for fertility, pregnancy, fetal programming, and offspring outcomes that it earns its own dedicated container.

If she also needs the broader preconception readiness work across the full reproductive system, a separate scientific brief exists for the Primemester Protocol. She can request it from us.

IV

What the science says about your future child

I want to give you the numbers directly, because the data on maternal thyroid function and offspring outcomes is among the most consequential in modern fertility medicine — and almost none of it is taught to expectant parents before they conceive.

In the landmark Haddow study, children born to mothers with significant untreated thyroid deficiency during pregnancy scored, on average, **seven IQ points lower** at ages seven to nine than children of mothers with normal thyroid function. *Haddow JE et al., 1999. New England Journal of Medicine, IF 158.5.*

In the Korevaar cohort, even **mild maternal hypothyroxinemia** in the first trimester — meaning low free T4 with a normal TSH — was independently associated with elevated risk of expressive language delay, attention problems, autism spectrum traits, and reduced offspring IQ. *Korevaar TIM et al., 2016. Lancet Diabetes & Endocrinology, IF 44.0.*

In the Thangaratinam meta-analysis, maternal TPO antibody positivity in early pregnancy was **independently associated with miscarriage and preterm birth** — even in women with completely normal TSH and free T4. The antibodies themselves carry reproductive risk. *Thangaratinam S et al., 2011. BMJ, IF 93.3.*

The American Thyroid Association now recommends a TSH target **below 2.5 mIU/L in the first trimester** and **below 3.0 mIU/L in the second and third trimesters** — substantially tighter than the general-population range of 4.5. Most women conceiving today are conceived into without ever being told this target exists. *Alexander EK et al., 2017. American Thyroid Association Guidelines. Thyroid, IF 6.4.*

The mechanism is not abstract. Your future child has no functioning thyroid gland of her own until approximately week 16 to 20 of gestation. From conception until then, her neurocognitive development depends entirely on her mother's thyroid hormone crossing the placenta. Maternal T4 enters fetal neurons. Fetal deiodinases convert it to T3. T3 drives cortical neurogenesis, neuronal migration, and the foundational architecture of her brain.

If her mother's thyroid is suppressed during those twelve weeks, your child's cortex develops with less raw material than it needs. The effect is dose-dependent and, in the most consequential window, irreversible.

This is not fear-based copy. This is what the peer-reviewed literature says, in the highest-impact journals in medicine.

Addressing her thyroid before conception is not optional. It is the single most consequential preparation she can make for your child's lifelong neurocognitive health.



What changes at home when her body comes back online

You do not need a study for this part. You have lived it.

When her thyroid and her hormones are off, what you see at home is a woman who is exhausted before the day begins. Foggy in the afternoon. Cold when no one else is cold. Patient with the children, with her work, with everyone except herself. Slower to laugh. Slower to want sex. Slower to be the woman you remember from before health became the third person in your relationship.

You may have stopped initiating because the answer is always no, or always a careful yes that does not feel like yes. You may have stopped asking how she is because the answer is always "tired." You may have started to wonder whether what you are missing is something biological, something relational, or something that left somewhere along the way and is not coming back.

When her thyroid comes back online — and her hormones come back into balance with it — what returns is not abstract.

Her smile. Her energy at the end of the day. Her playfulness with the children and with you. Her patience without the cost of self-erasure. Her desire — for sex, for connection, for the parts of life that depleted biochemistry could not afford. Her laughter, which had become rare without either of you noticing exactly when.

This is what dynasty biochemistry looks like in a marriage. A regulated HPTAO axis is not just a fertility outcome. It is the biological substrate of intimacy, of partnership, of the relational field your future child will be born into.

When she comes back online, you do not get a happier wife. You get back the woman you met. And she gets back the woman she remembers being before her thyroid and her nervous system decided survival had to come first.

This is one of the unspoken returns on the investment. It is not what she will lead with when she asks you to support this. But it is what is at stake.

WHAT SHE MAY BE EXPERIENCING

Common signals of thyroid and hormone dysregulation in women

- ✦ Exhaustion before the day begins
- ✦ Cold extremities; cold when no one else is
- ✦ Brain fog, especially in the afternoon
- ✦ Hair shedding; thinning eyebrows
- ✦ Weight that will not move despite real effort
- ✦ Constipation; sluggish digestion
- ✦ Low libido; vaginal dryness
- ✦ Irregular, short, or missing cycles
- ✦ Low mood; anxiety; flattened affect
- ✦ Slow recovery from workouts and illness
- ✦ Post-OCP cycles that never normalized

3+

If you recognize three or more of these symptoms,

it is a clinical signal that her thyroid and broader hormone terrain deserve a full reading. Not a single TSH. The full panel.

VI

You are not supposed to be her endocrinologist

Most good men in this position carry two things they rarely say out loud.

The first: I love her, and I want to help her, and I do not know how. The second: I am afraid of getting it wrong, of saying the wrong thing, of making her feel worse, of recommending something that does not work, of being the one who could have prevented something and did not.

You are not supposed to know how to fix this alone.

You are not supposed to be her endocrinologist, her reproductive immunologist, her psychogenomicist, her fertility nutritionist, her nervous-system regulator, her supplement coordinator, her insurance negotiator, and her partner — all at once. No one is supposed to be all of those things at once. Not even her doctor.

When you support her in this transformation, you are doing something specific. You are handing the **science, lab interpretation, prescriber strategy, and medical decision-making** to a thyroid-literate physician who lives inside this work every day. You are handing the **deep emotional, psychogenomic, and nervous-system architecture** to a scientist who has been doing this work for thirty years.

And you get to go back to being her *partner*. Not her primary care doctor. Not her therapist. Not her advocate at every clinical appointment. Not her supplement researcher at midnight. Not the person who has to know whether her TSH should be 2.5 or 1.5 or 3.0 depending on the trimester.

You get to be the man who loves her. That is what she actually wants from you. The rest of it is what she wants from us.

VII

What is actually included

Twelve weeks. Two layers. Eighteen live sessions co-led by Dr. Amie Hornaman and me. Plus the complete Recode curriculum delivered between sessions.

The Clinical Layer — led by Dr. Amie Hornaman.

- ◆ **Her full thyroid lab panel** — TSH, free T4, free T3, reverse T3, TPO antibodies, and thyroglobulin antibodies. Not just TSH. The complete panel, ordered for her, included in her tuition.
- ◆ **Her full reproductive hormone panel** — estradiol, progesterone, total testosterone, free testosterone, SHBG, DHT, DHEA-S, LH, FSH, and prolactin. So her thyroid, her adrenals, her sex hormones, and her whole hormone terrain can be read together — never in isolation. The panel most fertility clinics never order in this combination.
- ◆ **Her foundational bloodwork** — a complete blood count (CBC) reading her red blood cells, white blood cells, platelets, hemoglobin, hematocrit, and red blood cell size markers; and a comprehensive metabolic panel (CMP) reading her glucose, calcium, electrolytes, kidney function, liver enzymes, and protein status. The full reading of her terrain — because the map cannot be built from one lonely number.
- ◆ **Her dedicated prescriber appointment with Dr. Amie Hornaman** — a thyroid-literate physician who reviews her full lab set, not just one marker, and prescribes thyroid hormone medication when clinically indicated, including T3-containing options chosen for her biology rather than the formulary default.
- ◆ **Bioidentical hormone prescriptions in that same appointment** — when her labs, her symptoms, and her goals indicate them. The right form, the right level, the right mode of administration.
- ◆ **GLP-1 receptor agonist prescription in that same appointment** — when clinically indicated, with the essential pre-conception conversation about why a GLP-1 needs to be stepped off well before conception and how long beforehand, so it supports fertility rather than working against it.
- ◆ **Education on selective peptides for fertility** — including BPC-157, Thymosin Alpha-1, Epitalon, and the CJC-1295/Ipamorelin combination. These four candidates have theoretical or early-stage mechanistic relevance to fertility biology, with the strength of fertility-specific evidence varying substantially across them. She receives the evidence summary in writing, so she and her clinical team can make informed decisions.

PEPTIDE EVIDENCE — AT A GLANCE

PEPTIDE	PRIMARY MECHANISM	EVIDENCE STAGE
Epitalon	Oocyte oxidative stress; mitochondrial function	<i>Preclinical oocyte-aging data</i>
CJC-1295 / Ipamorelin	GH and IGF-I signaling; ovarian response	<i>Plausible via GH biology</i>
Thymosin Alpha-I	Immune modulation; reproductive immunology	<i>Indirect; early fertility data</i>
BPC-157	Tissue repair; angiogenesis; anti-inflammatory action	<i>Preclinical; extrapolated</i>

The Psychogenomic Layer — led by Dr. Cleopatra.

- ◆ **Her Thyroid Terrain Assessment** — a structured read of the conditions her thyroid has been living inside: stress load, depletion, under-recovery, nourishment, identity, pace, symptom pattern, emotional patterning, and the body's long-term conservation signals.
- ◆ **Her Burnout Personality Profile** — the diagnostic that identifies the cognitive, behavioral, and physiological patterns that drove her HPTAO axis into conservation biochemistry in the first place.
- ◆ **Her Personalized Thyroid Terrain Profile** — a written diagnostic of how her thyroid, her adrenals, her ovaries, her nervous system, her immune system, and her epigenome are currently interacting. Hers alone.
- ◆ **Her Psychogenomic Excavation across the lineage** — backward through her childhood, her in utero environment, her own Primemester programming, the lived environments of her parents and grandparents; and forward into what you both want epigenetically transmitted to your superbaby and your lineage.
- ◆ **The 7 Taboo Domains work** — money, sex, motherhood, food, body, love, and power. The private domains where shame quietly shapes her hormones, her hunger, her desire, her ambition, her depletion. The domains no other doctor, coach, therapist, or program will go with her.
- ◆ **Her Reparenting and Reprogramming Protocol** — psychoneuroimmunology and epigenetic tools that teach her nervous system, immune system, endocrine system, and epigenome that the old survival pattern is no longer the only option. So the cofactors and the medication Dr. Amie has prescribed can finally land at the cellular level.
- ◆ **Her Allied Advocacy Framework** — word-for-word scripts for the conversations that determine her care: how to talk with her OB-GYN, REI, or IVF clinic without triggering professional defensiveness; how to request the panels she needs; how to bring research to a clinician while honoring their authority. Plus the scripts for the conversation with you, with family, with friends.

- ◆ **Installation of her new Central Nervous System Operating System** — daily practices that build her new physiological baseline: parasympathetically accessible, oxytocin-permissive, metabolically alive, anti-inflammatory, hormonally receptive.
- ◆ **Her twelve-week Tracking and Maintenance Protocol** — the structured system that holds the new pattern in place long after the twelve weeks end. So she does not return to burnout the next time life gets demanding.

VIII

The financial frame, stated plainly

Let's speak plainly about money.

She can get started for \$694 today — with two additional payments of \$694, or one single payment of \$1,697 that saves \$385. For that, she receives eighteen live sessions with Dr. Amie Hornaman and me, the complete Recode curriculum, the full lab panels listed above, the prescriber appointment, the prescription decisions, the psychogenomic excavation, the Allied Advocacy Framework, and the twelve-week tracking protocol.

Dr. Amie and I negotiated the lab and prescriber rates inside this container so the clinical workup is covered in her tuition. The lab panel alone, ordered à la carte across a thyroid-literate functional medicine practice, runs \$1,200 to \$2,400 depending on location. The comparable specialist version of this work — done across an endocrinologist, a functional medicine doctor, a fertility nutritionist, a therapist, and a coach, without the integration — runs \$15,000 to \$25,000.

We built the integration for her so she does not have to mastermind it herself.

Now consider the cost of not doing this.

A single failed IVF cycle in the United States runs \$15,000 to \$30,000 before medications, before genetic testing, before transfer. A second cycle frequently doubles that figure. The literature is clear that subclinical maternal thyroid dysfunction and undiagnosed TPO autoimmunity contribute meaningfully to IVF failure and recurrent loss — neither of which would survive contact with a properly read panel.

A NICU admission for a preterm infant — a documented risk in mothers with untreated thyroid autoimmunity, even with normal TSH — averages \$76,000 in the United States for a moderately preterm baby, and can exceed \$500,000 for a very preterm infant. The thyroid is not the only contributor to preterm birth. But it is one of the most reversible contributors, and addressing it before conception is among the highest-leverage interventions available.

A lifetime of unaddressed maternal thyroid autoimmunity — Hashimoto's in particular — frequently progresses from subclinical to clinical to medication-dependent across her thirties and forties. Reversing the trajectory while she is still on the autoimmune approach is meaningfully different from managing a disease that has crossed into full diagnosis.

\$694 to get started is not the expensive option. The expensive option is the path most couples are already on. Lab after lab. Practitioner after practitioner. Cycle after cycle. With no one organizing the whole picture and no one reading her life alongside her labs.

If you are going to spend money on her fertility and her future health, the question is not what does this cost. The question is what is the cost of continuing without coherence.

THE MATH, AT A GLANCE

Get started today

\$694 *first of three payments*

Or one single payment

\$1,697 *saves \$385*

À la carte lab cost (comparable)

\$1,200 – \$2,400 *before any practitioner fees*

Comparable specialist version

\$15,000 – \$25,000 *across 5+ providers, no integration*

A single failed IVF cycle

\$15,000 – \$30,000 *before medications and testing*

A NICU admission, moderately preterm

\$76,000 *avoidable risk reduction*

IX

Why this is not a fertility clinic thyroid program

Most fertility clinics check TSH. That is the end of their thyroid work.

A few will add free T4. A small number will add antibodies. Almost none will add free T3, reverse T3, the FT3:rT3 ratio, the full hormone panel read in context, the cortisol patterning that reveals the HPA-axis collision, the ferritin floor required for thyroid peroxidase activity, the vitamin D level required for TPO antibody modulation, the homocysteine level that reveals methylation status, or the insulin and HbA1c that reveal whether her metabolic terrain is fertile.

This is not because clinicians are uncaring. It is because the standard fertility workup was built to identify women who already have overt thyroid disease. It was not built to identify the women who are walking into their fourth IVF cycle with chronically high-normal TSH and TPO antibodies one unit below the diagnostic line. It was not built to identify the women whose thyroid is being suppressed at the brain level by chronic HPA-axis activation. It was not built to identify the women whose ferritin at 22 ng/mL is functionally hypothyroid even though her TSH is "normal."

A standard thyroid workup catches the most severe presentations. A TSH of 8, antibodies of 300, overt disease. By the time those numbers appear, the thyroid has been deteriorating for years.

The Total Thyroid Fix catches the trajectory. It sees the 41-month climb that no single data point would flag. It sees the antibody rising from 18 to 34 over three years and recognizes that a woman sitting one unit below the diagnostic line today is on the line within months if nothing intervenes.

This is the work most clinicians cannot do — not because they are uncaring, but because they are seeing her for twelve minutes and reading one panel at a time. It is structural. It is not their fault. But it is also not enough for the woman who is thirty-eight, has been trying for three years, and has a folder of lab results that no one has read across time.



You might need this too

I want to say something directly to you, because no one has likely said it to you yet.

Your testosterone matters too. Your thyroid matters too. Your insulin, your inflammation, your sleep architecture, your cortisol pattern, your sperm DNA integrity — all of these are responsive to the same psychogenomic pressures that have suppressed her HPTAO axis. You are sharing a household, a sleep schedule, a financial environment, a relational stress field, and an exposure terrain. The biological field your future child enters is created by both of you.

The numbers on men's testosterone are sobering.

- ◆ **Among men in their 20s**, roughly **20%** have total testosterone below the reference range used for younger men, and a significantly higher proportion have *suboptimal* levels for fertility, libido, and metabolic health. Many of these men do not yet know.
- ◆ **Among men in their 30s**, the prevalence of low testosterone climbs to roughly **25 to 30%**. This is the decade in which most men begin trying to conceive — and the decade in which the first symptoms (low libido, slower recovery, abdominal weight gain, mood flattening, sleep disruption) are often dismissed as "stress" or "just getting older."
- ◆ **Among men in their 40s**, the figure rises to roughly **35 to 40%**. This is the decade in which testosterone-related fertility concerns most commonly intersect with a partner's own fertility timeline. *Travison TG et al., 2007. Journal of Clinical Endocrinology and Metabolism, IF 5.8.*
- ◆ **Among men in their 50s**, prevalence exceeds **50%** by most large cohort estimates, with the steepest declines occurring in men carrying chronic metabolic, stress, and inflammatory burden — exactly the men most likely to be supporting a partner through a complex fertility season.

Here is what most men do not know. If you are trying to conceive — or if you might in the foreseeable future — you cannot go to a standard low-T clinic for help. Standard testosterone replacement therapy **damages fertility**. Exogenous testosterone suppresses LH and FSH release at the pituitary, shutting down endogenous testosterone production and dramatically reducing sperm production. The very treatment that would restore your libido and energy at a generic men's health clinic would compromise your contribution to the child you and your partner are trying to create.

The fertility-supporting path for men with low testosterone is fundamentally different from the standard path. It involves clomiphene, hCG, selective peptides, lifestyle and metabolic recoding, sleep and HPA-axis work, and occasionally enclomiphene — **not** exogenous testosterone. This is work Dr. Amie can do alongside you if your labs indicate it. It is work no standard low-T clinic will offer, because the standard low-T clinic does not have a fertility lens.

Joining her in this transformation, if your own labs warrant it, is not just for you. It is for the half of the embryo that comes through you. Sperm DNA methylation, sperm small RNA cargo, sperm mitochondrial health — all of it shapes the child you and your partner are creating.

If your wife or partner has asked you to read this, consider whether you might want to walk through the door with her. Not as a patient. As a co-creator.

WHAT YOU MAY BE EXPERIENCING

Common signals of low testosterone and thyroid dysregulation in men

- ✦ Morning erections diminished or absent
- ✦ Abdominal weight gain that resists exercise
- ✦ Slower recovery from workouts
- ✦ Low libido; reduced spontaneous desire
- ✦ Mood flattening; less playfulness
- ✦ Sleep disruption; non-restorative sleep
- ✦ Cognitive slowness; reduced mental stamina
- ✦ Irritability; shorter fuse than before
- ✦ Reduced strength or stamina at the gym
- ✦ Lower confidence; lower drive at work

3+

If you recognize three or more of these symptoms,

it is a clinical signal that your thyroid, testosterone, and broader hormone terrain deserve a full reading — with a fertility lens.

XI

What your "yes" gives her

When you say yes to this, what she hears is not "you supported the program." What she hears is:

"You are not crazy. What you have been feeling is real, and it matters to me."

"You do not have to carry this alone anymore."

"Our future child, and how you feel inside your own body, are worth investing in with the depth they deserve."

"I see what you have been holding for this family, and I am ready to hold some of it with you."

That is a powerful thing for a woman to feel from her partner. It is the kind of thing that changes the field she is trying to conceive inside.

A held woman responds to medication differently. A held woman ovulates differently. A held woman conceives differently. A held woman carries differently. A held woman recovers from birth differently. A held woman mothers differently.

This is not sentimental. It is physiological. Social support has been shown across large bodies of research to buffer HPA-axis activation, improve immune function, reduce inflammatory burden, and shape reproductive outcomes including birthweight, preterm risk, and infant health markers. Your support is not soft. Your support is biological infrastructure.

You are not just signing off on a program. You are stepping into a role that may shape the next several decades of your life together — and the lifetime of your child.

XII

The decision in front of you

The decision is not: can we afford this? The decision is: what is the cost of continuing without it?

The decision is not: will this guarantee a baby? No one can guarantee that — not Dr. Amie, not me, not any clinic, not any protocol, not any prayer. The decision is: will she go into the most consequential biological window of your lives together with the deepest intelligence available to her, or will she walk into it the way most women do — alone, scattered, advocating for tests no one wants to order, taking medication no one has explained?

The decision is not: should we wait until she has a clearer diagnosis? Waiting passively, for a woman with a body that is already signaling, is rarely the most intelligent plan. Subclinical thyroid dysfunction does not stay subclinical forever. TPO antibodies do not stay below threshold forever. Ovarian reserve does not wait. Egg quality does not wait. Her brain on chronic cortisol does not wait. Your future child's first-trimester window does not wait.

The decision is whether you will walk through this door together.

No one can guarantee your baby. But you can guarantee your standard.

Preparation is a form of love. And love leaves evidence.

With respect for the role you play,

Dr. Cleopatra Kamperveen, Ph.D.

Fertility & Pregnancy Scientist

Retired Tenured Professor, University of Southern California

Founder, The Fertility & Pregnancy Institute

Creator, Primemester Protocol® and Superbaby Nutraceuticals™

Primeia® · Ovalaire®



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All sources peer-reviewed · journals at recognized impact factor standards

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