Deciphering Hormones with the Hormone Insights Test (HIT)

Charlotte Hunter
KBMO Diagnostics UK





What are we covering today?

- Recap of the FIT Test
- Why hormone testing?
- The Hormone Insights Test (HIT)
- Sample reports
- A note on support & education
- Please ask questions as we go...



Meet the UK Team



Charlotte Hunter Head of KBMO UK



Clare Kennedy Operations Manager



Linette Petrillo Customer Services



Natasha Khan Sales



Kelly Hutson Events



Emily Birch Clinical Support



The Food Inflammation Test (FIT)









Summary of Tests

FIT22

22 of the most common food sensitivities including gluten, cow's milk and egg.



176 foods including health foods such as honey, stevia and coconut oil plus the Gut Barrier Panel

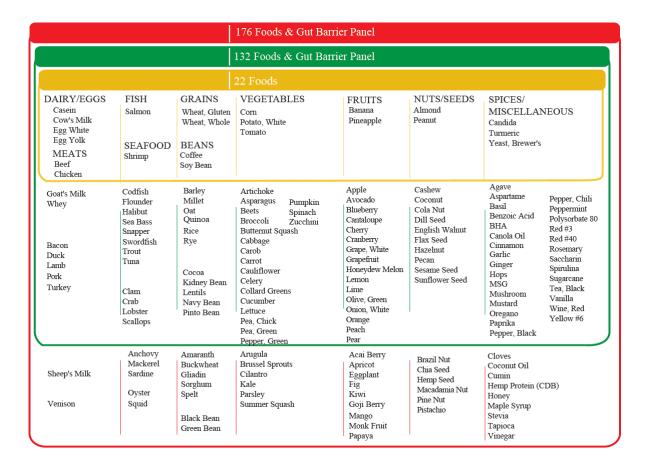
FIT132

132 foods and food additives plus the Gut Barrier Panel



Assess the integrity of the gut lining by measuring Candida, Zonulin, Occludin and Lipopolysaccharides (LPS)

Foods We Test

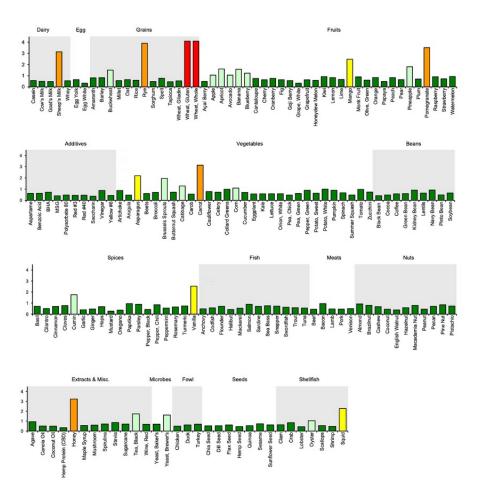


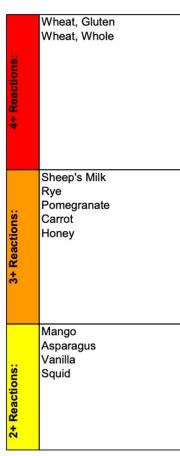
Food selection contains 'real-world' foods, encompassing raw and cooked.

The FIT132 and FIT176 both include the Gut Barrier Panel.



The Food Inflammation Test (FIT) Report





- Easy to read
- Colour coded
- Client-friendly
- Easy interpretation
- Efficient practice
- Easy to repeat tests



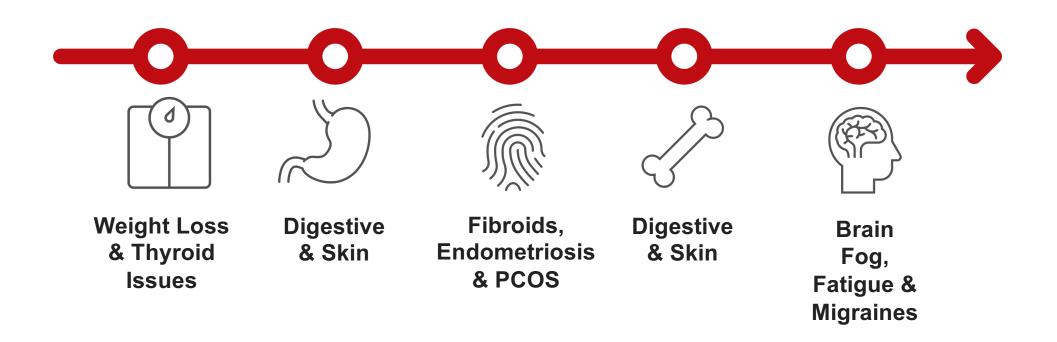
The Gut Barrier Panel

Gut Barrier Panel

KBMO has created a unique Gut Barrier Panel which in recognition that leaky gut occurs across a spectrum we have included the following gatekeeper markers: Candida, Zonulin and Occludin and LPS. For each marker, we measure IgG 1-4 /C3d in addition to IgA 1 and 2.

Gut Barrier Panel							
		lgG1-4+C3d	IgA1-2				
		Cut off		Cut off			
Candida	Negative		Positive				
Zonulin	Negative		Positive				
Occludin	Negative		Negative				
LPS	Negative		Positive				

Why test food sensitivities?





Hormone Testing

- Are your current hormone tests giving you what you need?
- Many hormone tests are:
 - Complicated to interpret
 - Expensive for clients
 - Less clear since report updates
 - Confusing and often contain errors in their algorithm-generated interpretations

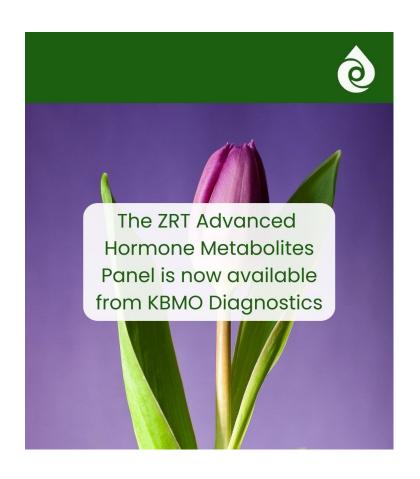


There IS Another Way!



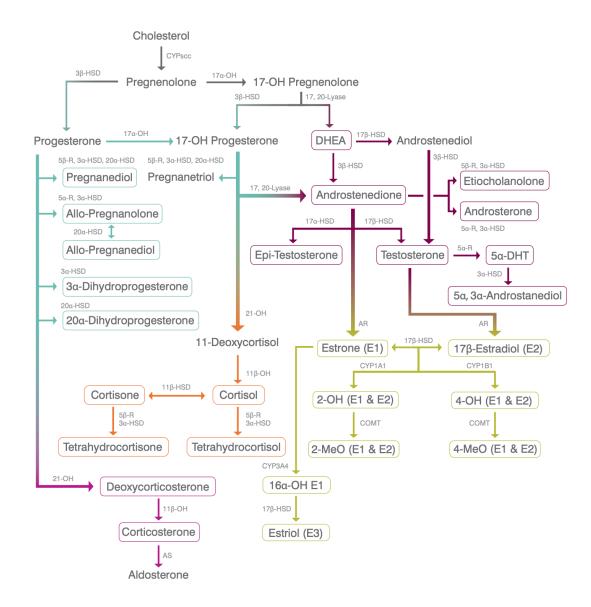
- The Hormone Insight Test (HIT): Powered by the Advanced Urine Hormone Metabolites Test by ZRT
- Measures 44 hormone-related markers
- 13 oestrogens, 8 androgens
- Diurnal cortisol & melatonin patterns
- Includes BPA (rarely assessed endocrine disruptor)

The Hormone Insights Test (HIT)



- Developed by ZRT laboratory pioneers in hormone testing
- The ORIGINAL urine metabolites test
- Built on decades of research and clinical use
- The HIT Test combines ZRT's scientific credibility with KBMO's practitioner-first support model.

The Steroid Hormone Cascade



Why Does the HIT Measure?

More Tests:

With 44 unique markers, ZRT's urine hormone profiles are more comprehensive than any other lab.

More Oestrogens:

ZRT assesses a total of 13 oestrogens, including 2-Methoxy and 2-Hydroxy, 4-Hydroxy and 4-Methoxy and 16a-Hydroxy oestrogens. We test more 4-Hydroxy metabolites than any other lab.

More Androgens:

ZRT assesses a total of 8 androgens, which is more than most labs offer.



Why Does the HIT Measure?

BPA: One of few labs to include an assessment of BPA in its urine hormone testing.

Diurnal Cortisol:

ZRT provides diurnal patterns for both cortisol and cortisone, which help get to the root of stress-related issues. It's also ideal for those unable to collect a saliva sample for diurnal cortisol.

Diurnal Melatonin:

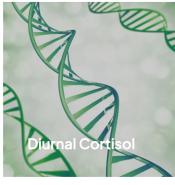
Diurnal Melatonin: ZRT is the only lab to include a diurnal pattern for melatonin, which helps evaluate sleep-related issues.



The ZRT Difference









- More Markers
- More Insight
- More clinical value

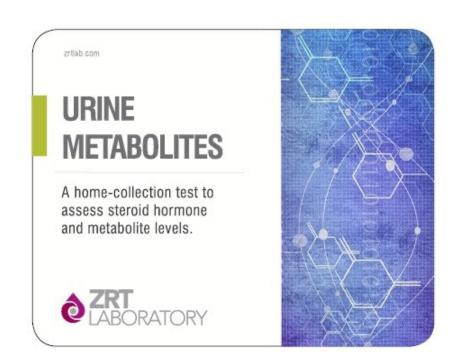
Why is HIT Different?

- Why should you be excited about the HIT?
- Relevant data especially around progesterone which is notoriously under-assessed in other tests)
- Clarity simple reference range bars and easy-to-read graphs
- At-home convenience competitive pricing for clients with good margins for practitioners



Why Use Urine – and Why Dried?

- Dried urine testing is a simple, noninvasive way to evaluate hormone activity and detoxification.
- Unlike other methods, it shows both levels and pathways. In other words, it reveals how hormones are used and cleared.
- ZRT pioneered this science-backed method and continues to lead the field with clinical precision and innovation.



When to Consider the HIT?

- Family history of hormone-dependent conditions or cancers
- Adrenal issues or hormonal imbalance such as weight gain and insomnia
- Symptoms of PCOS, such as acne and excess facial hair
- Symptoms of menopause and/or ready to begin hormone replacement therapy
- Symptoms of oestrogen dominance while on physiological replacement dosages of therapy





TEST REPORT

8605 SW Creekside Place
Beaverton, OR 97008
Phone: 503-466-2445 Fax: 503-466-1636

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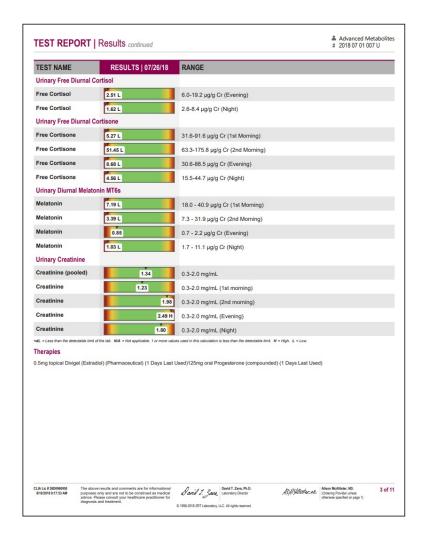
Ordering Provider: Getuwell Clinic Jim Getuwell, DO

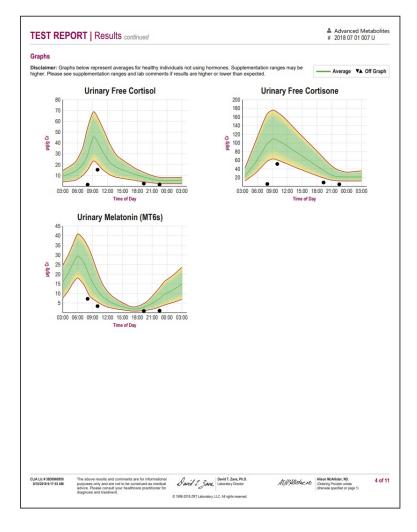
Samples Received 07/31/2018 Urine - 07/26/18 08:00 Urine - 07/26/18 10:00 Urine - 07/26/18 19:20 Urine - 07/26/18 22:30

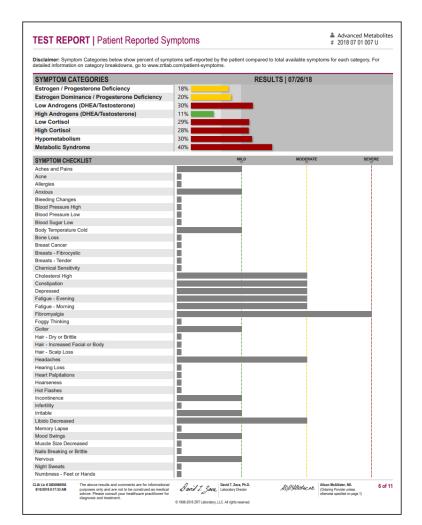
Patient Name: Advanced Metabolites Patient Phone Number: 555 555 5555

Gender Female	Last Menses Unspecified	Height 5 ft 3 in	
DOB 3/4/1960 (58 yrs	Menses Status s) Postmenopausal	Weight 165 lb	
TEST NAME	RESULTS 07/2	6/18	RANGE
Urinary Estrogens			
Estradiol	0.68 L		0.78-1.79 μg/g Cr Premeno-luteal or ERT
Estrone	2.85		2.27-5.22 μg/g Cr Premeno-luteal or ERT
Estriol	0.83		0.78-1.98 μg/g Cr Premeno-luteal or ERT
E3/(E1+E2)	0.24 L		>0.3 (> median value)
2-OH Estradiol	0.21		0.17-0.70 μg/g Cr Premeno-luteal or ERT
2-OH Estrone	1.09		0.70-2.54 μg/g Cr Premeno-luteal or ERT
4-OH Estradiol	0.15		0.10-0.18 μg/g Cr Premeno-luteal or ERT
4-OH Estrone		0.47	0.17-0.47 μg/g Cr Premeno-luteal or ERT
16α-OH Estrone	0.31 L		0.35-1.07 μg/g Cr Premeno-luteal or ERT
2-OH (E1 + E2)/16- OH E1	α- 4.19		1.29-5.49 Premeno-luteal or ERT
2-MeO Estradiol	0.06		0.03-0.08 μg/g Cr Premeno-luteal or ERT
2-MeO Estrone	0.51		0.26-0.68 μg/g Cr Premeno-luteal or ERT
2-MeO E1/2-OH E1		0.47 H	0.21-0.38 Premeno-luteal or ERT
4-MeO Estradiol		0.05 H	<0.04 µg/g Cr
4-MeO Estrone		0.12 H	<0.04 µg/g Cr
4-MeO E1/4-OH E1		0.26 H	0.05-0.13 Premeno-luteal or ERT
4-MeO E2/4-OH E2		0.33 H	0.10-0.29 Premeno-luteal or ERT
Bisphenol A	<di l<="" td=""><td></td><td>1.5-4.5 µg/g Cr Postmenopausal</td></di>		1.5-4.5 µg/g Cr Postmenopausal
8/10/2018 9:17:53 AM pt	ne above results and comments are for infor irposes only and are not to be construed as twice. Please consult your healthcare practit agnosis and treatment.	medical	David T. Zann. David T. Zenn. Ph.D. ADM/Shitsher AD. (Odering Positive unions) (Odering Specified on page 1)

TEST NAME	RESULTS 07/26/18	RANGE			
Urinary Progestogens					
Pregnanediol	8071 H	465-1609 μg/g Cr Premeno-luteal or PgRT			
Allopregnanolone	35.33 H	2.23-14.87 µg/g Cr Premeno-luteal or PgRT			
Allopregnanediol	148.31 H	14.65-76.71 μg/g Cr Premeno-luteal or PgRT			
3α- Dihydroprogesterone	3.29 H	0.67-2.03 μg/g Cr Premeno-luteal or PgRT			
20α- Dihydroprogesterone	15.13 H	3.93-11.62 µg/g Cr Premeno-luteal or PgRT			
Deoxycorticosterone	1.72	0.69-2.23 µg/g Cr Premeno-luteal or PgRT			
Corticosterone	1.48 L	3.19-9.59 µg/g Cr Premeno-luteal or PgRT			
Pgdiol/E2	11869.12 H	1000-1500 (Optimal Luteal Only)			
Urinary Androgens					
DHEA	6.44 L	8.63-37.28 μg/g Cr Postmenopausal			
Androstenedione	1.95 L	2.07-7.94 µg/g Cr Postmenopausal			
Androsterone	212	152-482 µg/g Cr Postmenopausal			
Etiocholanolone	195 L	239-777 µg/g Cr Postmenopausal			
Testosterone	0.91	0.66-2.89 μg/g Cr Postmenopausal			
Epi-Testosterone	0.78	0.39-1.32 μg/g Cr Postmenopausal			
T/Epi-T	1.17	0.5-3.0			
5α-DHT	0.24 L	0.26-0.98 μg/g Cr Postmenopausal			
5α,3α-Androstanediol	3.87	2.32-8.17 µg/g Cr Postmenopausal			
Urinary Glucocorticoids					
Total Cortisol	11.88 L	13.23-39.26 µg/g Cr Postmenopausal			
Total Cortisone	13.04 L	23.32-59.61 µg/g Cr Postmenopausal			
Cortisol/Cortisone	0.91 H	0.5-0.7			
Tetrahydrocortisol	192 L	281-711 μg/g Cr Postmenopausal			
Tetrahydrocortisone	530 L	551-1474 µg/g Cr Postmenopausal			
Urinary Free Diurnal Co	rtisol				
Free Cortisol	1.49 L	7.8-29.5 µg/g Cr (1st Morning)			
Free Cortisol	15.28 L	23.4-68.9 μg/g Cr (2nd Morning)			







TEST REPORT | Patient Reported Symptoms continued

Advanced Metabolites

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SYMPTOM CHECKLIST	MILD	MODERATE	SEVERE
Pulse Rate Slow	1		- 1
Rapid Aging			
Rapid Heartbeat			
Skin Thinning			
Sleep Disturbed			- 1
Stamina Decreased			1
Stress			
Sugar Cravings			1
Sweating Decreased			- 1
Swelling or Puffy Eyes/Face			1
Tearful			
Triglycerides Elevated			
Urinary Urge Increased			1
Uterine Fibroids			1
Vaginal Dryness			
Water Retention			
Weight Gain - Hips			
Weight Gain - Waist			

Lab Comments

PARENT ESTROGENS (ESTRADIOL-E2, ESTRONE-E1, ESTRIOL-E3)

The parent estrogens are within/near the expected median to 90 percentile of the reference ranges seen in postmenopausal women supplementing with topical estrogen replacement therapy (Note: transdermal and topical estrogens raise urinary estrogens very little as these estrogens are excreted predominately in bile and feces by this route of administration). This is commonly seen in postmenopausal women taking low dose topical estrogens (usually a topical estradiol or biestrogen containing estradiol + estriol). Topically delivered estrogens increase saliva and capillary blood levels of the supplemented estrogens, but increase urinary and serum levels much less. Topically delivered hormones are more likely to be excreted in bile/feces than in urine.

HYDROXYLATED (CATECHOL) ESTROGENS (2-OH E2 & E1, 4-OH E2 & E1, 16-OH E1) and 2-OH/16-OH RATIO

The hydroxylated estrogens are all within/near normal reference ranges for a postmenopausal woman supplementing with topical estrogen(s). Levels of the down-stream hydroxylated estrogens are usually within the low end of the reference ranges with topical ERT, as are the parent estrogens from which they are derived. Topically delivered estrogens raise the level of urinary estrogens very little, which is likely due to excrettion more in the hile/feces than in urine

The hydroxylation of estradiol and estrone represent the first phase of metabolism and elimination of these estrogens via urine. Following hydroxylation at the 2-, 4-, or -16 position, the estrogens undergo further modification (methylation, sulfation, glucuronidation) that increases their solubility and excretion in urine. In the laboratory these sulfate and glucuronide groups are removed by enzyme hydrolysis, which allows for measurement of the different types of hydroxylated estrogens, in addition to methylation of the hydroxyl groups (see below). The 2- and 4-hydroxylated E1 and E2 metabolites are referred to as catechol estrogens.

Research and clinical studies show that the 2-hydroxylated estrogens (2-OH E2 and 2-OH E1) are a safer pathway of hydroxylation than the 4hydroxyestrogens (4-OH E2 and 4-OH E1), which bind to and damage DNA, leading to mutations that are associated with increased breast cancer risk. For reviews see: Cavalleri EL, Rogan EG Future Oncol 6(1): 75-79, 2010; and Lee, JR, Zava DT What Your Doctor May Not Tell You About BREAST CANCER: How Hormone Balance Can Help Save Your Life: Chapter 7.

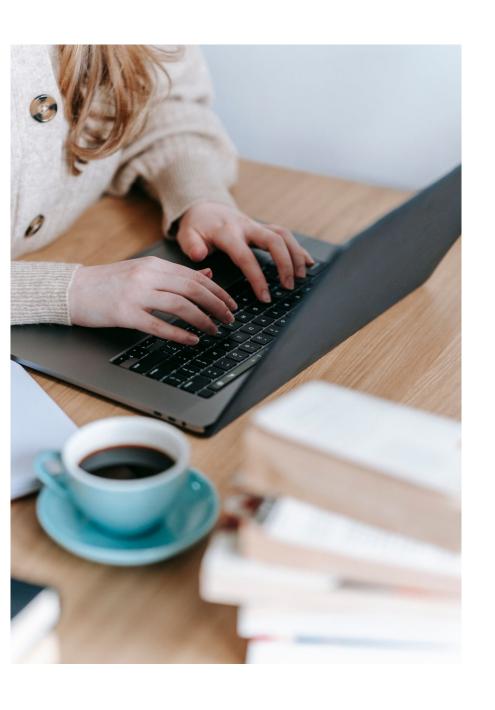
2-hydroxylated estrogen metabolism is increased with cruciferous vegetables and extracts of them, so higher consumption of these foods will enhance the safer 2-hydroxylation pathway for estrogen metabolism. The most commonly used concentrated extracts of cruciferous vegetables contain high levels of indoi-92-carbinot ((3C) and its metabolite diindoylmethane (DIM), Iodine also increases the 2-hydroxylation of estrogens, with a slight increase in 4-hydroxylation (Stoddard FR et.al. Int J Med Sci 5: 189-196, 2008), which is associated with the protective effects of higher dose iodine therapy for prevention of breast cancer. The more dangerous 4-hydroxylated estrogen metabolism is enhanced by exposure to environmental toxins, mostly petrochemical-based products but also heavy metals, that induce 4-hydroxylation pathway enzymes (1B1), and cause formation of Reactive Oxygen Species (ROS) that co-oxidize the catechol estrogens to quinones.

16-hydroxyestrone is another pathway of estrone metabolism and is a precursor to estriol (see Steroid Hormone Cascade). Early clinical research in humans suggested that a high urinary level of 16-hydroxyestrone relative to 2-hydroxylated estrogens (i.e. a low 2-OH E1 + 2-OH E2/16-OH E1 ratio), was associated with an increased risk of breast cancer in premenopausal women, but not in postmenopausal women. This has remained controversial and newer research suggests that while higher levels of 16-hydroxy estrone may indeed be slightly associated with

The above results and comments are for informational purposes only and are not to be construed as medical avivice. Please consult your healthcare practitioner for diagnosis and freatheast.

ADMISHISTOCAD. Alison McAllister, ND. (Ordering Provider unless otherwise specified on page 1)

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The Power of At-Home Testing

- Simple, non-invasive collection
- Less stressful, time-consuming and expensive!
- Better compliance
- Easy integration into your practice
- No need to stop hormones/supplements

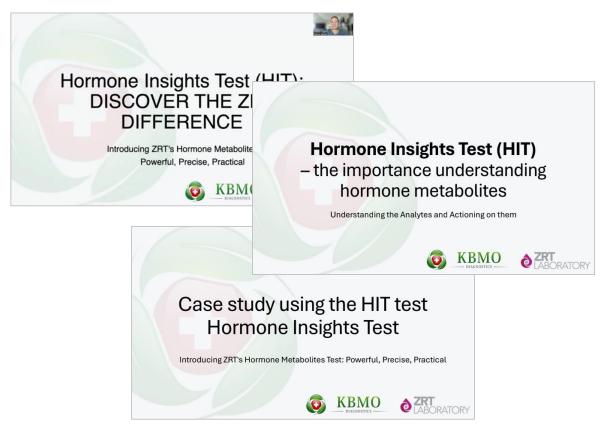
Support & Education

- 1st report → personal 1:1 walk-through
- Weekly Hormone Help Hour → group coaching, case studies and Q&A
- Supplement partnerships → practical, real-world application
- Education → one of KBMO's core values!
- And of course, we're always at the end of the phone or email!

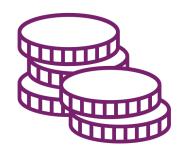




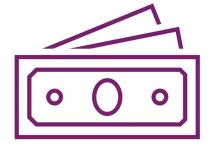
Dr Shania Seeber



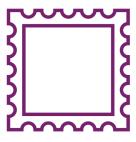
Pricing & Postage



Practitioner Price £249



Retail Price £299



FREE Postage & Returns

Your Next Steps with HIT

- Order your first test at www.kbmodiagnostics.co.uk
- Attend your first 1:1 session
- Bring your cases and questions to the Hormone Help Hour
- Enjoy the brilliant webinars with Dr Shania Seeber to build your knowledge on the HIT.



Smarter Testing, Better Outcomes



↑ HIT = insights that matter



Simple reports, relevant data



Support at every step



Future of hormone testing

Next Webinar: 16th September, 10am

Digestive Deep Dive:

The Dynamic Duo of Gut Testing in Action

with Dr Shania Seeber

Thank You



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