Nordic	Laboratories

TEST NUMBER: GENDER: Female AGE: 25

PATIENT

COLLECTED: 2022-03-24 06:01AM 2022-03-24 06:01AM 2022-03-24 06:01PM 2022-03-24 06:01PM

PRACTITIONER:

TEST REF:

TEST NAME: DUTCH Sex Hormone Metabolites

Test		Result	Units	Luteal*	Postmenopausal
Progesterone M	etabolites (Urine)			Range	Range
b-Pregnanediol	Below luteal range	285.0	ng/mg	600 - 2000	60-200
a-Pregnanediol	Below luteal range	48.6	ng/mg	200 - 740	15-50
Estrogens and M	Metabolites (Urine)				
Estrone(E1)	Above luteal range	33.6	ng/mg	12 - 26	1.0-7.0
Estradiol(E2)	Within luteal range	2.8	ng/mg	1.8 - 4.5	0.2-0.7
Estriol(E3)	Within luteal range	12.7	ng/mg	5 - 18	0.6-4.0
2-OH-E1	Within luteal range	7.7	ng/mg	5.1 - 13.1	0.3-2.0
4-OH-E1	Within luteal range	0.67	ng/mg	0 - 1.8	0-0.3
16-OH-E1	Within luteal range	1.41	ng/mg	0.7 - 2.6	0.2-0.6
2-Methoxy-E1	Above luteal range	8.08	ng/mg	2.5 - 6.5	0.3-1.4
2-OH-E2	Within luteal range	0.25	ng/mg	0 - 1.2	0-0.3
4-OH-E2	Within luteal range	0.2	ng/mg	0 - 0.5	0-0.1
Total Estrogen	High end of range	67.4	ng/mg	35 - 70	4.0-15
Androgens and	Metabolites (Urine)				
DHEA-S	Above range	940.6	ng/mg	20 - 750	
Androsterone	High end of range	1601.0	ng/mg	200 - 1650	
Etiocholanolone	Within range	816.0	ng/mg	200 - 1000	
Testosterone	Within range	6.8	ng/mg	2.3 - 14	
5a-DHT	Within range	3.15	ng/mg	0 - 6.6	
5a-Androstanediol	Above range	51.0	ng/mg	6 - 30	
5b-Androstanediol	Within range	31.9	ng/mg	20 - 75	
Epi-Testosterone	Within range	5.0	ng/mg	2.3 - 14	

*the Luteal Range is the premenopausal range. When patients are taking oral progesterone this range for progesterone metabolites is not luteal and reflects the higher levels expected when patients take oral progesterone. This test is intended to be taken in the luteal phase of the menstrual cycle (days 19-22 of a 28 day cycle) for premenopausal women. The ranges in the table below may be used when samples are taken during the first few days (follicular) of the cycle, during ovulation (days 11-14) or when patients are on oral progesterone. See the following pages for age-dependent ranges for androgen metabolites.

Additional Normal Ranges	Follicular	Ovulatory	Oral Pg (100mg)
b-Pregnanediol	100-300	100-300	2000-9000
a-Pregnanediol	25-100	25-100	580-3000
Estrone (E1)	4.0-12.0	22-68	N/A
Estradiol (E2)	1.0-2.0	4.0-12.0	N/A

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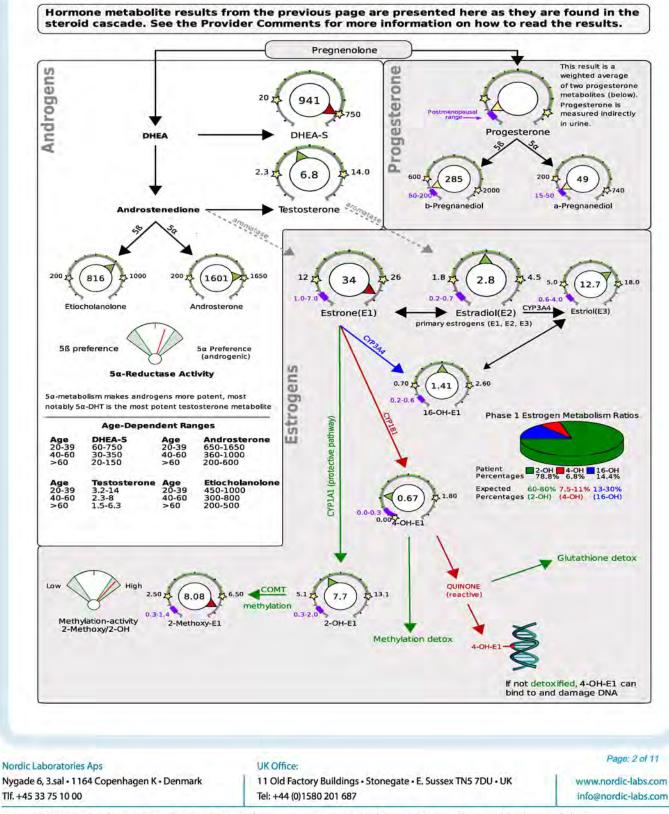
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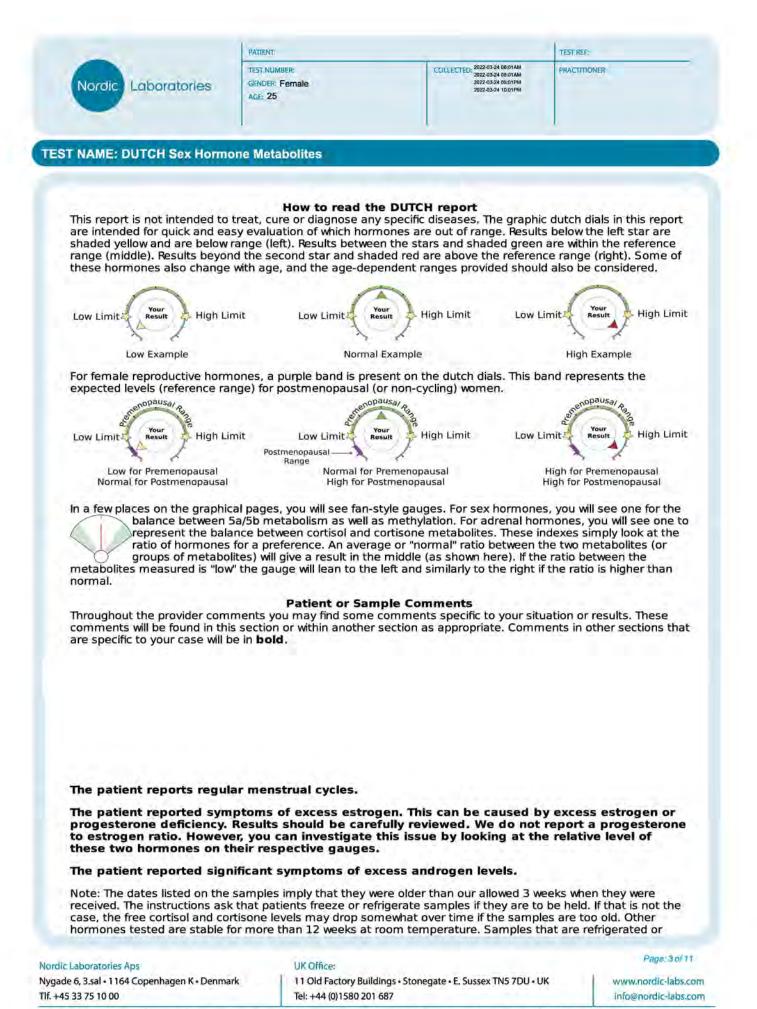
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1000	PATIENT:		TEST REF:
Nordic Laboratories	TEST NUMBER: GENDER: Female AGE: 25	COLLECTED: 2022-03-24 06:01AM 2022-03-24 06:01AM 2022-03-24 06:01PM 2022-03-24 10:01PM	PRACTITIONER:

TEST NAME: DUTCH Sex Hormone Metabolites







TEST NUMBER: GENDER: Female AGE: 25

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TEST NAME: DUTCH Sex Hormone Metabolites

frozen are stable for months.

Progesterone Metabolism

Progesterone is made predominately in the ovaries by the corpus luteum following the release of an egg. Progesterone metabolite levels will increase to the premenopausal luteal range (the range established as the green band between the two gold stars) only after the release of an egg. The level of progesterone metabolites seen on the DUTCH test can help determine if ovulation occurred 5-7 days prior to test collection.

The primary role of progesterone is to prepare the endometrium of the uterus for implantation. In addition, it may balance the effects of estrogen, it is a neurosteroid, it acts as a diuretic and raises basal body temperature.

We are measuring metabolites of progesterone 5b-pregnanediol and 5a-pregnanediol. 5b-pregnanediol has less activity in the body but does represent a larger percent of total progesterone metabolism overall. 5a-pregnanediol is often a metabolite of more interest, as it can cross the blood brain barrier and up-regulate GABA activity and is considered neuroprotective to the brain. In some women the 5a-pregnanediol is also the cause of PMDD and irritability due to issues with the GABA receptor's inability to adjust for sensitivity to fluctuating neurosteroids (Dr Briden).

If progesterone levels are in the low or lower end of the luteal reference range compared to estrogen levels, women may experience symptoms such as PMS, menorrhagia, mastaglia, moodiness, anxiety, and/or insomnia.

The metabolites of progesterone are excreted in urine (not the progesterone itself). When ordering the DUTCH Complete and DUTCH Plus reports, you will see a Progesterone Serum Equivalent on the summary page 1. The urine metabolites of progesterone have been proven to correlate strongly to serum progesterone. The Progesterone Serum Equivalent is most accurate with values in the luteal range and becomes more approximate at very low numbers in the postmenopausal range. Cycling women with very high progesterone metabolites may also decrease the accuracy of the serum equivalent calculation. NOTE: If progesterone is taken orally (also with sublingual), these metabolites are elevated from gut metabolism and results do NOT accurately reflect serum levels.

Progesterone metabolites are low for the luteal phase of the menstrual cycle. It is important to check in with the patient about the timing of the test in relation to menstruation before interpreting this result. If samples were collected too early or too late, the samples may not have been collected during the progesterone peak, which is reflected in the reference range. If the samples were collected about 5-7 days before menses, this likely indicates this cycle was anovulatory, indicating no fertile egg or progesterone production. Consider supporting the HPO axis (brain-to-ovary) communication and chasteberry extract, maca and B6 throughout the month to help with cycle regularity and ovulation.

Estrogen Metabolism

When evaluating estrogen levels, it is important to assess the following:

The status (low, normal or high?) of estrogen production:

Levels of the primary ovarian product, estradiol (the strongest estrogen), as well as "total estrogens" may be considered. For women not on HRT, consider the appropriate range (premenopausal or postmenopausal).

• Phase I Metabolism:

Estrogen is metabolized (primarily by the liver) down three phase I pathways. The 2-OH pathway is considered the safest because of the anti-cancer properties of 2-OH metabolites. Conversely, the 4-OH pathway is considered the most genotoxic as its metabolites can create reactive products that damage DNA. The third pathway, 16-OH creates the most estrogenic of the metabolites (although still considerably less estrogenic than estradiol) - 16-OH-E1. If overall estrogen levels are high, production of 16-OH-E1 may exacerbate high estrogen symptoms. Similarly, a woman with very low levels of estrogens, may have less low estrogen symptoms if 16-OH created to bone loss (a low estrogen symptom). Estriol is thought of as a safer (weaker) estrogen metabolite, but it is important to remember that estriol is actually 16-OH-E2, so generally patients that make a lot of the potentially protective/weak estriol may also make a lot of the estrogenic 16-OH-E1.

When evaluating phase I metabolism, it may be important to look at the ratios of the three metabolites to see which pathways are preferred relative to one another. It may also be important to compare these metabolites to the levels of the parent hormones (E1, E2). If the ratios of the three metabolites are favorable but overall levels

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TEST NAME: DUTCH Sex Hormone Metabolites

of metabolites are much lower than E1 and E2, this may imply sluggish phase I clearance of estrogens, which can contribute to high levels of E1 and E2. Similarly, patients with excessive phase I metabolism may have low E1 and E2 levels because of high rates of clearance (as opposed to simply not making a lot of estrogen). The pie chart will assist you in comparing the three pathway options of phase I metabolism compared to what is "normal." 2-OH metabolism can be increased by using products containing D.I.M. or I-3-C. These compounds are found (or created from) in cruciferous vegetables and are known for promoting this pathway.

Phase I metabolism shows a preference for 2-OH metabolism, which is the protective pathway. The 4-OH and 16-OH pathways may be out of range despite this preference, so careful assessment is necessary. Products to increase 2-OH metabolism may be considered if E1 and E2 are elevated or if the 4-OH or 16-OH estrogens are high relative to 2-OH estrogens. Products that push the 2-OH pathway may also lower E1 and E2 levels, so keep this in mind when considering therapy.

Methylation (part of phase II metabolism) of estrogens:

After phase I metabolism, both 4-OH and 2-OH (not 16-OH) estrogens can be deactivated and eliminated by methylation. The methylation-activity index shows the patient's ratio of 2-Methoxy-E1 / 2-OH-E1 compared to what is expected. Low methylation can be caused by low levels of nutrients needed for methylation and/or genetic abnormalities (COMT, MTHFR). The COMT enzyme responsible for methylation requires magnesium and methyl donors. Deficiencies in folate or vitamin B6 or B12 can cause low levels of methyl donors. MTHFR genetic defects can make it more difficult for patients to make sufficient methyl donors. Genetic defects in COMT can make methylation poor even in the presence of adequate methyl donors.

Androgen Metabolism

Androgen Metabolites: DHEA

DHEA and androstenedione are made almost exclusively by the adrenal gland (although a smaller amount is made in the ovaries for). These hormones appear in urine as DHEA-S (DHEA-Sulfate), androsterone and etiocholanolone.

DHEA peaks for men and women in their 20's and 30's, with a slow decline expected with age. DHEA mainly circulates throughout the body as DHEA-s, with interconversion to active DHEA as it reaches various tissues. DHEA is a weak androgen and will predominately convert to androstenedione, which will then convert to testosterone or estrogen. DHEA-s is made by sulfation, has a much longer half-life than DHEA and largely lacks a diurnal rhythm, which is why it is considered the best way to assess DHEA levels in the body. DHEA-s levels can be affected both by the total production as well as by the body's ability to sulfate DHEA.

The best way to assess the total production of DHEA is to add up these three metabolites. As DHEA production decreases quite significantly with age, we provide the age-dependent ranges. Adrenal DHEA serves as the main source of estrogen, progesterone and testosterone for post-menopausal women.

Androgen Metabolites: Testosterone

The DUTCH test measures the total of testosterone glucuronide and testosterone sulfate. These conjugates of testosterone are formed mostly from bioavailable testosterone that undergoes phase 2 metabolism to make it ready for urine excretion. Females make most of their DHEA in the adrenal gland and a fraction of that DHEA trickles down metabolically to testosterone. Testosterone is also made by the ovaries.

Testosterone glucuronide is mostly made by the UGT2B17 enzyme, which also makes the glucuronide forms of 5a-DHT and 5b-androstanediol. Genetic variants of this enzyme reduce the urinary levels of these hormones without affecting serum levels. The genetic variants of UGT2B17 vary in the population from 7-80% (variation dependent on genetic ancestry, with the highest rates in those of Asian descent). Heterozygous individuals show milder reductions in urinary testosterone than homozygous. For this reason, low and very low levels of urinary testosterone should be confirmed with serum testing before treatment is applied. Serum testing can include free and total testosterone and SHBG.

Testosterone levels may be better understood by also considering its downstream metabolites (5aandrostanediol, 5bandrostanediol). Technically, these metabolites can also be formed from DHEA metabolites without going through the testosterone pathway, but they generally tend to correlate with testosterone production.

Testosterone levels normally decline with age. Age dependent ranges are provided. Perimenopausal testosterone levels can transiently increase before declining again.

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Epi-testosterone (epi-T) is made at about the same rate as testosterone but is not androgenic. In cases where testosterone in urine is low, such as with the UGT2B17 deletion discussed above, epi-T may be used as a proxy for testosterone production, meaning that higher epi-T levels may indicate that a low testosterone result is falsely low. After menopause, epi-T production is less reliable as a marker of testosterone production as epi-T levels drop more sharply than does testosterone during the menopause transition. While epi-T may have limited utility in some cases, it does enhance the picture when taking androgen metabolites together as a whole. Androgens, specifically DHT and testosterone, help to support skin, connective tissue, bone and muscle integrity and promote dopamine conversion in the brain, which can help with mood and libido.

The testosterone level measured is 6.8ng/mg, which is within the overall normal range. If the patient complains of androgen imbalance symptoms, look at the metabolism and DHEA metabolites for further insight. Also, consider other causes. For example, hair loss, which can be androgenic, can also be caused by hypothyroidism, autoimmune disease, high stress or mineral deficiency. Acne, which can be androgenic, also has dietary triggers for some people, most commonly dairy and sugar.

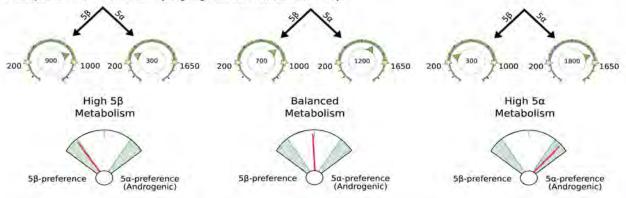
Androgen Metabolites: 5a-reductase versus 5b reductase

5a-reductase converts testosterone into 5a-DHT (DHT), which is even more potent (\sim 3x) than testosterone. High levels of DHT can lead to symptoms associated with too much testosterone, including scalp hair loss, hirsutism, acne and oily skin.

Metabolites created down the 5b-pathway are significantly less androgenic than their 5a counterparts.

The fan-style gauge below the hormones shows the 5a or 5b preference based on etiocholanolone (5b) and androsterone (5a) results. The gauge shows the relative ratio of 5a to 5b products but does not express the absolute value of DHT or if 5a-reductase inhibition is or is not indicated. Consider symptoms and look at the 5a-DHT result if high androgen symptoms are a concern. Progesterone metabolites are also metabolized by 5a and 5b enzymes and the balance between its two metabolites can be useful to confirm a 5a or 5b preference overall (or tissue specific preference).

Example of how to read fan-style gauge for 5a-reductase activity:



When assessing androgens in women, it is important to consider DHEA and testosterone production, 5ametabolism patterns as well as the patient symptoms. For example, a woman with higher levels of DHEA and testosterone will often have high androgen symptoms (facial hair, thinning scalp hair, etc.) exacerbated by 5ametabolism.

If, on the other hand, she prefers 5b-metabolism she may not express high androgen symptoms in spite of higher levels of testosterone because 5b is the less androgenic pathway.

You will also see levels of epi-testosterone, which is not androgenic like testosterone. It happens to be produced in about the same concentrations as testosterone (this is an approximate relationship). This can be helpful when assessing the validity of urinary testosterone testing in an individual patient. If epi-testosterone is much higher

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than testosterone, serum testosterone assessment should considered before initiated therapy for low testosterone. Epi-testosterone is suppressed when exogenous testosterone is given, which can serve as a proxy for assessing endogenous testosterone production which can be obscured by the exogenous hormone administration.

Urine Hormone Testing - General Information

What is actually measured in urine? In blood, most hormones are bound to binding proteins. A small fraction of the total hormone levels are "free" and unbound such that they are active hormones. These free hormones are not found readily in urine except for cortisol and cortisone (because they are much more water soluble than, for example, testosterone). As such, free cortisol and cortisone can be measured in urine and it is this measurement that nearly all urinary cortisol research is based upon. In the DUTCH Adrenal Profile the diurnal patterns of free cortisol and cortisone are measured by LC-MS/MS.

All other hormones measured (cortisol metabolites, DHEA, and all sex hormones) are excreted in urine predominately after the addition of a glucuronide or sulfate group (to increase water solubility for excretion). As an example, Tajic (Natural Sciences, 1968 publication) found that of the testosterone found in urine, 57-80% was testosterone-glucuronide, 14-42% was testosterone-sulfate, and negligible amounts (<1% for most) was free testosterone. The most likely source of free sex hormones in urine is from contamination from hormonal supplements. To eliminate this potential, we remove free hormones from conjugates (our testing can be used even if vaginal hormones have been given). The glucuronides and sulfates are then broken off of the parent hormones, and the measurement is made. These measurements reflect the bioavailable amount of hormone in most cases as it is only the free, nonprotein-bound fraction in blood/tissue that is available for phase II metabolism (glucuronidation and sulfation) and subsequent urine excretion.

Disclaimer: the filter paper used for sample collection is designed for blood collection, so it is technically considered "research only" for urine collection. Its proper use for urine collection has been thoroughly validated.

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PATIENT: TEST NUMBER:

GENDER: Female

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TEST NAME: DUTCH Sex Hormone Metabolites

Reference Range Determination (last updated 11.23.2021)

We aim to make the reference ranges for our DUTCH tests as clinically appropriate and useful as possible. This includes the testing of thousands of healthy individuals and combing through the data to exclude those that are not considered "healthy" or "normal" with respect to a particular hormone. As an example, we only use a premenopausal woman's data for estrogen range determination if the associated progesterone result is within the luteal range (days 19-21 when progesterone should be at its peak). We exclude women on birth control or with any conditions that may be related to estrogen production. Over time the database of results for reference ranges has grown quite large. This has allowed us to refine some of the ranges to optimize for clinical utility. The manner in which a metabolite's range is determined can be different depending on the nature of the metabolite. For example, it would not make clinical sense to tell a patient they are deficient in the carcinogenic estrogen metabolite, 4-OH-E1 therefore the lower range limit for this metabolite is set to zero for both men and women. Modestly elevated testosterone is associated with unwanted symptoms in women more so than in men, so the high range limit is set at the 80th percentile in women and the 90th percentile for men. Note: the 90th percentile is defined as a result higher than 90% (9 out of 10) of a healthy population.

Classic reference ranges for disease determination are usually calculated by determining the average value and adding and subtracting two standard deviations from the average, which defines 95% of the population as being "normal." When testing cortisol, for example, these types of two standard deviation ranges are effective for determining if a patient might have Addison's (very low cortisol) or Cushing's (very high cortisol) Disease. Our ranges are set more tightly to be optimally used for Functional Medicine practices. Below you will find a description of the range for each test:

Female Reference Ranges (Updated 11.23.2021) Low% High% High Low% High% High Low Low 90% 600 2000 20% 90% **b**-Pregnanediol 20% Cortisol A (waking) 10 50 a-Pregnanediol 20% 90% 200 740 Cortisol B (morning) 20% 90% 30 130 Cortisol C (~5pm) Estrone (E1) 20% 80% 12 26 20% 90% 7 30 Estradiol (E2) 20% 80% 1.8 4.5 Cortisol D (bed) 90% 0 0 14 20% 90% 40 Estriol (E3) 20% 80% 18 Cortisone A (waking) 120 5 2-0H-E1 20% 80% 5.1 13.1 Cortisone B (morning) 20% 90% 90 230 4-OH-E1 80% Ø Cortisone C (~Spm) 20% 90% 110 0 1.8 32 16-OH-E1 20% 80% 0.7 2.6 Cortisone D (bed) 0 90% 0 55 2-Methoxy-E1 20% 80% 2,5 Melatonin (6-OHMS) 20% 90% 10 6.5 85 2-OH-E2 0 80% 0 1.2 8-OHdG 0 90% 0 5.2 4-0H-E2 0 80% 0 0.5 Methylmalonate 0 90% 0 2.2 DHEA-S 20% 90% 20 750 Xanthurenate 0 90% 0 1.4 Androsterone 20% 80% 200 1650 0 90% 0 Kynurenate 7.3 Friocholanolone 20% 80% 200 1000 Pyroglutamate 10% 90% 37 60 Testosterone 20% 80% 2.3 14 Homovanillate 10% 95% 4 13 95% 5a-DHT 0 80% 0 6.6 Vanilmandelate 10% 2.4 6.4 5a-Androstanediol 20% 80% 6 30 **Calculated Values** 5h-Androstanediol 20% 80% 20 75 Epi-Testosterone 20% 80% 2.3 14 Total DHEA Production 20% 80% 500 3000 a-THF 80% 20% 90% 75 370 **Total Estrogens** 20% 35 70 **b-THF** 20% 90% 1050 2500 Metabolized Cortisol 20% 90% 2750 6500 1550 **b-THE** 20% 90% 3800 24hr Free Cortisol 20% 90% 65 200 24hr Free Cortisone 20% 90% 220 450 % = population percentile; Example - a high limit of 90% means results higher than 90% of the women tested for the reference

range will be designated as "high."

Provider Notes:

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