

RESULTS ENQUIRIES 13 39 36

24-25952360



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Collect date

HAYNE, LISA Patient

Shop 7, 63-65 Maitland St , Narrabri, NSW, 2390

HAYNE, LISA

GARBUTT, SONIA

21 RIVERSIDE DR NARRABRI NSW 2390

Phone 67933155

D.O.B Age F 15/06/1975 48 years Lab ref

Your ref Collect time 08:41 AM

11/03/2024

Reported 02:02 PM 18/03/2024

Tests requested RT3, STE, UTE, TFT, FE, I12, TAA, LIP, GLU, MBA, DVI

Clinical notes

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Clinical Notes : On Thyroxine last taken 05:50am 11/3/24.

Serum Reverse T3 (RT3) 363 pmol/L (170-539)

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PLASMA TRACE ELEMENTS

(RI)

Zinc (10.0-18.0)12.3 umo1/L Selenium 3.47 umol/L (0.80 - 1.90)

Excess selenium may cause toxicity with common symptoms including diarrhoea, fatigue, hair loss and nail discolouration. A recent Cochrane review (2018) found NO beneficial effect of selenium supplementation in reducing cancer risk in randomised controlled trials of more than 27,000 people.

RI = Reference Interval

Abnormal result/results reported above have been confirmed by repeat assay.

> SURGERY USE

Normal

No Action/File

Patient

Notified

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Further Tests

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URINE TRACE METALS

(RI) **BOEL** Total Volume Random

Urine Creatinine 10.5 mmol/L N/A

Urine Iodine (ug/L) ug/L N/A 242

Urine Iodine (Crt corrected) 186 ug/L (> 100)

WHO Criteria for assessing iodine nutrition based on median urinary iodine concentrations in 6 years or older:

- Severe Iodine deficiency : < 20 ug/L - Moderate Iodine deficiency : 20-49 ug/L 50-99 ug/L - Mild Iodine deficiency : 100-199 ug/L - Adequate Iodine intake : - Iodine intake above requirement : 200-299 ug/L - Excessive Iodine intake : >= 300 ug/L

Urine iodine (creatinine corrected) adjusts the measured result to account for changes in urine concentration. Please note: due to a high variability of an individual's urinary iodine concentration, spot urine iodine concentration is not useful for the diagnosis and treatment of individuals. For more detail refer to:

"https://apps.who.int/iris/bitstream/handle/10665/85972/WHO_NMH_NHD_EP G_13.1_eng.pdf"

BOEL = Biological Occupational Exposure Limit RI = Reference Interval

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THYROID PROFILE

Specimen Type: Serum

TSH 1.5 mIU/L (0.5-4.0)FT4 14 pmol/L (10-20)(3.5-6.5)FT3 pmol/L

Result(s) consistent with euthyroidism.

Please note the above reference intervals have been developed from a non-pregnant healthy general population study.

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IRON STUDIES

Specimen Type: Serum

(10-30)Serum Iron 17 umol/L Transferrin umol/L (32-48)30 Transferrin Saturation 29 % (13-45)Serum Ferritin 34 (30-165)ug/L

Transferrin may be decreased by inflammation (acute or chronic) or protein deficiency or loss.

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SERUM VITAMIN B12

Request Number 25952360 Date Collected 11 Mar 24 Time Collected 08:41

B12 (156-740)pmol/L 418

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THYROID AUTOANTIBODIES

Specimen Type: Serum

HAYNE, LISA

Anti-Thyroglobulin Abs (aTGII) (< 4.5)13 IU/mL Anti-Thyroidal Peroxidase Abs > 1300 IU/ml (< 60)

Over 90% of patients with autoimmune thyroiditis show moderate to high levels of Anti-Thyroidal Peroxidase Abs (anti-TPO) with Anti-Thyroglobulin Abs (anti-Tg) also present in about 90% of such patients. Up to 75% of patients with Graves' hyperthyroidism show increased anti-TPO with anti-Tg present in 50-60%. Low levels of both anti-TPO and anti-Tg may be found in up to 10% of "normal" asymptomatic adults. In most cases of autoimmune thyroid disease increased anti-TPO is the predominant finding although a small proportion of patients show a predominant increase in anti-Tg.

Please note that as of 08/09/2021, Laverty Pathology changed to a reformulated Atellica anti-thyroglobulin antibody (aTGII) assay. The reference interval has been updated. Differences in individual patient results may be observed compared to the previous method. If further information is required please contact a Chemical Pathologist on 9005 7000.

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LIPID STUDIES

Specimen Type: Serum

Reference intervals are included for reference only, and interpretation $\!\!/$ treatment goals should be guided by patient-specific cardiovascular risk assessment (see Australian Cardiovascular Risk Charts. Alternatively, the web-site www.cvdcheck.org.au can be accessed in order to complete a risk assessment for individual patients.)

Haemolysis Nil Nil **Icterus** Nil Lipaemia

Fasting status Random

Total Cholesterol (3.9-5.2)4.9 mmol/L (0.5-1.7)Triglycerides 0.9 mmol/L

NVDPA TARGET LIPID RANGES (MMOL/L) FOR PATIENTS AT HIGH / MODERATE RISK OF CARDIOVASCULAR DISEASE:

TOTAL CHOLESTEROL	<4.0
TRIGS (FASTING)	<2.0
HDL-C	
LDL-C	
NON HDL-C	<2.5
 LDL-C 	 <2.0

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SERUM/PLASMA GLUCOSE

Fasting status Random

Serum 4.9 mmol/L (3.4-7.7)

Normal glucose concentration.

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SERUM CHEMISTRY

SEROPI CHEMISTRI				
Specimen Type: Serum				
Haemolysis	Nil			
Icterus	Nil			
Lipaemia	Nil			
- 1:		7.4	(
Sodium	140	mmol/L	(135-145)	
Potassium	4.1	mmol/L	(3.6-5.4)	
Chloride	102	mmol/L	(95-110)	
Bicarbonate	26	mmol/L	(22-32)	
Anion Gap	16	mmol/L	(10-20)	
Urea	3.2	mmol/L	(2.5-8.0)	
Creatinine	55	umol/L	(45-90)	
eGFR	> 90		mL/min/1.73m^2	
Urate	0.34	mmol/L	(0.14-0.36)	
Bilirubin	19	umol/L	(< 15)	
AST	15	U/L	(< 30)	
ALT	20	U/L	(< 30)	
GGT	20	U/L	(< 35)	
Alkaline Phosphatase	62	U/L	(20-105)	
Protein	72	g/L	(60-82)	
Albumin	46	g/L	(38-50)	
Globulin	26	g/L	(20-39)	
Calcium	2.38	mmol/L	(2.10-2.60)	
Corrected Calcium	2.32	mmol/L	(2.10-2.60)	
Phosphate	1.25	mmol/L	(0.75-1.50)	
· ··oopiiaco	1.20	O 1 / L	(31.3 1100)	

eGFR >=90 mL/min/1.73m2 usually indicates normal kidney function but does not exclude patients with early kidney damage (those with albuminuria, haematuria or abnormal kidney imaging).

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VITAMIN D

Haemolysis Nil

Serum 25(OH) Vitamin D 79 nmol/L

Suggested decision limits for Vitamin D status:

Sufficiency 51 -200 nmol/L Mild deficiency 25 - 50 nmol/L Marked deficiency < 25 nmol/L Toxicity >250 nmol/L

References: Vitamin D and health in adults in Australia and New Zealand:

Position Statement. MJA 2012 June 18; 196(11),686-687.

SURGERY USE

Normal

No Action/File

Patient Notified

Make

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Required Speak with Dr.