

Report to **ELIZABETH JEAN WALKER**
22 TYTHERLEIGH ST
WANNIASSA 2903

Patient **WALKER, ELIZABETH JEAN**
22 TYTHERLEIGH ST
WANNIASSA 2903

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Phone
D.O.B 27/02/53 Age 72 Years Sex F

Ref. by/copy to Ref. by DR.SANJEEWA DON
KAMBAH MED CTR
KAMBAH, 2902
Phone: 0262316068
Collect date 23/04/25 Lab ref **25-11247648**
Collect time 08:19 Your ref 00393557
Printed 24/04/25

Tests requested Completed report. Thank you for your referral.
Completed - CHEM,LIP,IRON,GLU

Clinical notes

LIPID STUDIES

Request Number	24579396	10707780	11247648
Date Collected	7 Aug 23	28 Oct 24	23 Apr 25
Time Collected	10:49	08:55	08:19
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	Nil
Icterus	1+	Nil	Nil
Lipaemia	Nil	Nil	Nil

Fasting status	Fasting	Fasting	Fasting
Chol (3.9-5.2) mmol/L	4.8	5.7*	7.2*
Trig (0.5-1.7) mmol/L	0.7	1.0	0.8
HDL (1.0-2.0) mmol/L	1.5	1.5	1.7
LDL (1.5-3.4) mmol/L	3.0	3.7*	5.1*
Non-HDL (< 3.4) mmol/L	3.3	4.2*	5.5*
Chol/HDL(< 4.5)	3.2		4.2

*Went off
ezetimibe
late March*

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	>= 1.0
LDL-C	<2.0
NON HDL-C	<2.5

Increased LDL-C can be seen in hypothyroidism and nephrotic syndrome. Primary causes include polygenic hypercholesterolaemia, Familial Hypercholesterolaemia, Familial Defective ApoB-100 and Familial Combined Hyperlipoproteinaemia. Increased LDL-C is a risk factor for CVD. Assess overall risk and consider treatment.

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SERUM CHEMISTRY

Request Number	24579396	10707780	11247648
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Specimen Type: Serum			

Haemolysis	Nil	Nil	Nil
Icterus	1+	Nil	Nil
Lipaemia	Nil	Nil	Nil

Na	(135-145)	mmol/L	144	143	141
K	(3.6-5.4)	mmol/L	4.9	5.0	4.4
Cl	(95-110)	mmol/L	104	107	105
HCO3	(22-32)	mmol/L	24	28	29
An Gap	(10-20)	mmol/L	21*	13	11
Urea	(3.0-10.0)	mmol/L	7.1	4.8	5.6
Creat	(45-90)	umol/L	60	60	65
eGFR	mL/min/1.73sqM		89	88	80
Urate	(0.14-0.36)	mmol/L		0.33	
Bili	(< 15)	umol/L	30*	24*	22*
AST	(< 35)	U/L	27	30	27
ALT	(< 30)	U/L	20	22	18
GGT	(< 35)	U/L	< 7	9	9
Alk Phos	(30-115)	U/L	75	67	70
Protein	(60-82)	g/L	69	63	64
Albumin	(36-48)	g/L	44	43	43
Glob	(20-39)	g/L	25	20	21
Ca	(2.10-2.60)	mmol/L		2.40	
Corr Ca	(2.10-2.60)			2.40	
PO4	(0.75-1.50)	mmol/L		1.20	

eGFR values between 60 and 89 mL/min/1.73m2 should be interpreted with caution. These results are only consistent with CKD in the presence of other evidence such as microalbuminuria, proteinuria or haematuria.
Ref:Lamb EJ etal in Ann Clin Biochem 2005; 42:321-345.

SERUM/PLASMA GLUCOSE

Request Number	24579396	10707780	11247648	
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Time Collected	10:49	08:55	08:19	
Fasting status	Fasting	Fasting	Fasting	
Serum	(3.4-5.4) mmol/L	4.2	5.1	4.9

Normal glucose concentration.

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IRON STUDIES

Specimen Type: Serum
Serum Ferritin 146 ug/L (30-400)

No evidence of iron deficiency.

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Lipoprotein X may be seen in patients with cholestatic liver disease and lecithin-cholesterol acyl transferase deficiency.

Familial hypercholesterolaemia, an autosomal dominant cause of premature cardiovascular disease is likely (risk approximately 1 in 5). Recommend <http://www.athero.org.au/calculator> to calculate diagnostic score and <http://www.athero.org.au/physicians/what-is-familial-hypercholesterolaemia-fh> to assist with diagnosis and management, if required.

