

DAHLSTROM, VERA

For Surgery Use ☐ Urgent ☐ Ring Patient ☐ Make Appointment ☐ Note in Chart ☐ File ☐

Patient	DALEY, BRIGITTE	RPN 20 OAK ST MILLAA MILLAA QLD 4886	Requested	05/10/2023
Sex	F	Age 56 years	DOB	14/12/1966
Report For	DAHLSTROM, VERA		Collected	05/10/2023 10:45 AM
Ref. by/copy to	DAHLSTROM, VERA		Reported	06/10/2023 03:30 PM

Serum Zinc 10 umol/L (10-25)

Date 05/10/23
Time 10:45
Lab No 69985653

Thyroglobulin AbII 38 IU/mL (< 4.6)
Thy. Peroxidase Ab 180 IU/mL (< 60)

Please note that as of 06/9/2021, QML Pathology changed to a reformulated Atellica Thyroglobulin Antibody (TgAbII) 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 (07) 3121 4444.

CUMULATIVE SERUM BIOCHEMISTRY

Date	05/10/23
Time	10:45
Lab No	69985653
	FASTING
Sodium	140 mmol/L (137-147)
Potass.	4.4 mmol/L (3.5-5.0)
Chloride	108 mmol/L (96-109)
Bicarb	23 mmol/L (25-33)

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An.Gap	13.4	13	mmol/L (4-17)
Gluc		5.3	mmol/L (3.0-6.0)
Urea		5.3	mmol/L (2.5-7.5)
Creat		57	umol/L (50-120)
eGFR		> 90	mL/min (over 59)
Urate		0.38	mmol/L (0.14-0.35)
T.Bili		25	umol/L (2-20)
D.Bili		6	umol/L (0-8)
Alk.P		56	U/L (30-115)
GGT		16	U/L (0-45)
ALT		25	U/L (0-45)
AST		18	U/L (0-41)
LD		185	U/L (80-250)
Calcium		2.35	mmol/L (2.15-2.60)
Corr.Ca		2.31	mmol/L (2.15-2.60)
Phos		1.0	mmol/L (0.8-1.5)
T.Prot		74	g/L (60-82)
Alb		44	g/L (35-50)
Glob		30	g/L (20-40)
Chol		5.1	mmol/L (3.9-7.4)
Trig		1.8	mmol/L (0.3-2.2)
Lab No		69985653	
Date		05/10/23	

Pathology Report

CUMULATIVE SERUM HOMOCYSTEINE

Date 05/10/23
Time 10:45
Lab No 69985653

Homocysteine 10.7 umol/L (0.0-15.0)

69985653 High normal value.

With this level, the heterozygous state for defects of transsulphuration (homocysteinaemia) is unlikely. However the risk of coronary artery disease may be mildly elevated over the baseline. This is independent of other risk factors.

Homocysteine Related Risk

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Plasma level (umol/L)	Risk Average
Below 9.0	No increase
9.0 - 14.9	x 2
15.0 - 19.9	x 3
20.0 or greater	x 4.5

Risks approximated from New Eng J Med 1997 (337:230-236)

Serum Copper 14 umol/L (10-45)

CUMULATIVE SERUM COMPLEMENT AND C-REACTIVE PROTEIN (CRP)

Date	05/10/23
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Lab No	69985653

CRP < 5 mg/L(0-6)

C-reactive protein (CRP) is a non-specific indicator of tissue damage. The level rises rapidly (within 6-10 hours) after tissue injury, peaks at 48-72 hours and returns to normal within a few days. Common causes of markedly increased CRP include infection (particularly bacterial), trauma, surgery, myocardial infarction, many malignancies and inflammatory disorders.

CUMULATIVE IRON STUDIES

Date	05/10/23
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Iron	23	umol/L	(10-33)
TIBC	68	umol/L	(45-70)
Saturation	34	%	(16-50)
Ferritin	130	ug/L	(30-320)

CUMULATIVE SERUM VITAMIN D

Date 05/10/23
Time 10:45
Lab No 69985653
Vitamin D3 57 nmol/L (> 49)

69985653 We occasionally see evidence of functional vitamin deficiency and possibly accelerated bone loss with levels in the range of 50-70 nmol/L.

CUMULATIVE FULL BLOOD EXAMINATION

Date 05/10/23
Time 10:45
Lab No 69985653

Hb	139	g/L	(115-160)
RCC	4.2	x10 ¹² /L	(3.6-5.2)
Hct	0.42		(0.33-0.46)
MCV	101	fL	(80-98)
MCH	34	pg	(27-35)
Plats	471	x10 ⁹ /L	(150-450)
WCC	7.4	x10 ⁹ /L	(4.0-11.0)
Neuts	51 %	3.8 x10 ⁹ /L	(2.0-7.5)
Lymphs	37 %	2.7 x10 ⁹ /L	(1.1-4.0)
Monos	9 %	0.7 x10 ⁹ /L	(0.2-1.0)

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Eos	2 %	0.15	x10 ⁹ /L	(0.04-0.40)
Basos	1 %	0.07	x10 ⁹ /L	(< 0.21)

69985653 Automated Comment:

As per ISLH guidelines - Film not reviewed. If a film review is truly indicated, contact the laboratory within 24 hours of collection. Otherwise investigate any highlighted abnormalities as clinically appropriate.

Borderline Red Cell Macrocytosis: Recommend a review of haematinic profile (Iron/B12/Folate). Correlate Clinically.

Thrombocytosis may be seen in hyposplenic states, or occur as response to infection, inflammation, post-surgery/trauma, malignancy, bleeding, or in older patients an early Myeloproliferative Neoplasm (MPN). Correlate with E+LFTs, ESR/CRP, Iron Studies. If a MPN is a possibility, JAK2 molecular analysis is recommended.

** FINAL REPORT - Please destroy previous report **

Pathology Report