

# Elevated anti-Thyroglobulin Antibodies

<b>HOSP #</b>		<b>WARD</b>	Oudtshoorn Clinic
<b>CONSULTANT</b>	George van der Watt & David Marais	<b>DOB/AGE</b>	66y Male

## Abnormal Result

The screenshot shows a medical validation window titled "Medical Validation : XJ (Authorise By Episode)". It contains patient information, test details, and a table of results.

Test Set	Staff Notes	Test Item	Result	Units	Normal Values	Previous Result 1	Previous Result 2	Previous Result 3	Previous Result 4	Previous Result 5
ATGA		Anti-thyroglobulin Ab	1,944	U/mL	<115					
		ATGA clinician alert								

## Presenting Complaint

Mr. X, a 66 year old male, complained of chest pain, was seen at the Oudtshoorn Emergency department and a myocardial infarction was excluded by three serial point-of-care (POC) Troponin I results.

## History

- Known with hypothyroidism, but the cause was not defined yet.
- On Eltroxin 150 ug daily PO
- No other treatment.
- Various stool analyses had been sent in for culture, with no definitive result.

# Examination

Unfortunately not known.

# Laboratory Investigations

Free T4: 24.6 pmol/L (7.6 – 16.1 pmol/L)

Anti-Thyroglobulin Antibody levels were elevated at **1944 U/mL** (ref. <115 U/mL).

# Other Investigations

Later, by retrospective viewing of the patient's results it was revealed:

Total Cholesterol (TC) was elevated at 7.6 mmol/L. Hypothyroidism is associated with hypercholesterolemia. It can be concluded by the retrospective overview of results that upon an episode of hypothyroidism, the patient had hypercholesterolemia. This was most likely due to cessation of Thyroxine treatment, to whatever reason.

Test Item	15/04/2019 17:38	11/01/2019 17:44	31/10/2018 18:32	31/10/2018 00:24	30/10/2018 17:21	02/10/2018 15:03	03/09/2018 15:13	31/08/2018 18:19	23/02/2018 14:30
Comments									
Total chol	7,55							3,78	
Comment	CHOLC2							CHOLC2	
CRP								1	
Total PSA									
CEA									
Comment									
TSH	<b>δ+25,53 H</b>	<b>δ+ 1,34</b>			<.01 L			<.01 L	<.01 L
Free T4	<b>δ- &lt;3.2 L</b>	<b>δ- 8,9</b>			<b>δ- 15,9</b>	<b>δ- 24,6 H</b>	<b>34,3 H</b>		<b>27,6 H</b>
Free T3							<b>8,9 H</b>		
Anti-thyroglob Ab						<b>1944 H</b>			

Index sample marked by the yellow shade. TC result which is raised (upper left corner) corresponds to the severely hypothyroid episode as revealed by the low T4 on that same sample.

Investigations also confirmatory for auto-immune

hypothyroidism are:

- Anti-Thyroid peroxidase antibodies
- Anti-TSH receptor antibodies

## Final Diagnosis

Auto-immune hypothyroidism

## Take Home Messages

*Interestingly, numerous patients with hypothyroidism is diagnosed at our Lipid Clinic at Groote Schuur Hospital. Patients are being referred for hypercholesterolaemia. Generally referral to this clinic happens when TC > 7.5 mmol/L. These patients are referred as presumed to have familial hypercholesterolaemia, but upon further work-up it is found that many of these patients have long-standing untreated hypothyroidism.*

Prevalences of antithyroid antibodies as summarized by Up-to-date:



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## GFR by Iohexol

M.Med. discussion by Dieter van der Westhuizen

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# Thyroid Carcinoma

A case of Thyroid carcinoma

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## Hyperaldosteronism

<b>HOSP #</b>	<b>WARD</b>	Murraysburg Hospital, Female Ward
<b>CONSULTANT</b>	<b>DOB/AGE</b>	51 y female

## Abnormal Result

Aldosterone: 1380 pmol/L

Renin: 2.1 ng/L

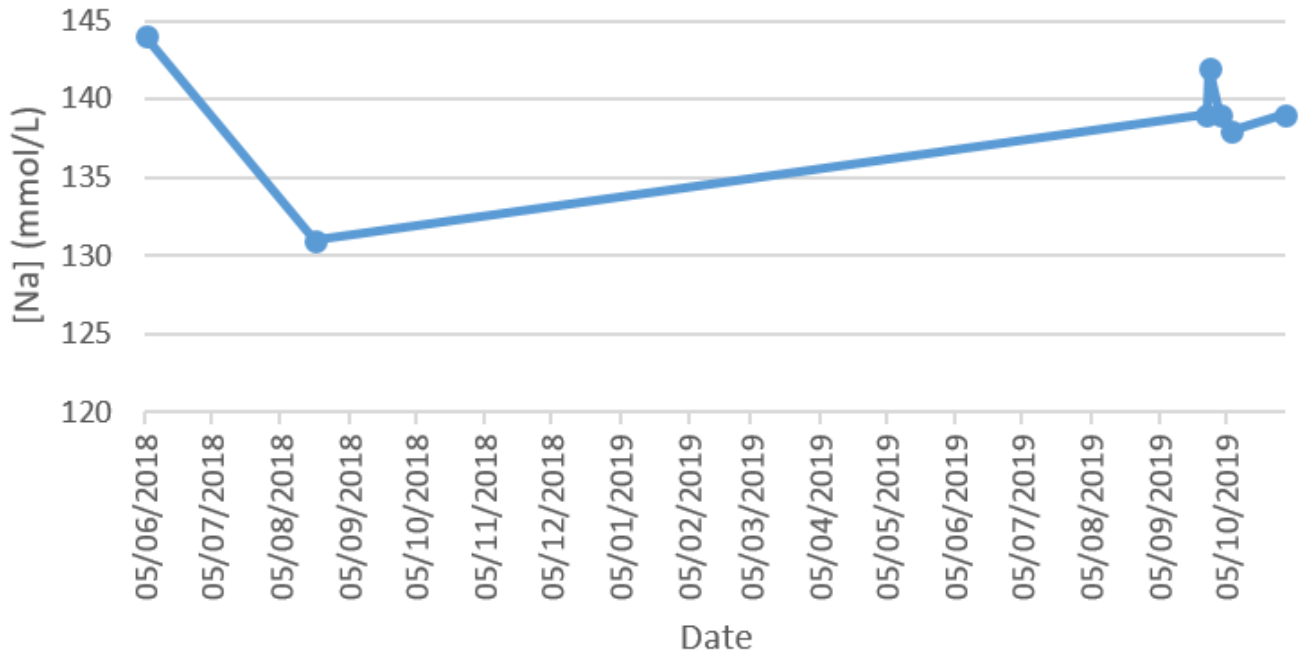
Aldosterone: Renin ratio: 657.14 pmol/ng

## Presenting Complaint

Uncontrolled Hypertension, unresolved on maximum dose of 3 antihypertensives.

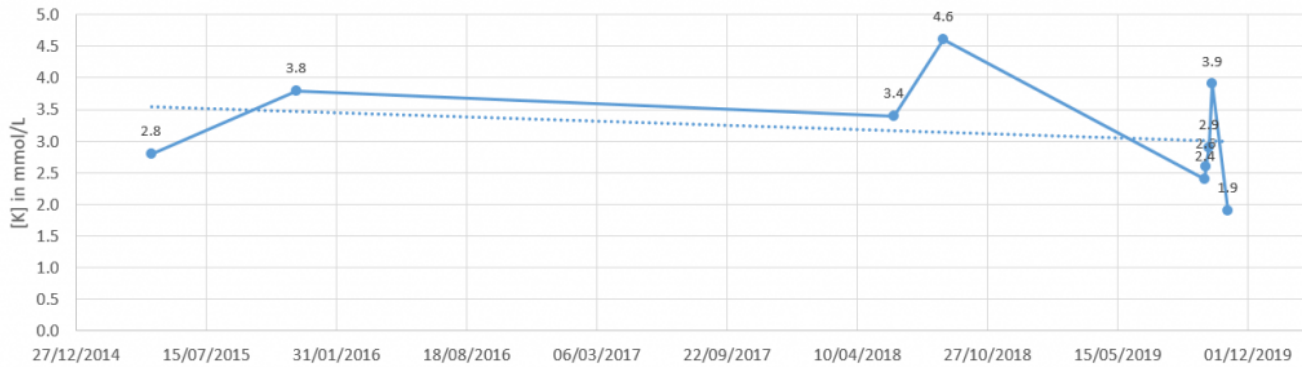
## History

## Sodium over time

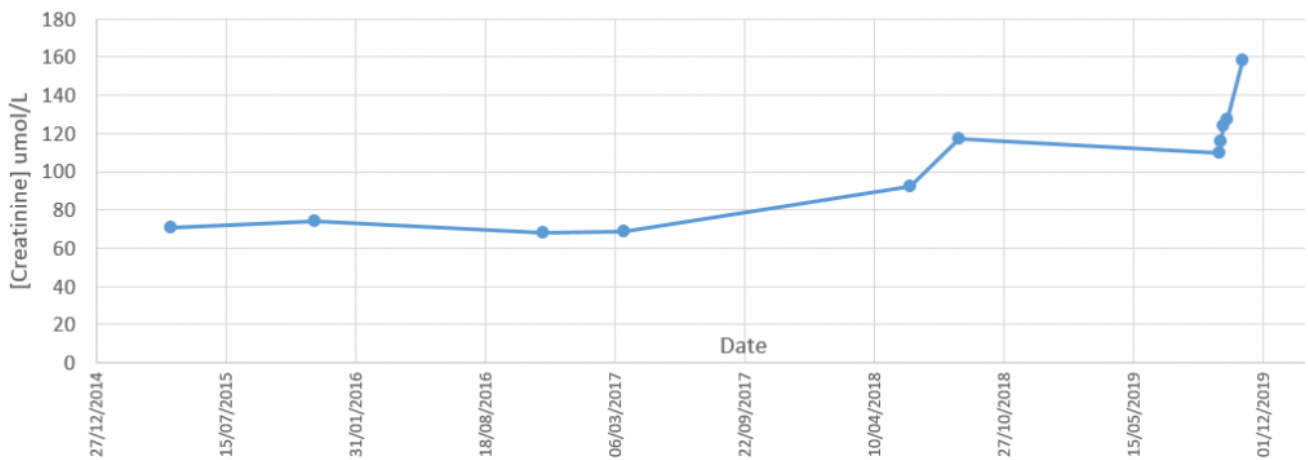


	31/10/2019	07/10/2019	02/10/2019	27/09/2019	26/09/2019	20/08/2018	05/06/2018	30/11/2015	21/04/2015
K	1.9	3.9	2.9	2.6	2.4	4.6	3.4	3.8	2.8

## Potassium plotted over time



## Creatinine over time





26/08/2018									
26/09/2019	139	2,4	45			0.81			
27/09/2019	142	2,6	43						
01/10/2019									
02/10/2019	139	2,9	40			CEGK			
03/10/2019									
07/10/2019	138	3,9	38						
31/10/2019	139	1,9	30	28					

## Urine metanephrines

Urine collection period	24 h	Reference value
Urine volume	3080 ml	
Ucreat	2,2 mmol/L	
Umetadren	160 nmol/L	
Unormetadren	870 nmol/L	
dUmetadren	493 nmol/24h	152-913
dUnormetadren	2680 nmol/24h	699-2643
Umetadren:cr	73 nmol/mmol creat	17-91
Unormetad:cr	395 nmol/mmol creat	75-309

## Final Diagnosis

Primary hyperaldosteronism causing secondary hypertension with accompanying renal injury.

## Take Home Messages

### Reference Ranges for Aldosterone:

- Upright 70 – 1066 pmol/L
- Supine 49 – 643 pmol/L

Screening for primary hyperaldosteronism: most sensitive when

>350 pmol/L

## Reference Ranges for Renin:

- Upright: 2.7 – 27.7 ng/L
- Supine: 1.7 – 23.9 ng/L

Beta-blockers suppress renin levels and should be stopped 2 weeks before testing.

## Aldosterone: Renin Ratio:

Most sensitive when the ratio is >118 pmol/ng.

## Effects of hyperaldosteronism

- One's expectation is a high serum sodium, but since it normalizes with an increase in fluid volume, hence hypertension as in this case, there is normal sodium.
- Low serum potassium due to loss in urine, although this can also be normal.
- Increased urine potassium concentration (>30 mmol/L) in a random urine specimen suggests increased mineralocorticoid effect.
- The renin:aldosterone ratio is used to compensate for the increase in aldosterone which is caused by an increase in renin (for instance which is caused by hypovolemia or low blood pressure).
- Some studies recently published are suggesting that the prevalence of hyperaldosteronism are significantly more than was (and is) thought, and hence urinary (24 hour) aldosterone measurement may be more accurate to screen for hyperaldosteronism. The authors of recent estimates of the prevalence of hyperaldosteronism are of opinion that hyperaldosteronism may be the cause of around 10% of unexplained "essential" hypertensives (see attached articles).

[Hyperaldo-prevalence-2020Download](#)

[Primary-hyperaldo-Editorial-2020Download](#)

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# **Drip line contamination – Ringers Lactate**

A case of drip line contamination.

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# **Albumin Assay – Bromocresol Green method**

Practical assay for albumin measurement

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# **Total Protein assay – Bradford**

A practical experiment to illustrate the measurement of total protein in serum using the Bradford assay

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# Hypernatremia

Case of a child with extreme hypernatremia.

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## Prolactin

HOSP #	WARD	ENT Clinic
CONSULTANT	DOB/AGE	35 Y Male

## Abnormal Result

Prolactin 10 986.0 ug/L (4-15.2)

Dilutions:

1/10 >4700;

1/100 = 10821;

1/50 = 10 986.

## Presenting Complaint

Epistaxis

## History

Patient with epistaxis referred to the ENT specialist clinic.  
No relevant medication history.

# Examination

35 y male with a large left post-nasal space mass, a vascular mass involving the pituitary fossa.

?NBL (non-benign lesion)

?Sinonasal malignancy

?Pituitary Tumour

# Laboratory Investigations

TSH 0.91 pmol/L (0.27-4.20)

Free T4 15.7 pmol/L (12-22)

FSH 0.8 IU/L ↓ (1.5-12.4)

LH 0.2 IU/L ↓ (1.7-8.6)

Testosterone 0.2 nmol/L ↓ (8.6-29.0)

PTH 1.7 pmol/L (1.6-6.9)

Prolactin measuring method:

The Elecsys prolactin sandwich immunoassay uses two monoclonal antibodies directed against human prolactin.

R1 = biotinylated antibody – recognizes the N-terminal end of the prolactin molecule

R2 – ruthenium complexed antibody probably reacts with a region in the middle of the prolactin molecule.

1<sup>st</sup> incubation: a biotinylated monoclonal prolactin-specific antibody and a monoclonal prolactin-specific antibody labeled

with a ruthenium complex form a sandwich complex.

2<sup>nd</sup> incubation: after addition of streptavidin-coated microparticles, the complex becomes bound to the solid phase via interaction of biotin and streptavidin.

Reaction mixture aspirated into the measuring cell where microparticles are magnetically captured into the surface of the electrode. Unbound substances are then removed with ProCell.

Application of a voltage to the electrode then induces chemiluminescent emission which is measured by a photomultiplier, results calculated by a standard curve.

## **Other Investigations**

Monomeric prolactin – 7744 ug/L (70% recovery after PEG precipitation)

Biopsy: confirmed tumour stained strongly positive with prolactin suggesting a prolactinoma.

## **Final Diagnosis**

Pituitary Macroprolactinoma

## **Take Home Messages**

Sandwich immunoassays are prone to high dose hook-effect. There are various ways to overcome this effect. (This will later be expanded on – see AFP

/ Beta-HCG).

Prolactin appears in the serum as:

1. Active monomeric prolactin ("little") (80%) 23kDa
2. Inactive dimeric prolactin ("big") (5-20%) 50-60kDa
3. Low activity tetrameric prolactin ("big-big") (0.5-5%) 150-170kDa

Precipitation by PEG yields the active monomeric prolactin, expressed as a percentage recovery after precipitation. Big-big prolactin consists of an antigen-antibody complex of monomeric prolactin-immunoglobulin G and is defined as macroprolactin. This has a long half-life in blood when compared to normal prolactin and gives false high readings of prolactin, leading to unnecessary investigations in certain cases. A high prolactin should thus be confirmed by doing a PEG precipitation.

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## **Fluid Triglycerides**

A case of high fluid triglycerides

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# ACTH

<b>HOSP #</b>		<b>WARD</b>	G16 Medical Ward
<b>CONSULTANT</b>		<b>DOB/AGE</b>	54 y Female

## Abnormal Result

21/08/2018 Two ACTH tests (referred to another laboratory) and two

Cortisol levels (at our laboratory) were done.

At first it was thought to be a dexamethasone suppression test, but then

realized the clinicians were suspecting hypopituitarism.

10h05: **ACTH 0.7 pmol/L ↓** (1.6-13.9) Cortisol 8 nmol/L ↓  
(Morning: 133- 537; Afternoon 68 – 327)

10h35: ACTH 1.8 pmol/L N (1.6-13.9)

Cortisol 68 nmol/L ↓ (Morning: 133- 537; Afternoon 68 – 327)

## Presenting Complaint

? hypopituitarism

## History

Known with a pituitary macroadenoma, previously seen at the Radiotherapy clinic in 2016.

## Examination

No clinical info available.

For Primary adrenal insufficiency one would expect:

Hyperpigmentation

(due to ↑ ACTH), +/- hyperkalemia/hyponatremia (aldosterone

effect), +/-  
virilization.

For Secondary adrenal insufficiency there is subtle symptoms, electrolytes are not deranged significantly because aldosterone function is preserved. See table on Bishop 7<sup>th</sup> ed. p. 459.

## Laboratory Investigations

Measurement of plasma ACTH concentration is used to assess Cushing's disease, adrenal tumors, ectopic ACTH-producing tumors, Addison's disease, Nelson's syndrome, and hypopituitarism.

The laboratory diagnosis of hypopituitarism, however is relatively straightforward. In contrast to the primary failure of an endocrine gland that is accompanied by dramatic increases in circulating levels of the corresponding pituitary tropic hormone, secondary failure (hypopituitarism) is associated with low or normal levels of tropic hormone. This is the diagnosis in this case with the history of previous radiotherapy which was given for a macro-adenoma.

## Other Investigations

Free T4 on 19/04/2018 was 7.8 pmol/L (12-22), also suggesting possible hypopituitarism, although a TSH would be helpful.

# Final Diagnosis

Hypopituitarism confirmed.

## Take Home Messages

Dexamethasone suppression test need only measurement of cortisol, not accompanying ACTH, except in extended work-up however, where a Cosyntropin (CRH) stimulation test can be done to distinguish between pituitary or hypothalamic insufficiency.

Evaluation of pituitary function need the Primary hormone (Cortisol) as well as the tropic hormones from the pituitary (ACTH).