

# PTH Summary

## PTH Summary

PTH (Intact) = active

1 84

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Heterogeneity

Fragments / Truncated

34 84

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37 84

1<sup>st</sup> Generation PTH assays

-Competitive

2<sup>nd</sup> Generation PTH assays

-Sandwich (detection against 84' end)

3<sup>rd</sup> Generation PTH assays

-Sandwich (detection against 84' end and 1-6' AA end)

Dr. Jody Rusch's simplistic explanation of PTH and its measurement

n-truncated are deficient in the first few aa's (7-84) – hence biologically inactive.

1-11 aa sequence is necessary for function.

Different assays:

- Intact PTH measurements: 7-84 and 1-84
- Bioactive PTH: 1-84
- CAP assay: cAMP inducible PTH. This assay determines the biologically active PTH by its ability to induce cAMP.
- PTHrP assays and PTH assays are exclusive to each other by design.

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# A Case of Neurodevelopmental Delay

<b>HOSP #</b>		<b>WARD</b>	Neurodevelopmental clinic – Inkosi Albert Luthuli Hospital
<b>CONSULTANT</b>	Prof. George van der Watt	<b>DOB/AGE</b>	2y male

## Abnormal Result

Urine organic acid analysis was performed upon which a big peak was seen, representative of phenylpyruvate.

## Presenting Complaint

The patient was a 2 year old male evaluated at a neurology clinic for neurodevelopmental delay.

## History

The patient's brother died at 3 or 4 years of age with similar neurodevelopmental delay.

## Examination

Unfortunately this information was unavailable. The clinician I got hold of at Inkosi Albert Luthuli hasn't seen the patient himself.

# Laboratory Investigations

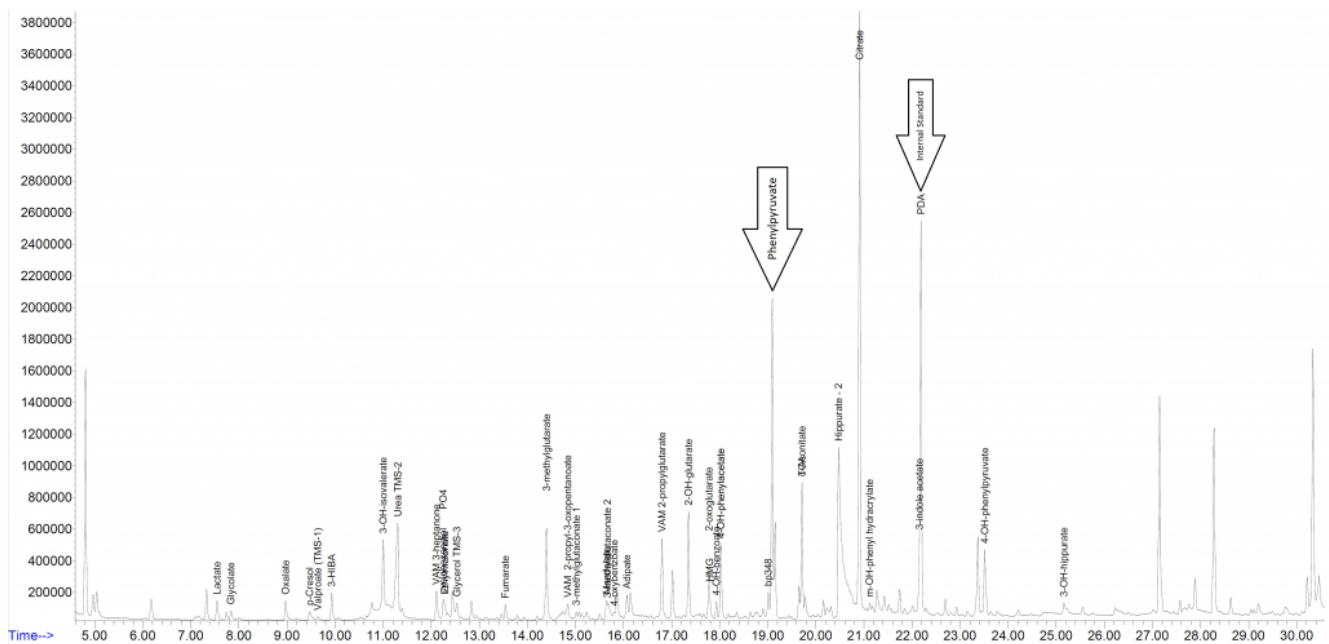


Fig 1 – Urine organic acid screening by GCMS demonstrates elevations of the phenylketones: phenylpyruvate and 4-OH phenylpyruvate. These findings are indicative of a diagnosis of phenylketonuria due to autosomal recessive deficiency of phenylalanine hydroxylase.

## Other Investigations

The urine amino acid analysis yielded a significantly raised phenylalanine: 672  $\mu\text{mol/L}$  (ref <67)

## Final Diagnosis

This is a case of phenylketonuria

The diagnosis is also supported by a plasma phenylalanine of 672  $\mu\text{mol/L}$  (ref < 67).

## Take Home Messages

Build-up of phenylalanine gets metabolised to phenylpyruvate (which is seen in urine at high levels).

Phenylalanine levels >600 umol/L in serum is highly indicative of phenylketonuria

Prof. George van der Watt

Biopterin cycling defects usually cause levels >125 umol/L.

This deficiency is 4-monooxygenase deficiency.

Management of PKU is with a phenylalanine restricted diet.

## 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 for a patient. The patient's name is Mr. X, and the MRN is MRN00100141. The patient is 66 years old, male, and was born on 22/04/1952. The test set is ATGA (Anti-thyroglobulin Ab), and the result is 1,944 U/mL, which is significantly above the normal value of <115 U/mL. The test was performed on 02/10/2018 at 11:55. The patient is currently at Oudtshoorn Clinic, ward not stated, and the doctor is SR BRINKHUIS.

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
Comment									
Total chol	7,55							3,78	
Comment	CHOLC2							CHOLC2	
CRP								1	
Total PSA									
CEA									
Comment									
TSH	δ+25,53 H	δ+ 1,34			<.01 L			<.01 L	<.01 L
Free T4	δ- <3.2 L	δ- 8,9			δ- 15,9	δ- 24,6 H	34,3 H		27,6 H
Free T3							8,9 H		
Anti-thyrogl Ab						1944 H			

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:

