



For Further Patient Information please contact Midland Public and Private Hospitals, 1 Clayton St, MIDLAND 6056, WA (08) 9462 4000

Nominated GP

Patient

IHI DOB

Sex

Phone

Address

ALDO RIQUELME Phone 08 9374 7000 Name MIDLAND GP SUPERCLINIC 6 08 9374 7099 Address Fax

60 BALFOUR ROAD, SWAN VIEW, WA, 6056

NATAYA NAKHONWONG

18 Oct 1986 (38yr)

0420665650

CENTENNIAL PL, RAILWAY WORKSHOP, MIDLAND, WA, 605

F

**Admission Details** 

Admit Date 30 Oct 2024 Disch Date

7 Days

AMO

DR REBECCA LOUISE DUGMORE

LOS AMO Ph

Patient

**General Medicine** 

Speciality Ward

3D - Medical

Disch To

Home

Name

Address 60 BALFOUR ROAD, SWAN VIEW, WA, 6056, Australia

Problems/Diagnoses Relevant To This Visit

Principal Diagnosis

Guillain-Barre syndrome

Problem/Complication

Progress/Summary

THERE WAS NO UNPLANNED RETURN TO SURGERY

\* indicates pirmary procedure

Start Date

MRN 222350

Mob 0420665650

**Clinical Synopsis** 

Presenting Problem

Guillain-Barre syndrome

Past Medical History

Eosinophilic lung disease

Chronic rhinosinusitis

Progress in Hospital / Summary of Stay

38 year old female presenting with bilateral lower limb and upper limb weakness.

On Sunday, noted bilateral lower limb weakness after work with numbness then on Monday noticed upper limb weakness with tingling sensation

Not settling, yesterday at work was not able to get up of chair, felt very lightheaded

Today was at work for 1 hour, felt very weak and passed out, was found by friend and asked brought to ED

Had viral illness a week ago, currently resolved

Has history of sinusitis

Denies fever/chills/rigors, abdo pain, nausea, vomiting, visual disturbances

Normal menstrual cycle, no history of menorrhagia

Lumbar Puncture: WCC: 4\*10^6/L (predominantly lymphocytes); protein 0.5 (predominantly antibodies)

Admitted under General Medicine (Dr Dugmore, Consultant)

#### Issues.

- 1. Ascending Weakness
- Neurology input
- Treated with IVIG
- MRI brain and spine:

Comment: Post-contrast sequences have been acquired supplement the MRI examination acquired yesterday, 30

October 2024. There is no pathological spinal cord, cauda

equina or leptomeningeal enhancement. There are no MRI

features to suggest Guillain-Barre syndrome.

- Allied health input
- Lower limb weakness improved
- Cleared by allied health, neuro, medically safe for discharge

- 1. Follow-up outpatient EMG studies in December (you will be provided with an appointment time)
- 2. Neurology follow-up in January (you will be provided with an appointment time)
- 3. If you experience a relapse in muscle weakness, difficulty breathing, or are otherwise concerned, please seek medical attention

We wish you the best with your recovery



NATAYA NAKHONWONG Patient

8003 6045 7347 7220 IHI

DOB 18 Oct 1986 (38yr) MRN 222350

Sex

Phone 0420665650 Mob 0420665650 Address 60 BALFOUR ROAD, SWAN VIEW, WA, 6056

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Alerts, Allergies, and Adverse Drug Reactions Type

Description Comments Discharge Summary Finalised Discharge

Start Date 06 Nov 2024

Summary Finalised

Alerts, Allergies, and Adverse Drug Reactions Description

Comments Start Date

Allergy Status Unknown

**Discharge Medications** 

Alert

Type

Drug Name, Form, Dose, Frequency, Quantity, Status, Change Reason, Reason For

1 Benralizumab, Unchanged, Subcut every 8 weeks

2 Budesonide/ Formeterol, 200/6 microg, As required, Unchanged

Able to Self Medicate

YES

Medications changed since admission

NO

Relevant Results

(For additional pathology information contact Australian Clinical Labs 1300 367 674) #22101589 : Red cell parameters suggest iron deficiency (chronic blood loss), impaired iron metabolis m (chronic disease) or thalassaemia.

Suggest iron studies, and if patient does not have a known history of thalassaemia, haemoglobinopathy screen. HAEMATOLOGY SPECIMEN: WHOLE BLOOD

Date: Coll. Time: Lab Number:	<b>06/11/24</b> 05/11/24 09:08 08:09 #22101589 22931453	08:46 result only)
HAEMOGLOBIN RBC HCT MCV MCH MCHC RDW WCC Neutrophils Lymphocytes Monocytes Eosinophils	124 121 5.07 4.99 0.38 0.37 * 75 * 75 * 24.5 * 24.2 326 327 13.4 13.6 5.5 6.1 2.9 3.2 1.9 2.4 0.6 0.5 0.0 0.0	123 (115 - 165) g/L 4.97 (3.80 - 5.50)×10 \\S\\12 /L 0.37 (0.35 - 0.47)  * 75 (80 - 99) fL  * 24.7 (27.0 - 34.0)pg 332 (310 - 360) g/L 13.4 (11.0 - 15.0)% 6.0 (4.0 - 11.0) ×10 \\S\\9 /L 3.7 (2.0 - 8.0) ×10 \\S\\9 /L 1.9 (1.0 - 4.0) ×10 \\S\\9 /L 0.4 (< 1.1) ×10 \\S\\9 /L
Basophils PLATELETS MPV	<ul> <li>0.1</li> <li>338</li> <li>9.5</li> <li>9.9</li> </ul>	< 0.1 (< 0.3) x10 \\S\\9 /L

#22101589 : Red cell parameters suggest iron deficiency (chronic blood loss), impaired iron metabolism (chronic disease) or thalassaemia. Suggest iron studies, and if patient does not have a known history of thalassaemia, haemoglobinopathy screen.

FBE-C ECU-W

This request has other tests in progress at the time of reporting Inpatient. Specialist management noted. GENERAL CHEMISTRY **SERUM** 

SPECIMEN:

Date: Coll. Time: Lab Number:	<b>06/11/24</b> 09:08 22101589	05/11/24 08:09 22931453	04/11/24 08:46 22931848		
Sodium Potassium Chloride Bicarbonate Anion Gap Urea Creatinine	135 3.8 101 26 12 4.6 65	* 134 3.8 100 28 10 3.4 62	136 3.9 102 26 12 3.7 64	(135 - 145) (3.5 - 5.2) (95 - 110) (22 - 32) (9 - 19) (3.0 - 7.0) (45 - 90)	mmol/L mmol/L mmol/L mmol/L mmol/L umol/L



Patient NATAYA NAKHONWONG

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Relevant Results

(For additional pathology information contact Australian Clinical Labs 1300 367 674)

eGFR

> 90

> 90

> 90 (> 59) mL/min/1.73m2

22101589 Inpatient. Specialist management noted.

FBE-R ECU-C

All tests on this request have now been completed In the setting of infection, CRP levels >100 mg/L are supportive of bacterial rather than viral aetio logy.

Note results from this CRP assay should not be used for cardiac risk assessment. Please request the high sensitivity assay (hsCRP) instead. BIOCHEMISTRY

### C REACTIVE PROTEIN (CRP)

SPECIMEN: SERUM

Date	Time	Lab No.		CRP	Units	Ref. Range
01/11/24 31/10/24 29/10/24 16/11/23 26/09/23 21/07/23	10:26 07:30 16:27 09:00 08:11 11:20	22931848 90353812 22097900 21970672 84650448 82350225 80664654 80728690	*	2.5 5.1 6.1 9.7 < 0.7 < 0.7 < 0.7	mg/L	(< 3.0)

In the setting of infection, CRP levels >100~mg/L are supportive of bacterial rather than viral aetiology.

Note results from this CRP assay should not be used for cardiac risk assessment. Please request the high sensitivity assay (hsCRP) instead.

CRP-C FBE-R ECU-W

This request has other tests in progress at the time of reporting MRI SPINE (RECALL)

Findings: The patient has returned for additional sequences to supplement the MRI examination acquired yesterday, 30 October 2024. The sequences acquired today are; T1 axial and T2 axial of the upper lumbar spine, T1 axial of the lower lumbar spine, T1 FS Dixon post-contrast sagittal of the upper and lower spine, T1 FS Dixon post-contrast axial of the upper and lower lumbar spine.

There is no abnormal thickening or enhancement of the cauda equina nerve roots. There is no pathological intramedullary enhancement. No leptomeningeal enhancement is shown.

Comment: Post-contrast sequences have been acquired supplement the MRI examination acquired yesterday, 30 October 2024. There is no pathological spinal cord, cauda equina or leptomeningeal enhancement. There are no MRI features to suggest Guillain-Barre syndrome.

Radiologist: Dr R. Brazel

View images and report in PRC Direct: https://app.prcdirect.com.au/admin/results/15364756 ACTH is measured by Immunoassay on a Siemens IMMULITE. ENDOCRINOLOGY

ADRENOC	ORTICOTR	OPHIC HORMONE		SPECIMEN: I	PLASMA	
Date	Time	Lab No.	ACTH	Units	Range	
31/10/2	4 07:30	22097900	11	pg/ml	(< 46)	



NATAYA NAKHONWONG Patient

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Units

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Relevant Results

(For additional pathology information contact Australian Clinical Labs 1300 367 674)

ACTH is measured by Immunoassay on a Siemens IMMULITE.

ASO-V ANF-W ENA-W CRY-W ANC-W CRP-R ACT-C FBE-R ECU-R COR-R

This request has other tests in progress at the time of reporting Cortisol less than 300 nmol/L may be seen with normal diurnal rhythm, exogenous steroids or adrenal i nsufficiency. Consider repeat cortisol with ACTH at 08:00 to clarify. Note; estrogens increase cortis ol binding globulin and thus total but not usually free cortisol. Cortisol measured on the Siemens Ce ntaur. Reference Ranges: AM 110 - 550 nmol/L

PΜ 70 - 400 nmol/L **ENDOCRINOLOGY** 

**CORTISOL STUDIES** 

SPECIMEN: SERUM/PLASMA/URINE

Coll. Date: 31/10/24 30/10/24 Coll. Time: 07:30 09:40

Ref. Lab Number: 22097900 24917362 Range --------

Serum Cortisol See below nmol/L

Reference Ranges: AM 110 - 550 nmol/L 70 - 400 nmol/L PM

22097900 Cortisol less than 300 nmol/L may be seen with normal diurnal rhythm, exogenous steroids or adrenal insufficiency. Consider repeat cortisol with ACTH at 08:00 to clarify. Note; estrogens increase cortisol binding globulin and thus total but not usually free cortisol. Cortisol measured on the Siemens Centaur.

ASO-W ANF-W ENA-W CRY-W ANC-W CRP-R ACT-W FBE-R ECU-R COR-C

This request has other tests in progress at the time of reporting

Neutrophil fluorescence alone is not a diagnostically specific assay. Results of the specific ELISA t ests for myeloperoxidase (MPO-ANCA) and proteinase 3 (PR3-ANCA) will follow. IMMUNOLOGY SPECIMEN: SERUM

### ANTI-NEUTROPHIL CYTOPLASMIC ANTIBODIES (ANCA)

C - A.N.C.A. **POSITIVE** P - A.N.C.A. Negative

COMMENT: Neutrophil fluorescence alone is not a diagnostically specific assay. Results of the specific ELISA tests for myeloperoxidase

(MPO-ANCA) and proteinase 3 (PR3-ANCA) will follow.

ENH-W ASO-V ANF-W ENA-V CRY-W ANC-C CRP-R ACT-R FBE-R ECU-R

This request has other tests in progress at the time of reporting SPECIAL CHEMISTRY

**PROTEIN STUDIES** SPECIMEN: SERUM

CRYOGLOBULINS Not detected CRYOFTBRINGEN Not detected

ENH-W ASO-V ANF-W ENA-V CRY-C ANC-R CRP-R ACT-R FBE-R ECU-R

This request has other tests in progress at the time of reporting

Negative ANA is not associated with SLE. TMMUNOLOGY

**ANTINUCLEAR ANTIBODIES** 

SPECIMEN: SERUM



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Phone 0420665650 Mob 0420665650 Address 60 BALFOUR ROAD, SWAN VIEW, WA, 6056

Relevant Results (For additional pathology information contact Australian Clinical Labs 1300 367 674)

Anti-nuclear Antibody titre : < 40

(< 160)

COMMENT: Negative ANA is not associated with SLE.

ENH-W ASO-V ANF-C ENA-V CRY-R ANC-R CRP-R ACT-R FBF-R FCU-R

This request has other tests in progress at the time of reporting

24-22097900: High titre anti-

DNase B antibodies indicate recent infection with Streptococcus pyogenes. Anti-

DNase B is more sensitive than ASOT for the diagnosis of post-

streptococcla glomerulonephritis and serious streptococcal skin infections. The titre peaks at 4-6 weeks, but high levels persist for longer than the ASOT.

High titre ASOT, or a two-

fold rise in titre, indicates recent infection. Antibodies are not detected until at least 3 weeks po st infection. (see RCPA Manual.)

Validated by Assoc. Prof. Louise A Smyth, BA MBBS GCUT DipHPE FRC Anti deoxyribonuclease-B (anti-DNAse B) titres greater than 200 U/mL in adults and 300 U/mL in school age children are considered as significant. **IMMUNOLOGY** SPECIMEN; SERUM

#### STREPTOCOCCAL ANTIBODIES

anti-Streptolysin 0 160 IU/mL (<201)

anti-DNA''ase B 211 II/mI

Anti deoxyribonuclease-B (anti-DNAse B) titres greater than 200 U/mL in adults and 300 U/mL in school age children are considered as significant.

24-22097900: High titre anti-DNase B antibodies indicate recent infection with Streptococcus pyogenes. Anti-DNase B is more sensitive than ASOT for the diagnosis of post-streptococcla glomerulonephritis and serious streptococcal skin infections. The titre peaks at 4-6 weeks, but high levels persist for longer than the ASOT. High titre ASOT, or a two-fold rise in titre, indicates recent infection. Antibodies are not detected until at least 3 weeks post infection. (see RCPA Manual.) Validated by Assoc. Prof. Louise A Smyth, BA MBBS GCUT DipHPE FRCPA, Immunopathologist.

ENH-W ASO-C ANF-R ENA-V CRY-R ANC-R CRP-R ACT-R FBE-R ECU-R

This request has other tests in progress at the time of reporting

This result may occur in treated, inactive or relapsing granulomatosis with polyangiitis (Wegener), mi croscopic polyangiitis (with its renal-limited variant) and EPGA (Churg-Strauss syndrome). This result also occurs in chronic infections, inflammatory bowel disease and othe r autoimmune diseases where the clinical significance is unclear. **IMMUNOLOGY** SPECIMEN: SERUM

#### ANTI-NEUTROPHIL CYTOPLASMIC ANTIBODIES (ANCA)

C - ANCA **POSITIVE** P - ANCA Negative

#### REFERENCE RANGES

Negative Equivocal Positive PR3 - ANCA < 2.0 IU/ml (< 2.0) (2.0 - 3.0)(> 3.0)MPO - ANCA (<3.5) (3.5-5.0)< 3.5 IU/ml

COMMENT This result may occur in treated, inactive or relapsing granulomatosis with polyangiitis (Wegener), microscopic



For Further Patient Information please contact

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Relevant Results (For additional pathology information contact Australian Clinical Labs 1300 367 674) polyangiitis (with its renal-limited variant) and EPGA (Churg-Strauss syndrome). This result also occurs in chronic

infections, inflammatory bowel disease and other autoimmune diseases where the clinical significance is unclear.

ENH-W ASO-R ANF-R ENA-V CRY-R ANC-R CRP-R ACT-R FBE-R ECU-R

This request has other tests in progress at the time of reporting Consistent with ongoing inflammation. Suggest QEP and SFLC. Further result to follow.

Validated by Assoc. Prof. Louise A Smyth, BA MBBS GCUT DipHPE FRC PROTEIN STUDIES

#### Immunoglobulins

IgG	<b>28.59</b> g/L	(6.50 - 16.00)
IgA	2.3 g/L	(0.4 - 3.5)
IgM	1.90 g/L	(0.50 - 3.00)

Consistent with ongoing inflammation. Suggest QEP and SFLC. Further result to follow.

Validated by Assoc. Prof. Louise A Smyth, BA MBBS GCUT DipHPE FRCPA, Immunopathologist.

ENH-W ASO-R ANF-R ENA-V CRY-R ANC-R CRP-R ACT-R FBE-R ECU-R

This request has other tests in progress at the time of reporting Clinical Details: 38-year-old female with ongoing lower limb weakness, tingling, episodes of visual changes, sensation changes to sole of foot and lateral calf and proximal upper limbs. Ascending weakness for 4 days and sensory deficit. ? Guillain-Barre syndrome. ? Demyelinating disorder.

### MRI BRAIN

Technique: Unenhanced MRI brain, demyelination protocol, at 1.5 Tesla; T2 axial, DWI axial, 3-D FLAIR sagittal and 3-D T1 sagittal. No prior imaging is available for comparison.

Findings: There is no intra or extra-axial mass or collection. There is no diffusion restriction or diffusion restricting lesion. Grey-white matter differentiation is preserved. There is no significant T2 or FLAIR signal alteration in the brain parenchyma. In particular, there are no lesions characteristic for demyelination. Parenchymal volume is age appropriate. No hydrocephalus.

The orbits, pituitary gland, midline commissural structures and posterior fossa are normal. There is no Chiari malformation. The cervicomedullary junction and visualised upper cervical cord are unremarkable. The major intracranial flow voids are preserved.

There is T2 hyperintense inflammatory mucosal thickening scattered throughout the paranasal sinuses, maximal in the left sphenoid sinus where there is also a small volume of bubbly secretions. The mastoid temporal bones are clear.

Comment: There is no acute intracranial abnormality or mass lesion. There is no evidence of intracranial demyelination.



For Further Patient Information please contact

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Relevant Results

(For additional pathology information contact Australian Clinical Labs 1300 367 674)

MRI WHOLE SPINE

Technique: Unenhanced MRI whole spine at 1.5 Tesla; T1 sagittal, T2 sagittal and STIR sagittal of the upper and lower spine. T2 axial cervical spine and T2 axial lower lumbar spine. No prior imaging is available for comparison.

Findings: There is conventional vertebral segmentation. The vertebral column is visualised from skull base to S1 (inclusive).

The cervicomedullary junction is normal. The spinal cord is normal in volume and signal intensity. There is no intrinsic cord signal abnormality. In particular, there are no cord lesions characteristic for demyelination. No intra or extra medullary mass or collection. The conus medullaris terminates at the level of the L1/2 intervertebral disc. The unenhanced cauda equina nerve roots are normal.

Alignment of the spine is normal. There is no infiltrative marrow process or destructive osseous lesion. There is mild disc degeneration at C5/6 and L4/5 where there is slight desiccation of the intervertebral discs and minimal annular bulging. There is no focal disc protrusion, canal stenosis or neural impingement at any level.

There is degenerative facet arthropathy bilaterally at T11/12 as well as throughout the lower lumbar spine. The imaged paravertebral soft tissues are unremarkable.

Comment: No significant abnormality demonstrated. There is no evidence for spinal cord demyelination.

Guillain-Barre syndrome is not excluded on unenhanced MRI. The patient has been recalled for post-contrast sequences and a supplementary report will be issued.

Radiologist: Dr R. Brazel

View images and report in PRC Direct: https://app.prcdirect.com.au/admin/results/15362374 Normal TSH level. ENDOCRINOLOGY

SPECIMEN: SERUM

#### THYROID FUNCTION TEST

Date: 30/10/24 21/06/24 24/11/23 Coll. Time: 09:40 10:15 10:24 Lab Number: 24917362 21386624 84650242

TSH 1.17 1.03 1.23 (0.40 - 4.00)mIU/L

24917362 Normal TSH level.

BFO-W FBE-R ECU-W LFT-W COR-C TFT-C

This request has other tests in progress at the time of reporting

Normal B12 and folate results.

 RANGES
 B12
 Serum Folate

 Normal
 > 180
 > 10.0

 Equivocal
 150 - 180
 5.0 - 10.0

 Deficient
 < 150</td>
 < 5.0</td>

 BIOCHEMISTRY

VITAMIN B12 AND FOLATE SPECIMEN: SERUM/BLOOD

Date: Time: **30/10/24** 09:40



For Further Patient Information please contact Midland Public and Private Hospitals, 1 Clayton St, MIDLAND 6056, WA (08) 9462 4000

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(For additional pathology information contact Australian Clinical Labs 1300 367 674) Relevant Results

Lab Number: 24917362

\_\_\_\_\_\_

Vitamin B12 572 pmo1/L Folate 31.4 nmol/L

24917362 Normal B12 and folate results.

RANGES B12 Serum Folate Normal > 180 > 10.0 5.0 - 10.0 Equivocal 150 - 180 Deficient < 5.0 < 150

BFO-C FBE-R ECU-C LFT-C COR-C TFT-C

All tests on this request have now been completed CHEST X-RAY

Clinical Details: Bilateral lower limb weakness and upper limb weakness with on and off chest pain.? Guilain Barre syndrome.? Infection or pneumothorax.

Comparison has been made with a previous chest x-ray from June 6, 2024.

Heart size and mediastinal contours appear normal. The lungs appear clear with no consolidation or collapse.

There are no pleural effusions or features to suggest pulmonary oedema.

There is no pneumothorax or free gas beneath the diaphragm.

No acute osseous abnormality is seen.

Comment:

The lungs appear clear.

Radiologist: Dr L. Gallagher

View images and report in PRC Direct: https://app.prcdirect.com.au/admin/results/15358598

MOLECULAR BIOLOGY SPECIMEN: Respiratory tract swab(s)

CORONAVIRUS COVID-19 / SARS-COV-2 NAAT

COVID-19 virus NAAT Not detected

PAI-R HUM-R CVI-C REV-R FLU-R ZRM-R

All tests on this request have now been completed

This assay will detect the presence of all current circulating Influenza A strains. This assay detect s Respiratory syncytial virus subtypes A and B.

MOLECULAR BIOLOGY SPECIMEN: Respiratory tract swab(s)

### RESPIRATORY VIRUSES PCR

Influenza A virus RNA Influenza B virus RNA			Detected Detected
Respiratory syncytial virus	RNA	Not	Detected
Parainfluenza virus Type 1	RNA	Not	Detected
Parainfluenza virus Type 2	RNA	Not	Detected
Parainfluenza virus Type 3	RNA	Not	Detected
Parainfluenza virus Type 4	RNA	Not	Detected
Human Metapneumovirus RNA		Not	Detected
Human Adenovirus DNA		Not	Detected



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**Relevant Results** 

(For additional pathology information contact Australian Clinical Labs 1300 367 674)

Human Rhinovirus RNA

Detected

This assay will detect the presence of all current circulating Influenza A strains.

This assay detects Respiratory syncytial virus subtypes A and B.

PAI-C HUM-C CVI-W REV-C FLU-C ZRM-C

This request has other tests in progress at the time of reporting Clear and colourless BIOCHEMISTRY SPECIMEN: CSF

CEREBROSPINAL FLUID

Tube No.

2

Macroscopic appearance

Clear and colourless

**CHEMISTRY** 

\* Protein Glucose **0.52** g/L 3.0 mmol/L

(0.15 - 0.45)

(2.8 - 4.2)

CSF-W CSP-C JMD-W

This request has other tests in progress at the time of reporting

TEST NAME: pending

**PATHWEST** 

QEII Medical Centre Hospital Avenue NEDLANDS WA 6909

NEDLANDS WA 6909 Ph: 13 7284 Fax: 08 9346 3354

NATA Accredited Laboratory Number: 14481

Test was referred on 29/10/24

pending

UNK-W CSF-W CSP-W MBF-W JMD-W

This request has other tests in progress at the time of reporting

TEST NAME: CMV Quantitative PCR

This test was performed by: Fiona Stanley Hospital Central Specimen Reception Ground Floor, Pathology Bldg 102-118 Murdoch Drive MURDOCH WA 6150

Test was referred on 30/10/24

Report was received on 01/11/2024 12:35 pm

REF LAB ID G630006792-

Results received from testing institution

Microbiology PCR/ Nucleic Acid Amplification Test (NAAT)

Specimen: Cerebrospinal Fluid

Collected: 30/10/2024 12:05 Received: 30/10/2024

12:05 CMV DNA

Not Detected



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#### Relevant Results

(For additional pathology information contact Australian Clinical Labs 1300 367 674)

MCP-R RCV-R HVM-R CVP-C JMD-C ABR-n RVM-R CSF-V CSP-R

This request has other tests in progress at the time of reporting Elevated CRP noted. Possible reduced iron stores in the presence of an inflammatory response. Ferriti n is not a good index of iron stores in the presence of inflammation. More reliable assessment can be obtained after recovery from intercurrent illness. BIOCHEMISTRY

IRON STUDIES SPECIMEN: SERUM

Date: Coll. Time: Lab Number:	<b>29/10/24</b> 16:27 21970672	10:24	21/07/23 11:20 80664654	
Iron	** <b>5.1</b> 2.85 ** <b>7</b> 36	12.9	29.7	(10.0 - 30.0) umol/L
Transferrin		* <b>2.00</b>	* 2.00	(2.10 - 3.80) g/L
Saturation		26	* 59	(15 - 45) %
Ferritin		121	* 258	(30 - 200) ug/L

21970672 Elevated CRP noted. Possible reduced iron stores in the presence of an inflammatory response. Ferritin is not a good index of iron stores in the presence of inflammation. More reliable assessment can be obtained after recovery from intercurrent illness.

CRP-R QUA-R TGP-R TMV-N COP-R FBE-R CPM-C ECU-C LFT-C TCG-N

All tests on this request have now been completed

Jarrad Hall - Resident		06 Nov 20	24
Finalised By		Date	
Document Details			Version 28