

**Allison Geffen**

201-3029 Carling Ave Ottawa Ontario K2B 8E8

Tel: 613-721-0978 Fax: 613-596-0496

## Consultation Request

Date:	2025-10-23	Patient:	ALBERT, MICHELLE
Status:	Non-Urgent	Address:	81-3415 Uplands Dr 81- 3415 Uplands Dr Gloucester, OT, ON
Service:	endocrinologist	Phone:	613-292-1018
Consultant:	Malakieh, Nadia	Work Phone:	
Phone:	613-505-9704	Cell Phone:	613-292-1018
Fax:	613-505-9707	Email:	michelle.albert@ocdsb.ca
Address:	Unit 208 4100 Strandherd Drive Ottawa ON K2J0V2	Birthdate:	1984-05-10 (y/m/d)
		Sex:	F
		Health Card No.:	(ON) 4392210656 JX
		Appointment date:	
		Time:	
		Chart No.:	

## Reason for consultation:

**REFERRING FOR PCOS, MENORRHAGIA W pHX IRON DEF ANEMIA, MILD HYPERPROLACTIN, OBESITY - difficulty with weight loss**

## Pertinent Clinical Information:

IDA 7/25 (ferritin 5, Hg 107) - new\* Cholecystectomy 4/24 QCH ER - biliary colic (Surg Dr Bruce Gay)  
 Abn Pap -ASCUS (9/16 colposcopy, 2015, 2008 Dysplasia clinic Riverside, dr Cargill) q3y BMI=33  
 (2/21, 11/22) (Phx GDM) - LEAF Weight clinic 11/23, 7/24 (20 lb wt loss) TMJ dysfx (bite guard) 2023  
 Gastroscopy = normal 3/17 (Bx neg, celiac neg, h pylori neg) \*GI=Dr SEKAR Endometrial Polyp resection 5/16 GYNE = Dr COCHEN Hysteroscopy/Curettage Endometrium Keloid scar (DERM =DrLaBerge) (\*Dr CARGILL - 11/19 ASCUS PAP - need repeat q6m\*=Nx3) PCOS/DUB/irregular cycles/dysmenorrhea G1 T1 L1 C/S 9/19 Civic TOH Emergency C/S Chorioamnionitis \*GDM 9/19 \*\*monitor A1c/FBG/weight\*\* Obesity (2011 weight loss) Prolactin elevated 34 NYD 12/18 - referred Endo R ear tube childhood 5/19 Incompetent cervix preg/Cerclage/Bedrest (\*D/C colposcopy clinic 11/16, Pap q3y) (\*Blood Transfusion post delivery TOH Civic) 6/20 Paronychia Toe (cerclage preg) Asthma - inactive (11/21 covid pos) cervical polypectomy Emergency C/Section for chorioamnionitis (2019 Dr Cargill, Yvonne)

## Current Medications:

MOUNJARO KWIKPEN 5 MG/0.6 ML WEEK 5-8: 5 MG SC INJ WEEKLY, THEN DECIDE IF STAY ON THIS DOSE OR INCREASE Qty:4 Repeats:2

TECTA DR 40 MG TABLET 1 po od for ? gerd (\*7 /25 trial) Qty:30 Repeats:1

\*\*DISCONTINUE SAXENDA, REPLACING WITH MOUNJARO

MOUNJARO KWIKPEN 2.5 MG/0.6 ML \*to REPLACE Saxenda 3/25: START 2.5 mg SC INJ WEEKLY X 4 WEEKS Qty:4 weeks Repeats:0

CIPRALEX 10 MG TABLET 1 po od for mood/anxiety Qty:90 Repeats:3

RUPALL 10 MG TABLET 1 po od Antihistamine Qty:30 Repeats:0

DYMISTA 137-50 MCG NASAL SPRAY 1 spray per nostril BID Qty:1 Repeats:0

CLAVULIN 875 TABLET 1 po bid for sinus infection x 10 days Qty:20 Repeats:1

OMNARIS 50 MCG NASAL SPRAY 1-2 nasal spray od-bid for acute sinusitis Qty:1 Repeats:0

## Allergies:

No Known Drug Allergies

Referring Practitioner : Allison Geffen (010193)

MRP : Geffen, Allison (010193)

Requesting Physician : Allison Geffen (010193)

Signature:

A handwritten signature in black ink, appearing to read "A. Geffen".

**LEAF Weight Management Clinic**

1980 Ogilvie Road  
Ottawa ON, Canada  
K1J 9L3

# Fax Cover Sheet

To: Dr. Allison Sigal Geffen  
Phone: (613) 596-9840  
Fax: (613) 596-0496

Date: 2024-Apr-18

From: Dr. Colette Hansen  
Phone: (613) 701-1222  
Fax: (613) 701-1223

Number of pages including cover: 3

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Re: Michelle Albert

[www.leafwmc.com](http://www.leafwmc.com)

Phone: 613-701-1222  
Fax: 613-701-1223  
Email: [Info@leafwmc.com](mailto:Info@leafwmc.com)  
Address: 1980 Ogilvie Road  
Ottawa, ON K1J 9L3

2024-Apr-18

Dr. Geffen  
201-3029 Carling Ave  
Ottawa, ON K2B 8E8

**Patient:** Mrs. Michelle Albert  
**PHN:** 4392 210 656JX  
**Birthdate:** 1984-May-10

Dear Dr. Geffen,

I met with Michelle by video in follow-up on 2024-Apr-18. She completed our ROOT program today. This is an update on her progress and a summary of her program results.

Informed consent was obtained from this patient to communicate and provide care using virtual and other telecommunications tools. This patient has been explained the risks related to unauthorized disclosure or interception of personal health information and steps they can take to help protect their information. We have communicated that care provided through video or audio communication cannot replace the need for physical examination or an in-person visit for some disorders or urgent problems and the patient understands the need to seek urgent care in an emergency department as necessary.

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## PROGRAM SUMMARY

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Starting Weight: 222 pounds  
Ending Weight: 205 pounds  
Weight Change: -18 pounds

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## ASSESSMENT

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**Reason for Visit**

Follow up on weight management care plan.

Meal pattern: three meals per day, healthy snack in the morning

Program outcomes: improved portion control and meal structure, eliminated habitual evening snacking, decreased intake of restaurant sourced meals, better awareness of triggers for emotional eating and strategies for stress management, developed a better routine with physical activity

**Measurements**

**Self-Reported Height:** 67 Inches (2024-Apr-18)  
**Self-Reported Weight:** 205 lbs (2024-Apr-18)  
**Calculated BMI:** 32.1 Kg/m<sup>2</sup> (2024-Apr-18)

**Weight Classification:** None Recorded

**Physical Activity:**

Cardiovascular - walks during work break 3x week, afternoon walks with daughter

**Meals:**

First Meal - ProFlex protein shake - self made - banana, protein , ice , PB  
Second Meal - leftovers typically or homemade protein bowls and veggies  
Third Meal - chicken or beef with vegetable, rice, roommate orders Hello Fresh (planning to stop)

**Snacks:**

First Snack - morning: fruit  
Second Snack - afternoon: fruits or veg or nuts  
timing - evening , btu also in the dya if someone brings in donuts  
choices - chips. salty food  
reasons - stress makes her eat fast food, cravings

**Meal Replacement Consumption**

**Protein Supplement:** 7 servings per week

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**DIAGNOSIS****Nutrition Diagnosis**

Excessive energy intake and undesirable food choices related to competing values for behaviour change as evidenced by estimated intake inconsistent with estimated need and intake of high calorie density or large portions.

**From Previous:** unresolved and improving.

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**INTERVENTION****Nutrition Prescription**

Energy modified (1400 to 1600 calories), nutrient-dense, structured meal plan with meal replacement as commercial product at one meals per day positioned at breakfast.

**Nutrition & Health Behaviour Education**

energy and calorie awareness.

**Resources Provided**

meal planning aid

**Nutrition Counselling**

self monitoring and stimulus control.

**Goals**

Maintaining positive behaviours change

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**MONITORING AND EVALUATION**

The patient will follow-up with LEAF physician with an appointment on May 23, 2024.

Thank you for letting me participate in the care of your patient.

Colette Hansen, MS, RD, CBE  
CDO: 15055

Detail Results: Patient Info				Results Info		
Patient Name:	MICHELLE ALBERT	Home Phone:	(613)292-1018	Date of Service:	2025-10-20 16:46	
Date of Birth:	1984-05-10	Work Phone:		Date Received:	2025-10-21 21:02	
Age:	41 years	Sex:	F	Report Status:	Final	
Health #:	4392210656	Patient Location:	GDML	Client Ref. #:		
				Accession #:	YU-56302473	
Requesting Client: A.S. GEFFEN		cc: Client:				

**CHEMISTRY**

Test Name(s)	Result	Abn	Reference Range	Units	Date/Time Completed	Status
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**FERRITIN**

FERRITIN	8	L	30 - 125.00	ug/L	2025-10-21 20:53:31	F
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Result consistent with Iron Deficiency

For guidance, see [www.hemequity.com/raise-the-bar](http://www.hemequity.com/raise-the-bar)

TSH	2.84	N	0.35 - 5.00	mIU/L	2025-10-21 20:53:31	F
HEMOGLOBIN A1c	5.4	N	<= 5.99	%	2025-10-21 20:53:31	F
	NON-DIABETIC:	< 6.0 %				
	PREDIABETES:	6.0 - 6.4 %				
	DIABETIC:	> 6.4 %				
	OPTIMAL CONTROL:	< 7.0 %				
	SUB-OPTIMAL CONTROL:	7.0 - 8.4 %				
	INADEQUATE CONTROL:	> 8.4 %				

LH	24.5	N		IU/L	2025-10-21 20:53:31	F
	Follicular:	1.9 - 14.6				
	Ovulatory:	12.2 - 118.0				
	Luteal:	0.7 - 12.9				
	Post-menopausal:	5.3 - 65.4				

FSH	5	N		IU/L	2025-10-21 20:53:31	F
	Follicular:	3 - 15				
	Ovulatory:	5 - 23				
	Luteal:	1 - 9				

Post-menopausal: 16 - 157

**PROLACTIN**

PROLACTIN	25	H	<= 23.99	ug/L	2025-10-21 20:53:31	F
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Roche Diagnostics Electrochemiluminescence Immunoassay

(ECLIA).

Values obtained with different assay methods or  
kits may not be comparable and cannot be  
used interchangeably.

Testing for the potential presence of macroprolactin is suggested in asymptomatic hyperprolactinemic patients. If clinically indicated, "Macroprolactin" testing should be requested on an OHIP laboratory requisition. A fresh blood sample (serum) will need to be collected.

**HEMATOLOGY**

Test Name(s)	Result	Abn	Reference Range	Units	Date/Time Completed	Status
HEMOGLOBIN	119	N	110 - 147.000	g/L	2025-10-21 20:53:31	F
HEMATOCRIT	0.37	N	0.33 - 0.440	L/L	2025-10-21 20:53:31	F
RBC	4.7	N	3.8 - 5.200	x 10*12/L	2025-10-21 20:53:31	F
RBC INDICES: MCV	79	N	76 - 98.000	fL	2025-10-21 20:53:31	F
MCH	25	N	24 - 33.000	pg	2025-10-21 20:53:31	F
MCHC	318	N	313 - 344.000	g/L	2025-10-21 20:53:31	F
RDW	17.7	H	12.5 - 17.3		2025-10-21 20:53:31	F
WBC	6.3	N	3.2 - 9.400	x 10*9/L	2025-10-21 20:53:31	F
PLATELETS	271	N	155 - 372.000	x 10*9/L	2025-10-21 20:53:31	F
MPV	9.1	N	4.0 - 14.000	fL	2025-10-21 20:53:31	F

**DIFFERENTIAL WBC'S**

NEUTROPHILS	3.6	N	1.4 - 6.3	x 10*9/L	2025-10-21 20:53:31	F
LYMPHOCYTES	2.1	N	1.0 - 2.9	x 10*9/L	2025-10-21 20:53:31	F
MONOCYTES	0.3	N	0.2 - 0.8	x 10*9/L	2025-10-21 20:53:31	F
EOSINOPHILS	0.1	N	0.0 - 0.5	x 10*9/L	2025-10-21 20:53:31	F
BASOPHILS	0.00	N	0.00 - 0.09	x 10*9/L	2025-10-21 20:53:31	F

**INR**

INR	0.9 - 1.3	2025-10-21 20:53:31	F
GUIDELINES:	SUGGESTED INR RANGE:		
1. Most clinical conditions indicating anticoagulant therapy	2.0 - 3.0		
2. Patients with mechanical prosthetic heart valves	2.5 - 3.5		

\*\*\*\*\*  
\* NOTE: SPECIMEN RECEIVED NSQ. TESTING NOT PERFORMED \*  
\* DUE TO INAPPROPRIATE ANTICOAGULANT/BLOOD RATIO. \*  
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END OF REPORT
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Detail Results: Patient Info				Results Info		
Patient Name:	MICHELLE ALBERT	Home Phone:	(613)292-1018	Date of Service:	2025-07-14 14:27	
Date of Birth:	1984-05-10	Work Phone:		Date Received:	2025-07-16 17:46	
Age:	41 years	Sex:	F	Report Status:	Final	
Health #:	4392210656	Patient Location:	GDML	Client Ref. #:		
				Accession #:	YU-55038500	
Requesting Client:	A.S. GEFFEN	cc: Client:				

CHEMISTRY						
Test Name(s)	Result	Abn	Reference Range	Units	Date/Time Completed	Status
GLUCOSE SERUM RANDOM	5.7	N	3.6 - 7.70	mmol/L	2025-07-15 17:23:15	F
3.6 - 7.7 Normal random glucose						
7.8 - 11.0 Potentially at risk for diabetes; consider repeat fasting						
>11.0 Provisional diagnosis of Diabetes Mellitus						
CREATININE						
CREATININE	62	N	50 - 100.00	umol/L	2025-07-15 17:23:15	F
eGFR	111	N	60 - 99999.99	ml/min/1.73 m <sup>2</sup>	2025-07-15 17:23:15	F
eGFR is calculated using the CKD-EPI 2021 equation which does not use a race-based adjustment.						
An eGFR result >=60 ml/min/1.73m**2 rules out CKD stage 3-5. Assessment of urine ACR is required to definitively rule out or confirm CKD diagnosis. The KidneyWise toolkit (kidneywise.ca) recommends remeasuring eGFR and urine ACR annually for people with diabetes mellitus and less frequently in others unless clinical circumstances dictate otherwise.						
HOURS FASTING	NA	N		hrs	2025-07-15 17:23:15	F
CHOLESTEROL	4.75	N	<= 5.19	mmol/L	2025-07-15 17:23:15	F
Total cholesterol and HDL-C used for risk assessment and to calculate non-HDL-C.						
TRIGLYCERIDES	1.29	N	<= 1.69	mmol/L	2025-07-15 17:23:15	F
If nonfasting, triglycerides <2.00 mmol/L desired.						
HDL CHOLESTEROL	1.31	N	1.30 - 9999.00	mmol/L	2025-07-15 17:23:15	F
F: >=1.30 mmol/L HDL-C <1.30 mmol/L indicates risk for metabolic syndrome.						
LDL CHOLESTEROL CALC.	2.91	N	<= 3.49	mmol/L	2025-07-15 17:23:15	F
LDL-C was calculated using the NIH equation.						
For additional LDL-C and non-HDL-C thresholds based on risk stratification, refer to 2021 CCS Guidelines.						
NON-HDL-CHOLESTEROL(CALC)	3.44	N	<= 4.19	mmol/L	2025-07-15 17:23:15	F
TC/HDL-C RATIO	3.6	N			2025-07-15 17:23:15	F
VITAMIN B12	308	N	221 - 918	pmol/L	2025-07-15 17:23:15	F
60% of symptomatic patients have a						

**CHEMISTRY**

Test Name(s)	Result	Abn	Reference Range	Units	Date/Time Completed	Status
hematologic or neurologic response to						
B12 supplementation at a level < 148 pmol/L						
Vitamin B12 Deficiency: < 148 pmol/L						
Vitamin B12 Insufficiency: 148 to 220 pmol/L						

**FERRITIN**

FERRITIN	5	L	30 - 125.00	ug/L	2025-07-15 17:23:15	F
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Result consistent with Iron Deficiency

For guidance, see [www.hemequity.com/raise-the-bar](http://www.hemequity.com/raise-the-bar)

SODIUM	138	N	136 - 146.00	mmol/L	2025-07-15 17:23:15	F
POTASSIUM	3.8	N	3.7 - 5.4	mmol/L	2025-07-15 17:23:15	F
ALT	21	N	<= 35.99	U/L	2025-07-15 17:23:15	F
TSH	1.58	N	0.35 - 5.00	mIU/L	2025-07-15 17:23:15	F
HEMOGLOBIN A1c	5.5	N	<= 5.99	%	2025-07-15 17:23:15	F
NON-DIABETIC: < 6.0 %						
PREDIABETES: 6.0 - 6.4 %						
DIABETIC: > 6.4 %						
OPTIMAL CONTROL: < 7.0 %						
SUB-OPTIMAL CONTROL: 7.0 - 8.4 %						
INADEQUATE CONTROL: > 8.4 %						
RHEUMATOID FACTOR	<10	N	<= 13.99	IU/mL	2025-07-15 20:09:49	F
CRP	2.4	N	<= 7.99	mg/L	2025-07-15 20:09:49	F
GLUCOSE SERUM RANDOM	5.7	N	3.6 - 7.70	mmol/L	2025-07-16 17:37:01	F
3.6 - 7.7 Normal random glucose						
7.8 - 11.0 Potentially at risk for diabetes; consider repeat fasting						
>11.0 Provisional diagnosis of Diabetes Mellitus						

**CREATININE**

CREATININE	62	N	50 - 100.00	umol/L	2025-07-16 17:37:01	F
eGFR	111	N	60 - 99999.99	mL/min/1.73 m <sup>2</sup>	2025-07-16 17:37:01	F

eGFR is calculated using the CKD-EPI 2021 equation

which does not use a race-based adjustment.

An eGFR result >=60 mL/min/1.73m\*\*2 rules out CKD stage 3-5. Assessment of urine ACR is required to definitively rule out or confirm CKD diagnosis. The KidneyWise toolkit ([kidneywise.ca](http://kidneywise.ca)) recommends remeasuring eGFR and urine ACR annually for people with diabetes mellitus and less frequently in others unless clinical circumstances dictate otherwise.

HOURS FASTING	NA	N	hrs	2025-07-16 17:37:01	F
CHOLESTEROL	4.75	N	<= 5.19	mmol/L	2025-07-16 17:37:01
Total cholesterol and HDL-C used					
for risk assessment and to calculate non-HDL-C.					
TRIGLYCERIDES	1.29	N	<= 1.69	mmol/L	2025-07-16 17:37:01
If nonfasting, triglycerides <2.00 mmol/L desired.					

**CHEMISTRY**

Test Name(s)	Result	Abn	Reference Range	Units	Date/Time Completed	Status
HDL CHOLESTEROL	1.31	N	1.30 - 9999.00	mmol/L	2025-07-16 17:37:01	F
F: >=1.30 mmol/L indicates risk for metabolic syndrome.						
LDL CHOLESTEROL CALC.	2.91	N	<= 3.49	mmol/L	2025-07-16 17:37:01	F
LDL-C was calculated using the NIH equation.						
For additional LDL-C and non-HDL-C thresholds based on risk stratification, refer to 2021 CCS Guidelines.						
NON-HDL-CHOLESTEROL(CALC)	3.44	N	<= 4.19	mmol/L	2025-07-16 17:37:01	F
TC/HDL-C RATIO	3.6	N			2025-07-16 17:37:01	F
VITAMIN B12	308	N	221 - 918	pmol/L	2025-07-16 17:37:01	F
60% of symptomatic patients have a hematologic or neurologic response to B12 supplementation at a level < 148 pmol/L						
Vitamin B12 Deficiency: < 148 pmol/L						
Vitamin B12 Insufficiency: 148 to 220 pmol/L						

**FERRITIN**

FERRITIN	5	L	30 - 125.00	ug/L	2025-07-16 17:37:01	F
Result consistent with Iron Deficiency For guidance, see <a href="http://www.hemequity.com/raise-the-bar">www.hemequity.com/raise-the-bar</a>						
SODIUM	138	N	136 - 146.00	mmol/L	2025-07-16 17:37:01	F
POTASSIUM	3.8	N	3.7 - 5.4	mmol/L	2025-07-16 17:37:01	F
ALT	21	N	<= 35.99	U/L	2025-07-16 17:37:01	F
TSH	1.58	N	0.35 - 5.00	mIU/L	2025-07-16 17:37:01	F
HEMOGLOBIN A1c	5.5	N	<= 5.99	%	2025-07-16 17:37:01	F
NON-DIABETIC: < 6.0 %						
PREDIABETES: 6.0 - 6.4 %						
DIABETIC: > 6.4 %						
OPTIMAL CONTROL: < 7.0 %						
SUB-OPTIMAL CONTROL: 7.0 - 8.4 %						
INADEQUATE CONTROL: > 8.4 %						

**HEMATOLOGY**

Test Name(s)	Result	Abn	Reference Range	Units	Date/Time Completed	Status
HEMOGLOBIN	107	L	110 - 147.000	g/L	2025-07-15 17:23:15	F
HEMATOCRIT	0.35	N	0.33 - 0.440	L/L	2025-07-15 17:23:15	F
RBC	4.6	N	3.8 - 5.200	x 10*12/L	2025-07-15 17:23:15	F
RBC INDICES: MCV	75	L	76 - 98.000	fL	2025-07-15 17:23:15	F
MCH	23	L	24 - 33.000	pg	2025-07-15 17:23:15	F
MCHC	308	L	313 - 344.000	g/L	2025-07-15 17:23:15	F
RDW	18.8	H	12.5 - 17.3		2025-07-15 17:23:15	F
WBC	6.1	N	3.2 - 9.400	x 10*9/L	2025-07-15 17:23:15	F

**HEMATOLOGY**

Test Name(s)	Result	Abn	Reference Range	Units	Date/Time Completed	Status
E.S.R.	8	N	0 - 20.000	mm/hr	2025-07-15 17:23:15	F
PLATELETS	257	N	155 - 372.000	x 10 <sup>9</sup> /L	2025-07-15 17:23:15	F
MPV	8.7	N	4.0 - 14.000	fL	2025-07-15 17:23:15	F

**DIFFERENTIAL WBC'S**

NEUTROPHILS	3.4	N	1.4 - 6.3	x 10 <sup>9</sup> /L	2025-07-15 17:23:15	F
LYMPHOCYTES	2.2	N	1.0 - 2.9	x 10 <sup>9</sup> /L	2025-07-15 17:23:15	F
MONOCYTES	0.3	N	0.2 - 0.8	x 10 <sup>9</sup> /L	2025-07-15 17:23:15	F
EOSINOPHILS	0.1	N	0.0 - 0.5	x 10 <sup>9</sup> /L	2025-07-15 17:23:15	F
BASOPHILS	0.06	N	0.00 - 0.09	x 10 <sup>9</sup> /L	2025-07-15 17:23:15	F

**SMEAR**

CONSULTANT'S REPORT TO FOLLOW.

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HEMOGLOBIN	107	L	110 - 147.000	g/L	2025-07-16 17:37:01	F
HEMATOCRIT	0.35	N	0.33 - 0.440	L/L	2025-07-16 17:37:01	F
RBC	4.6	N	3.8 - 5.200	x 10 <sup>12</sup> /L	2025-07-16 17:37:01	F
RBC INDICES: MCV	75	L	76 - 98.000	fL	2025-07-16 17:37:01	F
MCH	23	L	24 - 33.000	pg	2025-07-16 17:37:01	F
MCHC	308	L	313 - 344.000	g/L	2025-07-16 17:37:01	F
RDW	18.8	H	12.5 - 17.3		2025-07-16 17:37:01	F
WBC	6.1	N	3.2 - 9.400	x 10 <sup>9</sup> /L	2025-07-16 17:37:01	F
E.S.R.	8	N	0 - 20.000	mm/hr	2025-07-16 17:37:01	F
PLATELETS	257	N	155 - 372.000	x 10 <sup>9</sup> /L	2025-07-16 17:37:01	F
MPV	8.7	N	4.0 - 14.000	fL	2025-07-16 17:37:01	F

**DIFFERENTIAL WBC'S**

NEUTROPHILS	3.4	N	1.4 - 6.3	x 10 <sup>9</sup> /L	2025-07-16 17:37:01	F
LYMPHOCYTES	2.2	N	1.0 - 2.9	x 10 <sup>9</sup> /L	2025-07-16 17:37:01	F
MONOCYTES	0.3	N	0.2 - 0.8	x 10 <sup>9</sup> /L	2025-07-16 17:37:01	F
EOSINOPHILS	0.1	N	0.0 - 0.5	x 10 <sup>9</sup> /L	2025-07-16 17:37:01	F
BASOPHILS	0.06	N	0.00 - 0.09	x 10 <sup>9</sup> /L	2025-07-16 17:37:01	F

**SMEAR**

SMEAR REFERRED TO PATHOLOGIST FOR REVIEW.

\*\* PLEASE REVIEW THIS REPORT AS ADDITIONAL TESTING HAS BEEN ADDED/REPORTED.

**CONSULTANT'S REPORT**

QUERY IRON STATUS.

REVIEWED BY DR. LINDA LACROIX M.D., F.R.C.P.

**AUTOIMMUNE/IMMUNOHEMATOLOGY**

Test Name(s)	Result	Abn	Reference Range	Units	Date/Time Completed	Status
<b>ANTINUCLEAR AB. (ANA)</b>						
ANTINUCLEAR AB. (ANA)	NEGATIVE	N	NEGATIVE		2025-07-15 20:09:49	F

A negative test is strong evidence against a diagnosis of rheumatic disease but is not conclusive. Results should be interpreted in conjunction with other serological tests and clinical findings.

MICHELLE ALBERT

END OF REPORT