



Patient Referral Form

Barrie Mid-Toronto Brampton Etobicoke Markham Oakville Thornhill

PATIENT INFORMATION:

Hammoud, Mohammad Ahmad RS 10812

Name: 429 FAITH DR
Mississauga, ON L5R 3Y7
Health Card: 647-469-5797(M)

DOB: M Mar 6, 1960
8513 425 887 FW

(dd/mm/yyyy)
Uninsured Specify:

Address: _____

(unit) _____

(city) _____

(postal code) _____

(e-mail address) _____

(home #) _____

(work # with extension) _____

(other #) _____

DIABETES/ENDOCRINOLOGY PLEASE SPECIFY:**The following investigations would be helpful:**

<input type="radio"/> Diabetes	<input type="radio"/> Type 1	<input checked="" type="radio"/> Type 2	<input type="radio"/> GDM	<input type="radio"/> FPG, A1C, Lipids, Renal Function, uACR
<input type="radio"/> Newly Diagnosed Diabetes Rapid Triage (1-2 weeks)				<input type="radio"/> FPG, A1C, Lipids, Renal Function, uACR

Consultation & shared care Consultation only

<input type="radio"/> Thyroid	<input type="radio"/> Thyroid function, Relevant imaging
<input type="radio"/> Osteoporosis	<input type="radio"/> BMD report <2 years, other relevant labs
<input type="radio"/> Lipids	<input type="radio"/> TC, LDL, HDL (<3 months), A1C
<input type="radio"/> PCOS	<input type="radio"/> LH, FSH, estrogen, testosterone, A1C
<input type="radio"/> Other (please specify):	

Notes: 65 yr. old ♂ with DM₂ Current Medications: Janumet XR 50-1000 mg
CAD, angioplasty & RA (Valsartan 80 mg OD, Forxiga 10 mg PO OD)
Review medication (Naproxen 375 mg BID, atorvastatin 20 mg
methotrexate 25 mg TID, ezetimibe 10 mg OD, Lyncia bisoprolol 2.5 mg OD)
Folic acid
Referred By: Dr. Reem Z. Salim Referring Physician Billing #: 031044

Referring Physician Signature: R

Date: 22 Oct 2025

Oct 14, 2025

LIFELABS ONTARIO Lab Data (Updated)

RS

Accession Number
 Collection Date
 Ordering Physician: FLORICA, BRANNICKA
 Result Copy To: FLORICA, BRANNICKA
 Result Copy To: Salim, Reem

2025-EA2670752
 Oct 14, 2025 10:41AM

ESR

Report Date Oct 14, 2025 7:17PM
 Lar Licence #: 5407
 Testing location name and address: KENNEDY 6500 Kennedy Road Mississauga Ontario L5T 2X4 Canada B
 ESR 5 2 + 30

Complete Blood Count

Report Date	Oct 14, 2025 6:33PM	
WBC	5.7	4.0 - 12.0
RBC	5.20	4.90 - 6.10
Hb	146	117 - 177
Hct	0.452	0.400 - 0.500
MCV	87	80 - 100
MCH	28.1	27.0 - 33.0
MCHC	323	308 - 340
RDW	15.0 (N)	14.0 - 18.0
Platelets	224	100 - 400
Neutrophils #	4.0	0.0 - 7.5
Lymphocytes #	1.1	0.0 - 3.5
Monocytes #	0.4	0.2 - 3.0
Eosinophils #	0.1	0.0 - 0.5
Basophils #	0.1	0.0 - 0.5
Immature Granulocytes #	0.0	0.0 - 0.1
Nucleated RBC's as % of WBCs	0	

Urinalysis Chemical

Report Date	Oct 14, 2025 5:19PM	
Collection Date	Oct 14, 2025	
Collection Time	10:41	
Urine Appearance	CLEAR	TRANSPARENT
Urine Colour	YELLOW	TRANSPARENT
Urine Specific Gravity	>=1.030	
Urine pH	6.0	5.0 - 8.0
Urine Protein	0.3 (N)	Negative
Urine Glucose	>=55 (N)	Negative
Urine Ketones	NEGATIVE	Negative
Urine Erythrocytes	NEGATIVE	Negative
Urine Nitrite	NEGATIVE	Negative
Urine Leukocyte Esterase	NEGATIVE	Negative

TEST COMMENT Please see <https://tests.lifelabs.com/s/article/URINALYSIS-CHEMICAL-Ontario-for-alternative-reporting-units>.

Report Date Oct 14, 2025 6:53PM
 FBS 6.7 (N) 5.0 - 6.7

Fasting Glucose greater than or equal to 7.0 mmol/L after an 8 hr fast can be used as a provisional diagnosis of diabetes mellitus. If asymptomatic, a repeat confirmation test using Fasting Glucose, HbA1c, or A1c (G7) must be done.

Report Date Oct 14, 2025 6:53PM
 Albumin 44 mg/dL 20 - 250
 Report Date Oct 14, 2025 6:53PM
 ALT 37 U/L 0 - 40

Hemoglobin A1c

Report Date Oct 14, 2025 8:21PM
 Hb A1c 9.0 (N) 5.0 - 6.0

Diabetes Canada 2018 Guidelines:

Screening and Diagnosis:

< 5.7 Normal

5.5% - 5.9% At risk
 6.0% - 6.4% Prediabetes
 ADR= 6.5% Diabetes Mellitus***

***Regarding diagnosis: in the absence of symptomatic hyperglycemia, if a single laboratory test result is in the diabetes range, a repeat confirmatory laboratory test (HbA1c, A1C, 2hIG in a 75 g OGTT) must be done on another day for diagnosis confirmation.

 Monitoring: <IF= 7.0 %
 Target in adults without comorbidities. Other targets may be more appropriate in children, elderly and patients with comorbidities.

 Results may not accurately reflect mean blood glucose in patients with hemoglobin variants, disorders associated with abnormal erythrocyte turnover, severe renal and liver disorders.

Creatinine/GFR

Report Date	Oct 14, 2025 6:53PM		
Cr	79	77 - 117	mg/dL
eGFR	95	63.4 - 99.9	ml/min/1.73m ²

Reference interval: =>60 mL/min/1.73m²

eGFR is calculated using the CKD-EPI GFR equation which does not use a race-based adjustment.

Lipid Assessment

Report Date	Oct 14, 2025 6:53PM		
Hours after a Meal	12		
TG	1.40		
FASTING:	<1.70 mmol/L		
NON-FASING:	<2.00 mmol/L		
CHOL	3.07	3.00 - 4.00	
Total cholesterol and HDL-C used for risk assessment and to calculate non-HDL-C.			
HDL	1.07	1.00 - 1.60	
HDL-C <1.00 mmol/L indicates risk for metabolic syndrome.			
NON-HDL	2.00	1.00 - 3.00	
Non-HDL-Cholesterol is not affected by the fasting status of the patient.			
LDL	1.43	1.00 - 1.60	
LDL-C is calculated using the NCEP equation.			

For additional LDL-C and non-HDL-C thresholds based on risk stratification, refer to 2021 CCS Guidelines. Can J Cardiol. 2021;37:1119-1150.

CHOL/HDL

Cholesterol/HDL-C is not included in the 2021 CCS guideline as a lipid initiation or treatment target but is recognized as an indicator of high CVI risk at Cholesterol/HDL-C ratio >6.0

Albumin Creatinine Ratio Urine Random

Report Date	Oct 16, 2025 4:49PM		
5 Year FPRF	NOT APPLICABLE	-	%

Results indicate mild to moderate albuminuria reflecting increased risk of CVD progression. If this is the first result with an ACR >3, confirm with at least 2 of 3 elevated results within 3 months.

If there is hematuria (>20rbc/hpf confirmed on urine microscopy), refer to nephrology.

Remeasure eGFR and urine ACP annually for patients with diabetes mellitus.

See the KidneyWise toolkit (kidneywise.ca) for further management recommendations including when to refer to nephrology.

Microalbumin

154

No reference interval has been established for this test.

Urine Creatinine Random

3.1

No reference interval has been established for this test.

Microalbumin/Creatinine Ratio

16.9 (H) 0.0

Oct 14, 2025 6:53PM

C-Reactive Protein

1.6 0.0

Test method: Abbott Alinity CRP, suitable for cardiovascular disease assessment and detection of active inflammation.

CRP >1.0 mg/L is a risk-enhancing factor for cardiovascular disease, as defined in the Guidelines of the American Heart Association and the American College of Cardiology (JACC 2019; 74: e177).

Report Date

Oct 14, 2025 6:53PM

RF

43 (A)

Reference range:

Negative: <30 IU/mL

Intermediate: 30-50 IU/mL

Positive: >50 IU/mL

Feb 9, 2024

Ontario Laboratories Information System Lab Data RS



Accession Number

2024-EA6401226

Collection Date

Feb 9, 2024 1:42PM

Ordering Physician: FLORICA, EPANDUSA

Result Copy To: FLORICA, EPANDUSA

Copy To physician was not subsequently identified.

Erythrocyte Sedimentation Rate - WHOLE BLOOD (Final)

- ESR	4	0.0 - 20
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Complete Blood Count - WHOLE BLOOD (Final)

- WBC	6.7	4.0 - 11.0
- RBC	5,00	4,30 - 6,30
- Hb	146	110 - 150
- Hct	0.435	0.390 - 0.460
- MCV	87	80 - 100
- MCH	29.2	27.0 - 31.0
- MCHC	336	310 - 360
- RDW	14.7 (B)	11.0 - 14.5
- Platelets	196	100 - 400
- Neutrophils #	4.4	1.0 - 7.5
- Lymphocytes #	1.4	0.8 - 2.5
- Monocytes #	0.7	0.2 - 1.0
- Eosinophils #	0.1	0.0 - 1.0
- Basophils #	0.1	0.0 - 0.2
- Immature Granulocytes #	0.0	0.0 - 0.1
- Nucleated RBC's as % of RBCs	0	

Creatinine - SERUM (Final)

- Cr	95	0.0 - 117
- eGFR	84	

An eGFR from 60-89 mL/min/1.73 m² mL is consistent with mildly decreased kidney function. However, in the absence of other evidence of kidney disease, eGFR values in this range do not fulfill the KDIGO criteria for chronic kidney disease. Interpret results in concert with ACR measurement.

Effective May 4 2018, eGFR is calculated using the CKD-EPI 2009 equation.

KDIGO 2012 guidelines highlight the importance of eGFR and urine albumin:creatinine ratio (ACR) in screening, diagnosis and management of CKD. Results for eGFR should be interpreted in concert with ACR.

Alanine Aminotransaminase - SERUM (Final)

- ALT	24	0-40
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C Reactive Protein - SERUM (Final)

- C-Reactive Protein	1.9	0.0
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Test method: Abbott Alinity CRP, suitable for cardiovascular disease assessment and detection of active inflammation. CRP >=2.0 mg/L is a risk-enhancing factor for cardiovascular disease, as defined in the Guidelines of the American Heart Association and the American College of Cardiology (JACC 2016; 74: e177).