

COVID-19 Prophylaxis Clinic Referral Form

Scarborough Health Network (Centenary site)	2867 Ellesmere Rd (Virtual Clinic)	Clinic Tel: 416-284-8131 ext 7788 Fax: 416-623-1213
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Evusheld™ (Tixagevimab and cilgavimab) is indicated for the pre-exposure prophylaxis of COVID-19 in adults and adolescents 12 years of age and older weighing at least 40 kg.

Current patients of the Scarborough Health Network (SHN)'s Oncology/Haematology program who have blood cancer (e.g. CAR T-cell therapy recipients, stem cell transplant, CD-20 inhibitors, receiving Bruton Tyrosine Kinase (BTK) inhibitors, venetoclax or other treatment for malignant hematology) will be identified and assessed for Evusheld eligibility by the patient's SHN Oncologist/Haematologist and do not require a referral to the SHN COVID-19 Prophylaxis Clinic.

Patient Information			
Name: _____		Gender: <input type="checkbox"/> M <input type="checkbox"/> F	Date of Birth (DD/MM/YY): _____
Address: _____			
Postal Code: _____		Phone Number: _____	HCN: _____
Allergies: <input type="checkbox"/> No known drug allergies		Vaccination Status: <input type="checkbox"/> 3 or more doses of COVID-19 vaccine received <input type="checkbox"/> Greater than 2 weeks since last dose of COVID-19 vaccine received <input type="checkbox"/> Other (specify): _____	
Weight (kg): _____		Patient Consent: <input type="checkbox"/> SHN Evusheld Consent Form has been fully reviewed with patient and the patient has provided verbal/signed consent. (Attach signed consent form to this referral form if completed).	
CRITERIA FOR USE			
ALL fields from A to E must be completed to be eligible for treatment for Evusheld			
A	<input type="checkbox"/> Patient 12 years and older		
B	<input type="checkbox"/> Weight equal or greater than 40 kg		
C	<input type="checkbox"/> Patient is NOT currently infected with COVID-19		
D	<input type="checkbox"/> Patient has not had a known recent exposure to an individual infected with COVID-19 (within 8 days)		
E		Solid Organ Transplant (Patients 12 years and older)	Malignant Hematology
			Adults (age 18 years and older)
			Children (age 12 to 17 years)
	Criteria 1	<input type="checkbox"/> All lung transplant <input type="checkbox"/> Recent transplant less than 6 months <input type="checkbox"/> B-cell depletion (rituximab) within the last 6 months <input type="checkbox"/> Plasmapheresis/ATG rejection (excluding patients with ongoing plasmapheresis) within last 3 months	<input type="checkbox"/> CAR T-cell therapy <ul style="list-style-type: none"> Ideally within 1 year of CAR T-cell therapy OR Could be given prior to CAR T-cell therapy <input type="checkbox"/> Allogeneic Stem Cell Transplant <ul style="list-style-type: none"> Ideally within 1 year of allogeneic stem cell transplant, or if patient remains on prednisone or other therapy with significant immuno-suppression effects OR Could be given prior to allogeneic stem cell transplant conditioning <input type="checkbox"/> Treated with CD-20 Inhibitors <ul style="list-style-type: none"> Patients on active treatment OR Within 6 months of treatment
Criteria 2	<input type="checkbox"/> All organs transplant patients age 60 years and older	<input type="checkbox"/> Treated with Bruton Tyrosine Kinase (BTK) inhibitors or venetoclax <ul style="list-style-type: none"> Patients on active treatment OR within 6 months of treatment <input type="checkbox"/> Autologous Stem Cell Transplant <ul style="list-style-type: none"> Ideally within 6 months of autologous stem cell transplant OR Could be given prior to allogeneic stem cell transplant conditioning 	<i>Not routinely recommended for autologous stem cell transplant patients</i>
Criteria 3	<input type="checkbox"/> All organs transplant patients age less than 60 years	<input type="checkbox"/> Other Malignant Hematology Patients <ul style="list-style-type: none"> Patients on active treatment OR Within 3 months of active therapy 	<input type="checkbox"/> Other Malignant Hematology Patients <i>Based on discretion of subspecialist and within 3 months if considered</i>



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Criteria 4	<input type="checkbox"/> Patients receiving anti-B-cell therapy (e.g. rituximab) (all patients 12 years and older)
Criteria 5	<input type="checkbox"/> Patients with significant primary immunodeficiency (all patients 12 years and older) * (see <i>Appendix A</i> for examples of common primary immunodeficiencies on page 3)
Prescriber/Referee Attestation (Must be completed to be eligible for treatment)	
<input type="checkbox"/> I affirm that my patient meets above criteria for use	
Prescriber/Referee Name (print): _____ Direct Contact Number (not office line): _____	
Prescriber/Referee Signature: _____ Date/Time: _____	
CPSO# (if applicable): _____	
Referral Physician's Billing # (if applicable): _____	
<input type="checkbox"/> RN/RPN (if applicable): _____	
<input type="checkbox"/> Referred from SHN COVID-19 Vaccine Clinic (if applicable): _____	

Appendix A: Examples of common primary immunodeficiencies (*this list only includes the most common conditions and does not include all types of primary immunodeficiencies*)¹

- Activated PI3K Delta Syndrome (APDS)
- Agammaglobulinemia: X-Linked and Autosomal Recessive
- Ataxia Telangiectasia
- Cartilage Hair Hypoplasia (CHH)
- Chronic Granulomatous Disease (CGD) and Other Phagocytic Cell Disorders
- Chronic Mucocutaneous Candidiasis (CMC)
- Chronic Neutropenia
- Comel-Netherton Syndrome
- Common Variable Immune Deficiency (CVID)
- Complement Deficiencies
- Congenital Athymia
- DiGeorge Syndrome
- Drug-Induced Antibody Deficiency
- Heavy Chain Deficiencies
- Hemophagocytic Lymphohistiocytosis (HLH)
- Hoyeraal-Hreidarsson Syndrome (Dyskeratosis Congenita)
- Hyper IgE Syndrome
- Hyper IgM Syndromes
- IgG Subclass Deficiency
- Immunodeficiency with Centromeric Instability and Facial Anomalies (ICF)
- Immunodeficiency with Thymoma (Good's Syndrome)
- Innate Immune Defects
- Kappa Chain Deficiency
- Leukocyte Adhesion Deficiency (LAD)
- NEMO Deficiency Syndrome
- Post-Meiotic Segregation (PMS2) Disorder
- Schimke Syndrome
- Selective IgA Deficiency
- Selective IgM Deficiency
- Severe Combined Immune Deficiency (SCID) and Combined Immune Deficiency
- Specific Antibody Deficiency
- Transient Hypogammaglobulinemia of Infancy
- Transcobalamin II Deficiency
- Unspecified Hypogammaglobulinemia
- Veno-occlusive Disease (VODI)
- WHIM Syndrome (Warts, Hypogammaglobulinemia, Infections, and Myelokathexis)
- Wiskott-Aldrich Syndrome
- X-linked Lymphoproliferative (XLP) Syndromes 1 and 2
- X-linked Immune Dysregulation with Polyendocrinopathy (IPEX) Syndrome

Reference:

1. Immune Deficiency Foundation. Specific PI Diagnoses. Available from: <https://primaryimmune.org/specific-pi-diagnoses> [Accessed 21st October 2022]