Essai Clinique

Généré le 16 mai 2025 à partir de

Titre	Capturing Canadian Real-World Data for AML Chemotherapy Ineligible Patients on Venetoclax
Protocole ID	LIVEN (P23-363)
ClinicalTrials.gov ID	NCT05424562
Type(s) de cancer	Leucémie myéloïde aiguë (LMA)
Phase	Autres
Type étude	Autre
Institution	CENTRE UNIVERSITAIRE DE SANTE MCGILL SITE GLEN 1001 boul. Décarie , Montréal, QC, H4A 3J1
Ville	
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Statut	Actif en recrutement
Date d'activation	27-01-2023
But étude	Acute Myeloid Leukemia (AML) is a cancer of the blood and bone marrow and is the most common acute leukemia in adults. This study will evaluate how well venetoclax works to treat AML in adult participants who are ineligible for intensive induction chemotherapy in Canada.Venetoclax is a drug approved to treat Acute Myeloid Leukemia (AML). All study participants will receive Venetoclax as prescribed by their study doctor in accordance with approved local label. Adult participants with a new diagnosis of AML who are ineligible for intensive induction chemotherapy will be enrolled. Around 270 participants will be enrolled in the study in approximately 15 sites in Canada.Participants will receive venetoclax tablets to be taken by mouth daily according to the approved local label. The duration of the study is approximately 36 months. There is expected to be no additional burden for participants in this trial. All study visits will occur during routine clinical practice and participants will be followed for 36 months.
Critères d'éligibilité	• Diagnosis of Acute Myeloid Leukemia (AML). • Ineligible for intensive chemotherapy, determined by the physician's assessment of age, Eastern Cooperative Oncology Group Performance Status (ECOG-PS), comorbidities, regional guidelines, and institutional practice. • Participant for whom the physician has decided to initiate venetoclax treatment in accordance with the local label. The decision to treat with venetoclax is made by the physician prior to any decision to approach the participant to participate in this study.
Critères d'exclusion	• Participation in an interventional clinical trial within 30 days prior to venetoclax treatment initiation.