Essai Clinique

Généré le 28 avr. 2024 à partir de

Titre	Registre de l'hémoglobinurie paroxystique nocturne (HPN)
Protocole ID	Registre PNH (M07-001)
ClinicalTrials.gov ID	NCT01374360
Type(s) de cancer	Syndrome myélodysplasique
Type étude	Autre
Institution	CIUSSS DE L'EST-DE-L'ILE-DE-MONTREAL PAV. MAISONNEUVE/PAV. MARCEL-LAMOUREUX 5415 boul. de l'Assomption, Montréal, QC, H1T2M4
Ville	Montréal
Investigateur principal	Dr Thomas KIss
Coordonnateur	Johanne Blais 514-252-3400 poste 3295
Statut	Fermé
But étude	This study is a collection of data to evaluate safety and characterize progression of Paroxysmal Nocturnal Hemoglobinuria (PNH).
Critères d'éligibilité	• Patients of any age, including minors, with a diagnosis of PNH or a detected PNH clone, including patients previously treated with Soliris and withdrawn from treatment. (Subjects under the age of eighteen years must have parent/legal guardian consent. Upon turning eighteen years of age, these subjects must be re-consented). • Ability to comprehend and sign consent to have data entered in the PNH Registry.
Critères d'exclusion	