

Essai Clinique Généré le 09 mai 2025 à partir de

Titre	Phase IV Observational Study to Collect Safety and Outcome Data in Real-World Setting in Adult Patients With Extensive Stage Small Cell Lung Cancer (SCLC) Receiving Zepzelca
Protocole ID	EMERGE 402 (JZP712-402)
ClinicalTrials.gov ID	<u>NCT04894591</u>
Type(s) de cancer	Poumon à petites cellules
Phase	Phase IV
Stade	Maladie avancée ou métastatique
Type étude	Autre
Médicament	Zepzelca
Institution	CISSS DE CHAUDIERE-APPALACHES H HOTEL-DIEU DE LEVIS 143 rue Wolfe, Lévis, QC, G6V 3Z1
Ville	
Investigateur principal	Dre Danièle Marceau
Coordonnateur	Pierre Bédard 418-835-7121
Statut	Fermé
But étude	To assess the effectiveness and safety of Zepzelca in adult participants with extensive stage small cell lung cancer (SCLC) in real-world clinical practice.
Critères d'éligibilité	 Patient has initiated or will be receiving Zepzelca treatment in line with the local Zepzelca prescribing information. Decision to initiate treatment with Zepzelca was made as per investigator's routine treatment practice prior to enrollment in the study. Patient, or a legally acceptable representative, signed the informed consent before any study-related activities are undertaken.
Critères d'exclusion	 Patients who discontinued a prior Zepzelca treatment due to adverse events. Patient who received more than 2 cycles of Zepzelca treatment in their current treatment schedule. Patient received treatment with any investigational agent within 30 days prior to first Zepzelca infusion or plans to use another investigational agent while receiving Zepzelca.