

# Essai Clinique

Généré le 07 mai 2024 à partir de

Titre	ROMAN : Réduction de la mucosite buccale avec le Avasopasem Manganese (GC4419) - Étude de phase 3 chez les patients recevant de la chimioradiothérapie dans le traitement du cancer de la tête et du cou localement avancé et non métastatique
Protocole ID	GTI-4419-301
ClinicalTrials.gov ID	<a href="#">NCT03689712</a>
Type(s) de cancer	ORL
Phase	Phase III
Stade	Localement avancé
Type étude	Support
Médicament	Superoxide Dismutase Mimetic GC4419
Institution	<p>CIUSSS DE LA MAURICIE-ET-DU-CENTRE-DU-QUEBEC  <b>H</b> CHAUR  1991 Boulevard du Carmel, Trois-Rivières, QC, G8Z 3R9</p>
Ville	Trois-Rivières
Investigateur principal	Dr François Vincent
Coordonnateur	Marie-Ève Caron 819-697-3333 poste 63238
Statut	Fermé
But étude	The purpose of the phase 3, clinical study is to determine if GC4419 administered prior to intensity-modulated radiation therapy (IMRT) reduces the severity of radiation induced oral mucositis in patients who have been diagnosed with locally advanced, non-metastatic squamous cell carcinoma of the head and neck.
Critères d'éligibilité	<ul style="list-style-type: none"> <li>• squamous cell carcinoma of the head and neck</li> <li>• treatment plan to receive IMRT delivered as single daily fractions of 2.0 to 2.2 Gy with a cumulative radiation dose of 60-72 Gy</li> <li>• Treatment plan to receive standard cisplatin monotherapy</li> <li>• Age 18 years or older</li> <li>• Eastern Cooperative Oncology Group (ECOG) performance status ≤ 2</li> <li>• Adequate hematologic, renal and liver function</li> <li>• Negative serum pregnancy test</li> <li>• Use of effective contraception</li> </ul>
Critères d'exclusion	<ul style="list-style-type: none"> <li>• Tumor of the lips, larynx, hypopharynx, nasopharynx, sinuses, or salivary glands</li> <li>• Metastatic disease</li> <li>• Prior radiotherapy to the region of the study cancer or adjacent anatomical</li> <li>• Prior induction chemotherapy</li> <li>• Receiving any approved or investigational anti-cancer agent other than those provided for in this study</li> <li>• Concurrent participation in another interventional clinical study</li> <li>• Inability to eat soft solid food at baseline</li> <li>• Malignant tumors other than HNC within the last 5 years</li> <li>• Active infectious disease excluding oral candidiasis</li> <li>• Presence of oral mucositis at baseline</li> <li>• Known history of HIV or active hepatitis B/C</li> </ul>

- Female patients who are pregnant or breastfeeding
- Known allergies or intolerance to cisplatin and similar platinum-containing compounds
- Requirement for concurrent treatment with nitrates or other drugs that may create a risk for a precipitous decrease in blood pressure