

# Medication Administration Protocol: LETAIRIS

## Document Control Information

Document ID:	MED-PROT-5262
Effective Date:	
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Therapeutic Class:	Adverse Event Report
Risk Level:	Medium

## Standard Dosing Protocol

**Current Standard Dose:** Standard dosing per label

**Monitoring Requirements:** Standard warnings

## Administration Guidelines

**Route of Administration:** IM

**Frequency:** Twice daily

**Duration:** 7 days

**Patient Population:** Patients similar to case: 2, age 74

## Contraindications and Precautions

**Contraindications:** Enhanced screening for risk factors

**Adverse Events:** Decreased appetite, Fatigue, Abdominal pain

# Detailed Clinical Considerations

## Patient Monitoring Protocol

Regular monitoring is essential for safe administration of LETAIRIS. Current protocols require Standard warnings. Healthcare providers should assess patient response and adjust therapy accordingly. Baseline laboratory values should be obtained prior to initiation. The mechanism of action involves LETAIRIS mechanism with identified safety concern. Clinical efficacy has been demonstrated in Decreased appetite, Fatigue, Abdominal pain.

## Special Populations

**Elderly Patients:** Dose adjustment may be required based on renal function and overall health status.

**Pediatric Patients:** Safety and efficacy not established in patients under 18 years.

**Pregnancy:** Use only if potential benefit justifies potential risk to fetus.

**Nursing Mothers:** Exercise caution when administering to nursing women.

*Last revised: April 2024*