LoW Dose-Intensity vs. Standard Dose-Intensity COntinuous Renal ReplaceMent Therapy in Critically Ill Patients (WISDOM)

NCT06446739

Status	RECRUITING
Phase	Not Applicable
Sponsor	University of Alberta
Enrollment	100 participants

Key Eligibility Criteria

Inclusion (5)

- age e 18 years
- weight e 55 kg
- plan to initiate CRRT or within 24 hours of having started CRRT for AKI
- expected to survive and receive CRRT for a duration of e 48 hours
- able to provide informed consent or have an authorized representative provide consent after being informed on the details and risks of the trial unless a deferred consent process is approved by local Research Ethics Board (REB).

Exclusion (4)

- indication for sustained higher dose-intensity CRRT as designated by the attending clinicians
- end-stage kidney disease receiving maintenance dialysis
- receipt of any RRT for AKI during the current hospitalization
- inability to comply with the requirements of the study protocol.

Locations (7 total)

Mayo Clinic, Rochester, Minnesota, United States
University of Alberta Hospital, Edmonton, Alberta, Canada
Sturgeon Community Hospital, St. Albert, Alberta, Canada
... and 4 more locations