ARDS in Children and ECMO Initiation Strategies Impact on Neurodevelopment (ASCEND)

Status: RECRUITING

Eligibility Criteria

Age: 14 days to 20 years old

This study is NOT accepting healthy

Healthy Volunteers: volunteers

Inclusion Criteria:

* Time between intubation and ECMO cannulation is less than 240 hours (10 days) * ECMO support type is respiratory (VV or VA cannulation) * Chest radiograph with bilateral lung disease * Moderate or severe pediatric ARDS as measured by oxygenation index or oxygen saturation index after intubation and prior to ECMO cannulation: One $OI \ge 16$ or Two $OIs \ge 12$ and ≤ 16 at least four hours apart or Two $OSIs \ge 10$ at least four hours apart

Exclusion Criteria:

* Previously enrolled in PROSpect * Perinatal related lung disease * Congenital diaphragmatic hernia or congenital/acquired diaphragm paralysis * Respiratory failure caused by cardiac failure or fluid overload * Cyanotic congenital heart disease * Cardiomyopathy * Primary pulmonary hypertension (PAH) * Unilateral lung disease * Intubated for status asthmaticus * Obstructive airway disease * Bronchiolitis obliterans * Post hematopoietic stem cell transplant * Post lung transplant * Home ventilator dependent * Neuromuscular respiratory failure * Head trauma: (managed with hyperventilation) * Intracranial bleeding * Unstable spine, femur or pelvic fractures * Acute abdominal process/open abdomen * Family/medical team have decided to not provide full support * Enrolled in interventional clinical trial: not approved for co-enrollment; does not include cancer protocols. * Known pregnancy

Conditions & Interventions

Interventions:

DEVICE: ECMO support, OTHER: PROSpect protocolized therapies

Conditions:

Acute Respiratory Distress Syndrome, Extracorporeal Membrane Oxygenation

Keywords:

ARDS, Acute Respiratory Distress Syndrome, Extracorporeal Membrane Oxygenation, ECMO, Extracorporeal Life Support, ECLS, Pediatric, Quality of Life, Functional Status

More Information

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Phase: IRB Number:

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