Individualized Fortification of Human Milk for Infants Born d1250 g (MaxiMoM-InForM)

NCT05308134

Status	RECRUITING
Phase	Not Applicable
Sponsor	The Hospital for Sick Children
Enrollment	615 participants

Key Eligibility Criteria

Inclusion (3)

- d1250g birth weight or Gestational Age \<30+0 weeks and \<1500 g birth weight.
- Parental/guardian consent to participate.
- Consent for the use of pasteurized donor milk if mother's milk is not available.

Exclusion (5)

- Infant received fortifier or formula before Study Day 1.
- Study Day 1 anticipated to occur after postnatal day 21.
- Infants with congenital or chromosomal anomalies or brain injury that may affect growth or neurodevelopment.
- Enrollment in any other clinical study affecting nutritional management during the feeding intervention.
- Reasonable potential that the infant will be transferred to a NICU where the study protocol will not be continued before they have completed at least 4 weeks of the feeding intervention.

Locations (20 total)

University of Alberta, Edmonton, Alberta, Canada
William Osler Health System-Brampton Civic Hospital, Brampton, Ontario, Canada
William Osler Health System-Etobicoke General Hospital, Etobicoke, Ontario, Canada
... and 17 more locations