

Feeding Tolerance and Growth of Preterm Infants Consuming a Supplement Containing Two Human Milk Oligosaccharides (HMOs)

NCT06212427

Status	RECRUITING
Phase	Not Applicable
Sponsor	Société des Produits Nestlé (SPN)
Enrollment	188 participants

Key Eligibility Criteria

Inclusion (6)

- Written informed consent has been obtained from at least one parent (or other legally acceptable representative \[LAR\], if applicable)
- Infant's parent(s)/LAR is of legal age of majority, has parental authority, must understand the consent form and other relevant study documents, and is willing and able to fulfil the requirements of the study protocol
- Infant gestational age is \geq 34 weeks as determined by the first day of the mother's last menstrual period or by fetal ultrasound
- Infant birth weight \geq 2500g
- Infant postnatal age \geq 14 days
- ... and 1 more (see full listing online)

Exclusion (8)

- Infant is clinically unstable, for example:
- Infant has hemodynamic instability as evidenced by clinical signs of sepsis, hypotension (MAP $<$ 5th percentile for age for at least three hours), or is receiving vasopressor drugs
- Infant has received an exchange transfusion within the past 48 hours
- Infant has had an episode of severe asphyxia at birth (PH less than 7.0)
- Infant has signs of necrotizing enterocolitis according to modified Bell staging criteria (stage IIA or higher)
- ... and 3 more (see full listing online)

Locations (6 total)

Kepler Universitätsklinikum Linz, Linz, Austria
Evangelisches Waldkrankenhaus Spandau, Berlin-Spandau, Germany
Kinderklinik Darmstadt, Darmstadt, Germany
... and 3 more locations

<https://clinicaltrials.gov/study/NCT06212427>

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