Pilot Trial Investigating Every Other Day Dosing of Oral Iron in Premature Infants (IQONic)

NCT06555315

Status	RECRUITING
Phase	Not Applicable
Sponsor	CHRISTUS Health
Enrollment	100 participants

Key Eligibility Criteria

Inclusion (5)

- Children (Minor \< 18 years of age)
- Neonates
- Hospitalized
- Premature infants who are on full enteral feeds and are started on oral iron
- Premature infants who completed 26 0/7 to 32 6/7 weeks' gestation at birth

Exclusion (1)

- • Infants with known congenital anomalies or chromosomal abnormalities (such as Trisomy 18 or Trisomy 21), conditions that affect iron metabolism (such as thalassemia or hemochromatosis), bleeding disorders or coagulopathy, and received iron parenterally prior to randomization

Locations (1 total)

CHRISTUS Children's, San Antonio, Texas, United States