

IV Iron-induced Hypophosphatemia After RYGB

NCT06350955

Status	RECRUITING
Phase	Phase 4
Sponsor	Lucie Favre
Enrollment	94 participants

Key Eligibility Criteria

Inclusion (8)

- Patients over 18 years
- Signed informed consent
- Patients with previous RYGB surgery performed \geq 12 months ago
- Failed response to oral iron supplementation
- Established diagnosis of iron deficiency by ferritin $<$ 50 ug/l or serum ferritin \geq 100ug/l and low transferrin saturation (TSAT) \leq 30%
... and 3 more (see full listing online)

Exclusion (16)

- Patients with known hypersensitivity to iron preparation and/or anaphylaxis from any cause
- Patients for whom a treatment with one of the IV iron is contra-indicated (based on product summary of product characteristics)
- Women who are pregnant or breastfeeding
- Intention to become pregnant during the course of the study
- Renal failure, chronic kidney disease stage 3b or worse (eGFR \leq 45 ml/min/1.73m²)
... and 11 more (see full listing online)

Locations (1 total)

CHUV, Lausanne, Canton of Vaud, Switzerland