Stryker's Surgeon iD Mandible Reconstruction Plates: A Retrospective Post Market Follow Up

NCT07221916

Status	RECRUITING
Sponsor	Stryker Craniomaxillofacial
Enrollment	50 participants

Key Eligibility Criteria

Inclusion (2)

- Patients who underwent primary or secondary mandibular reconstruction using a Stryker Surgeon iD Plate implant as per routine clinical practice.
- Patients for whom data on the primary outcome variable is available.

Exclusion (4)

- Patients with active local infections at the time of surgery.
- Patients with known metal allergies and/or foreign body sensitivity at the time of surgery.
- Potentially non-compliant patients who were unwilling or incapable of following post-operative care instructions.
- Patients with inadequate bone quantity or quality necessary for plate fixation or stabilization at the time of surgery

Locations (2 total)

UCSF Otolaryngology Head and Neck Surgery, San Francisco, California, United States
Heinrich-Braun-Klinikum gGmbH, Zwickau, Germany