Stryker Universal Midface and Upper-Face Fixation System: A Retrospective Post Market Follow Up

NCT07245719

Status RECRUITING
Sponsor Stryker Craniomaxillofacial
Enrollment 120 participants

Key Eligibility Criteria

Inclusion (4)

- Skeletally mature patients at the time of surgery.
- Patients who underwent a craniomaxillofacial procedure using implants of the Upper-Face and/or Mid-Face modules (1.2 / 1.7 modules) of the Stryker Universal CMF System as per routine clinical practice.
- Patients for whom data on the primary outcome variable is available.
- Patients with data available from follow-up visits.

Exclusion (5)

- Patients with active infections at the time of surgery.
- Patients with known metal allergies and/or foreign body sensitivity at the time of surgery.
- Subjects with non-reducible and unstable fractures (except reconstruction plates) at the time of surgery that were treated with the subject devices.
- Patients who underwent secondary reconstructions with non-secondary reconstruction plates
- Patients with limited blood supply or insufficient quality or quantity of bone at the time of surgery.

Locations (1 total)

Weill Cornell Medicine Oral and Maxillofacial Surgery 525 East 68th Street, F-2132, New York, New York, United States