Bone Substitutes Outcomes - Post Market Follow-up

NCT06374342

Status	RECRUITING
Sponsor	Teknimed
Enrollment	425 participants

Key Eligibility Criteria

Inclusion (5)

- Be 18 years or older.
- Be willing to sign an informed consent approved by IRB or EC (where applicable) or not being opposed to the use of their clinical data in the study (France) and
- Be considered for a surgery where bone filling with one of the TEKNIMED bone substitutes comprised in this study is needed and intended to be used according to the IFU.
- Have undergone a surgery with a TEKNIMED bone substitute used according to the IFU, between the 1st January 2015 and the date of the site initiation visit
- Be informed of the study and not being opposed to the use of their clinical data in the study (France) or be willing to sign an informed consent (where applicable) during the first follow-up visit following the site initiation.

Exclusion (11)

- Patients presenting one of the following conditions will not be included:
- Under trusteeship or guardianship
- Pregnancy or breast-feeding women
- According to contraindications per IFU:
- Procedures other than those stated in the INDICATIONS section

... and 6 more (see full listing online)

Locations (6 total)

Pôle Rachis Hôpital Privé d'Eure et Loir, Mainvilliers, Eure et Loir, France
Hôpital Joseph Ducuing, Toulouse, Haute Garonne, France
Clinique Médipole Garonne, Toulouse, Haute Garonne, France
... and 3 more locations

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