Hand & Wrist Pyrocarbon Implants Outcomes Clinical Study

NCT04137237

Status	RECRUITING
Sponsor	Stryker Trauma and Extremities
Enrollment	417 participants

Key Eligibility Criteria

Inclusion (5)

- years or older at the time he/she receives the information and signs the informed consent (when applicable as per local regulatory requirements),
- Informed on the study and willing to sign an informed consent form approved by Institutional Review Board or Ethics Committee,
- Willing and able to comply with the requirements of the study protocol,
- Follow-up visits (at least the last two) must be prospective,
- Patient must have complete information available for each completed visit

Exclusion (3)

- Patient pertaining to one of the categories referred to as "vulnerable population" in the French Law (articles L. 1121-5 to L. 1121-8 of the French Public Health Code), or as "particularly vulnerable persons" in the Swiss Federal Law (Chapter 3, Sections 1 to 4 of CC 810.30 Federal Act on Research involving Human Beings - Human Research Act, HRA),
- Patient unable to comply with the study procedures based on the judgment of the investigator (e.g. cannot comprehend study questions, inability to keep scheduled assessment times),
- Any medical condition that could impact on the study outcomes functional signification at the investigator's discretion (e.g., neuropathy, allergy…)

Locations (4 total)

Clinique du Parc, Lyon, France
Espace Médical Vauban, Paris, France
Institut de la Main Nantes-Atlantique - Pôle Santé-Atlantique, Saint-Herblain, France
... and 1 more locations