

Multigen Plus CCK and AMF TT Cones Follow Up Study

NCT05314491

Status	RECRUITING
Sponsor	Limacorporate S.p.a
Enrollment	68 participants

Key Eligibility Criteria

Inclusion (5)

- Male or female.
- Age e 18 years old.
- Given written informed consent approved by the reference Ethics Committee (EC).
- Subjects in whom a decision has already been made to perform a Total Knee Arthroplasty with Multigen Plus system as per Indication For Use. The decision to implant a Multigen Plus CCK system, alone or involving the AMF TT cones, must be taken prior to, and independently from, the decision to enrol the subject. This decision should be made in accordance with routine clinical practice at the study site concerned.
- Subjects able to comply with the Study Protocol.

Exclusion (5)

- Age < 18 years old.
- Subjects with any Multigen Plus system contraindication for use, or any AMF TT cones contraindication for use when used in combination with the Multigen Plus CCK, as reported in the current Instruction For Use.
- Any clinically significant pathology based on the medical history that the Investigator feels may affect the study evaluation.
- Female subjects who are pregnant, nursing, or planning a pregnancy.
- Previous knee replacement on the contralateral side within the last year and whose outcome is achieving an KSS < 70 points.

Locations (3 total)

Hospital Senhora da Oliveira in Guimarães, Guimarães, Portugal
Saints Cyril and Methodius Hospital, Bratislava, Slovakia
Royal Devon and Exeter Hospital, Exeter, United Kingdom