

Evaluation of the Vivo Isar Stent System in Routine Clinical Practice

NCT06420505

Status	RECRUITING
Sponsor	Translumina GmbH
Enrollment	3,000 participants

Key Eligibility Criteria

Inclusion (6)

- e18years old
 - Ability to provide signed informed consent form.
 - Male or non-pregnant female patient (Pregnancy status to be confirmed verbally from the female patient of reproductive age)
 - Presentation with acute coronary syndrome or chronic coronary syndrome with stable angina or angina equivalent symptoms or with a positive noninvasive test for ischemia or evidence of a functionally significant coronary stenosis.
 - Patient having social security number.
- ... and 1 more (see full listing online)

Exclusion (7)

- Concurrent participation in another clinical trial.
 - Having benefited from an angioplasty of d 1 month with a stent other than Vivo ISAR.
 - Planned elective surgery in next 6 months
 - Cardiogenic shock/ hemodynamically unstable patients
 - Concurrent medical condition with a life expectancy of less than 12 months
- ... and 2 more (see full listing online)

Locations (19 total)

Clinique Axiom, Aix-en-Provence, France
CHU CAEN, Caen, France
Hôpital A. Schweitzer - GHCA, Colmar, France
... and 16 more locations