Development of a Device for Evaluating Primary Hemostasis Under Whole Blood Flow Conditions

NCT03773159

Status	RECRUITING
Sponsor	Centre Hospitalier Universitaire Dijon
Enrollment	200 participants

Key Eligibility Criteria

Inclusion (5)

- person who has given oral consent
- adult
- blood donor at EFS Bourgogne Franche-Comté
- or patient with von Willebrand disease or major constitutional thrombopathy followed by the Resource and Competence Centre (CRC) - Constitutional Hemorrhagic Diseases of Dijon or Besançon
- or patient on antiplatelet drugs consulting for thrombosis at the Dijon Bourgogne or Besançon Hospital

Exclusion (5)

- a person who is not affiliated to or not a beneficiary of national health insurance
- person subject to court-ordered protection (curatorship, guardianship)
- pregnant, parturient or breastfeeding woman
- a person who is unable to consent
- person on anti-inflammatory treatment and serotonin reuptake inhibitor antidepressants (platelet function disorders)

Locations (2 total)

CHU de Besançon, Besançon, France
CHU Dijon Bourgogne, Dijon, France