A Study of ETHIZIA Versus SURGICEL Original in Controlling Soft Tissue Bleeding During Open Surgery

NCT06664788

Status	RECRUITING
Phase	Not Applicable
Sponsor	Ethicon, Inc.
Enrollment	108 participants

Key Eligibility Criteria

Inclusion (8)

- Pre-operative
- Participant is scheduled to undergo an elective open, abdominal, retroperitoneal, pelvic, thoracic (non-cardiac) or extremity surgical procedure
- Participant is willing and able to give written informed consent for the clinical investigation participation
- Intra-operative
- Participant in whom the Investigator can identify and visualize a target bleeding site for which any applicable conventional means for hemostasis (e.g., suture, ligature, or cautery) are ineffective or impractical

... and 3 more (see full listing online)

Exclusion (8)

- Pre-operative
- Participant is scheduled for another planned surgery within the follow-up period and the subsequent surgery would jeopardize the ETHIZIA or SURGICEL Original application
- Participant is taking multiple antithrombotic therapies in therapeutic dosage up to the time of surgery, but allowing exclusive use of acetylsalicylic acid
- Participant has an active or suspected infection at the bleeding site
- Participant is pregnant, planning on becoming pregnant, or actively breastfeeding during the 28-day follow-up period

... and 3 more (see full listing online)

Locations (8 total)

Keck Hospital of USC, Los Angeles, California, United States
Washington University Barnes Jewish Hospital, St Louis, Missouri, United States
Capital Health Medical Center - Hopewell, Pennington, New Jersey, United States
... and 5 more locations