

qDSA Blood Flow Measurement in Patients Undergoing TAE of the Liver

NCT06971991

Status	RECRUITING
Phase	Not Applicable
Sponsor	University of Wisconsin, Madison
Enrollment	20 participants

Key Eligibility Criteria

Inclusion (1)

- Give voluntary, written informed consent to participate in this study and willing to comply with study-related evaluation and procedure schedule

Exclusion (5)

- Pregnancy or breastfeeding
- Patients with an acute kidney injury or stage IV or V chronic kidney disease (Cr greater than 2.4 or estimated glomerular filtration rate (eGFR) less than 30), unless anuric and on dialysis without expected return of renal function.
- Patients with an iodinated contrast allergy who cannot be adequately premedicated to receive contrast as part of the embolization procedure.
- Patients with a physical or psychological condition that would impair study participation.
- The patient is judged unsuitable for study participation by the Investigator for any other reason

Locations (1 total)

UW Hospital and Clinics, Madison, Wisconsin, United States