

Study to Compare the Clinical Effectiveness and Cost Utility of an All-in-one Procedure With an At-home Screening Trial

NCT06996574

Status	RECRUITING
Phase	Not Applicable
Sponsor	Academisch Medisch Centrum - Universiteit van Amsterdam (AMC-UvA)
Enrollment	180 participants

Key Eligibility Criteria

Inclusion (4)

- Diagnosed with chronic, neuropathic pain with an approved aetiology in the Netherlands PSPS (persistent spinal pain syndrome) PDN (painful diabetic neuropathy) CRPS (complex regional pain syndrome) SFN (small fibre neuropathy)
- Routinely selected for CL-SCS
- Be willing to complete additional trial related questionnaires
- Be willing and capable of giving informed consent.

Exclusion (4)

- Drug abuse Pregnancy Coagulation disorder / Use of anticoagulation therapy which cannot be stopped temporarily Active infection Life expectation <12 months Intake questionnaires "PROMS neuromodulatie" not completed by the patient
- Have an existing drug pump and/or SCS system or another active implantable device such as a pacemaker, deep brain stimulator (DBS), or sacral nerve stimulator (SNS)
- Current or previous treatment with an implanted pain reduction device
- Participation in another clinical trial that may confound the results of this trial

Locations (1 total)

Amsterdam UMC, Amsterdam, Netherlands