Perineural Incobotulinumtoxin-A for Complex Regional Pain Syndrome - An Open-label Feasibility Study

NCT07473635

Status	RECRUITING
Phase	Not Applicable
Sponsor	Bo Biering-Soerensen
Enrollment	20 participants

Key Eligibility Criteria

Inclusion (7)

- Are over the age of 18
- Have a diagnosis of CRPS type 1 or 2 in either one upper or one lower extremity which fulfils the Budapest research criteria
- Have had the condition for at least 6 months
- Rate CRPS as their primary pain condition
- Have been on a stable analgesic regimen, including any rescue medications, for at least 1 month prior to the study and intend to maintain this regimen throughout the study

... and 2 more (see full listing online)

Exclusion (11)

- Are allergic to botulinum toxin A
- Have been treated with botulinum toxin A for any indication within 3 months prior to study start
- Are diagnosed with myasthenia, Lambert-Eaton syndrome, amyotrophic lateral sclerosis, or any other condition which makes differentiation of CRPS-specific pain difficult
- Have an ongoing infection in the affected limb
- Do not intend to start physical therapy, psychotherapy, or any other non-pharmaceutical intervention aimed at reducing pain

... and 6 more (see full listing online)

Locations (1 total)

CRPS- and Nerve Pain Clinic, Rigshospitalet Glostrup, Glostrup, Region Sjælland, Denmark