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Title

A randomised feasibility study of serial magnetic resonance imaging to reduce treatment times in Charcot neuroarthropathy in people with diabetes (CADOM): A protocol.

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Abstract

Background

Charcot neuroarthropathy is a complication of peripheral neuropathy associated with diabetes which most frequently affects the lower limb. It can cause fractures and dislocations within the foot, which may progress to deformity and ulceration. Recommended treatment is immobilisation and offloading, with a below knee non-removable cast or boot. Duration of treatment varies from six months to more than one year. Small observational studies suggest that repeated assessment with Magnetic

Resonance Imaging improves decision making about when to stop treatment, but this has not been tested in clinical trials. This study aims to explore the feasibility of using serial Magnetic Resonance Imaging without contrast in the monitoring of Charcot neuroarthropathy to reduce duration of immobilisation of the foot. A nested qualitative study aims to explore participants' lived experience of Charcot neuroarthropathy and of taking part in the feasibility study.

Methods

We will undertake a two arm, open study, and randomise 60 people with a suspected or confirmed diagnosis of Charcot neuroarthropathy from five NHS, secondary care multidisciplinary Diabetic Foot Clinics across England. Participants will be randomised 1:1 to receive Magnetic Resonance Imaging at baseline and remission up to 12 months, with repeated foot temperature measurements and x-rays (standard care plus), or standard care plus with additional three-monthly Magnetic Resonance Imaging until remission up to 12 months (intervention). Time to confirmed remission of Charcot neuroarthropathy with off-loading treatment (days) and its variance will be used to inform sample size in a full-scale trial. We will look for opportunities to improve the protocols for monitoring techniques and the clinical, patient centred, and health economic measures used in a future study. For the nested qualitative study, we will invite a purposive sample of 10-14 people able to offer maximally varying experiences from the feasibility study to take part in semi-structured interviews to be analysed using thematic analysis.

Discussion

The study will inform the decision whether to proceed to a full-scale trial. It will also allow deeper understanding of the lived experience of Charcot neuroarthropathy, and factors that contribute to engagement in management and contribute to the development of more effective patient centred strategies.

Trial registration ISRCTN, ISRCTN, 74101606. Registered on 6 November 2017, http://www.isrctn.com/ISRCTN74101606?q=CADom&filters=&sort=&offset=1&totalResults=1&page=1&pageSize=10&searchType=basic-search

Keywords Charcot neuroarthropathy, diabetes, MRI, temperature monitoring, X-ray, patient experience, feasibility study.

Background

Charcot neuroarthropathy (CN) is a complication of peripheral neuropathy associated with diabetes which most frequently affects the lower limb. It can cause fractures and dislocations within the foot, which may progress to deformity and ulceration. The symptoms include redness, warmth and swelling in the foot and/or leg. This inflammation can lead to fractures in the bones and can damage joints, affecting the shape and function of the foot. It was first described 140 years ago (1), however it remains a poorly understood and frequently overlooked complication of diabetes (2).

Population-based studies have estimated a life time cumulative incidence for CN of 0.4% to 1.3% in people with diabetes, rising to 13% in people at high risk who attend diabetic foot speciality clinics (3). In 2018 a regional survey of 205,033 people with diabetes in the East Midlands, UK reported a point prevalence of 0.04% (4). CN is associated with increased length of stay and use of medical resources (5).

The aim of treatment is to stop the inflammatory process, relieve pain and maintain foot architecture reducing the risk of future ulceration and amputation (6). The current international consensus is that the foot should be immobilised in a below knee non-removable cast or boot, with weekly or fortnightly review by healthcare professionals working in specialist multidisciplinary diabetic foot clinics (7). The immobilisation minimises the potential for any further damage to the foot structure. Immobilisation is continued until

remission, defined as the absence of clinical signs of inflammation, measured using skin surface infra-red thermography, and X-rays showing signs of bone healing and union (8).

The evidence base for the treatment of CN is weak. It is based on studies from a few centres which used retrospective designs and case note review methods using small sample sizes, typically in the range of 9-55 participants (3,9–13). Many studies failed to standardise monitoring, treatment and outcomes, which makes direct comparison between studies difficult.

Studies from the UK have shown a median time to remission of 9-12 months (9,13,14). However, US studies report considerably shorter time to remission of 3-5 months (3,10–12). Studies from Brazil and Germany show remission times of 3-12 months and 3-6 months, respectively (15,16). Shorter treatment times could be related to reported differences in the relapse rates for CN, between 12-33% (13,17–19), but without clear and consistent definitions for remission and relapse this is unknown. There is also variation in the reported annual major amputation rates in people with CN from two different case series from hospitals in the USA: 2.7% and 6.6% (20,21)

The reasons for the variation are not understood but could include people's characteristics at the start of the treatment, different techniques for monitoring CN, different protocols for the same monitoring techniques, variations in approach to off-loading, and variability in study design. These could either underestimate or overestimate treatment duration.

Temperature difference between the feet is one of the most frequently used methods to monitor CN. It is recommended in the 2015 National Institute for Health and Care Excellence guidance on diabetic foot problems (22). The most recent systematic review (8) published in 2013 recommends that immobilisation is continued until the temperature difference between the feet is less than 1-2 °C, and no further radiological changes on imaging have occurred. However this recommendation is only based on level IV evidence, i.e. case series (8). There is variability in the protocols used to measure the temperature difference between the feet. The most detailed protocol for measuring temperature discrepancy requires a 15 minute acclimatisation period, controlled ambient air temperature, and readings collected from nine different places on each foot (23). In addition, plain X-rays demonstrate damage to the bone and joints rather than disease activity (inflammation).

Studies show inconsistency in the methods for monitoring and monitoring devices used (13,17–19,23–25). These factors may overestimate or underestimate the degree of inflammation, so treatment may be discontinued too early or continued for longer than necessary. The presence of simultaneous bilateral foot disease or the absence of a contralateral limb through prior amputation invalidates the use of temperature measurement as a tool for identifying disease remission.

The National Institute for Health and Care Excellence recommends the use of MRI in determining a diagnosis of CN in the early stages of disease when no signs are evident on

plain radiology (30). However serial MRI is not widely used in routine clinical practice as a tool to monitor for signs of disease remission in CN (27). One prospective study using MRI with contrast reported that mean healing times were associated with contrast uptake assessed at baseline (28). A further two retrospective studies looked at bone marrow oedema. One study reported decreasing bone marrow oedema in 69% of follow up images (29) and the second study found a significant positive correlation between intensity of bone marrow oedema on MRI and clinical measures (30). This emerging evidence suggests that MRI may be useful for the surveillance of active CN. The findings from MRIs could be adopted as the criterion standard for establishing disease activity and remission.

The use of MRI in monitoring CN therefore needs to be formally evaluated in a trial (29). However, the evidence to support a full randomised controlled trial is presently insufficient. We will conduct a randomised feasibility study to understand the proportion of people who meet the eligibility criteria, the number of eligible participants recruited, the number of participants who receive an alternative diagnosis, and the proportion of participants who withdraw. Time to MRI confirmed remission of CN with off-loading treatment (in days) and its variance will be used to inform sample size in a main trial. We will look for opportunities to improve the protocols for monitoring techniques in a future trial. We will examine the feasibility of a range of clinical, patient centred, and health economic measures We are using a randomised controlled trial as it is considered the gold standard for evaluating efficacy in clinical research (31).

As part of the feasibility study we will carry out a qualitative study to further the understanding of people's experiences of living with CN and the factors that contribute to people's engagement in their treatment. Previous qualitative studies have demonstrated the importance of people's perspectives in order to promote engagement in the prevention and management of diabetic foot ulcerations (32–34). What may be people's views and experiences of CN is an under-researched area (35). In the UK treatment times for CN are between 9-12 months (14), which is longer than those for foot ulceration, where treatment times are no more than 12 weeks for half of the people (36). This means that evidence on people's experiences of foot ulceration may not transfer to CN.

In summary, there is a lack of evidence to support the use of monitoring techniques in CN. Healthcare professionals rely on methods and devices which do not accurately reflect disease progression, and decision making about discontinuing or prolonging immobilisation is challenging. A lack of understanding on people's experiences of living with CN, means their needs and wishes may be neglected with current treatments, and are not being considered when developing new treatment strategies and pathways.

Aim and objectives

This study aims to explore the feasibility of using serial MRI without contrast in the monitoring of CN to reduce duration of immobilisation of the foot, in order to decide whether a large-scale trial is warranted. We will assess eligibility, recruitment, retention and withdrawal rates. Time to MRI confirmed remission of CN with off-loading treatment (days)

and its variance will be used to inform sample size in a main trial. We will also examine the feasibility of collecting clinical, patient centred and health economic measures. The nested qualitative study aims to explore the dimensions of lived experience of CN and the participants' experiences of taking part in the feasibility study.

Methods

Study Design **(Figure 1)**

This is a two-arm, open, randomised controlled trial, investigating the feasibility of using serial MRI to monitor CN. The study will last for a maximum of 3 ½ years. The study is divided into two phases. Phase one, the active phase, will last until the CN is in remission, or a maximum of 12 months. Phase two, the follow-up phase, will last for six months after remission (Figure 1). The maximum time a participant will be in the trial is 18 months.

The decision to use an open label design was pragmatic: the MRIs will be reported by radiologists and interpreted by the healthcare professionals working in multidisciplinary specialist diabetic foot clinics. As the reporting of MRIs relies on comparison to previous images, this will indicate the trial arm the participant has been randomised to.

The trial has been reviewed and approved by East Midlands - Derby Research Ethics Committee, 04/10/2017, ref: 17/EM/0288.

Setting

The setting will be multidisciplinary specialist diabetic foot services at five NHS Hospital Trusts in England.

Randomisation

A randomisation scheme has been generated by the trial statistician. Allocation will be stratified by centre. Participants will be randomised using a web-based randomisation process on a 1:1 basis to: (a) Immobilisation discontinued on the basis of clinical remission determined by skin temperature measurement, which triggers an MRI **(standard care plus)** or (b) Standard care plus and additionally the serial use of MRI at 3, 6, 9 and 12 months to identify disease remission and thus discontinuation of immobilisation **(intervention)**.

Sample size

As this is a feasibility study a power calculation is not required. An allowance has been made for up to 10-15% of participants to be withdrawn from the study due to an alternative diagnosis. The sample size will be 60 people with 30 participants per arm, based on recommended sample sizes between 24 – 50 for a feasibility study (37,38). We will invite a purposive subsample of 10-14 participants from the feasibility study to take part in the qualitative study.

Participants – Inclusion and exclusion criteria

Participants will be people with diabetes as defined by the World Health Organisation (39) and a suspected or confirmed diagnosis of CN who are attending NHS multidisciplinary specialist diabetic foot services. They will be identified, recruited and consented by the healthcare professionals working in the foot clinics, these will include podiatrists, nurses and doctors. The full inclusion and exclusion criteria are shown in Table 1. The main exclusion criteria were selected because: 1) they are contra-indications to having an MRI scan, 2) bilateral disease prevents temperature comparison with the contra-lateral limb, and 3) co-morbidities may alter people's inflammatory response. A confirmed diagnosis of CN can take several weeks, so participants will be recruited as early as possible to accurately collect length of time in below knee non-removable cast or boot. If the clinical team decides on an alternative diagnosis during the trial, then the participant will exit the study. We anticipate that alternative diagnosis will include infection, gout, arthritis, soft tissue injuries, or deep vein thrombosis. Follow-up care will be provided by the appropriate clinical team.

For the qualitative study we have identified five participant characteristics which will purposively inform the sampling framework and will seek to maximise variation in gender, age, history of previous foot complications, duration of treatment for the current episode of CN, and employment status. In addition to these factors we will also ensure that participants equally represent both study arms.

Outcomes

We will measure a range of feasibility, clinical and patient centred outcomes (Table 2). We will record time to MRI confirmed remission of CN with off-loading treatment (days) and its variance will be used to inform the sample size for a full-scale trial.

For participants in the standard care arm remission is defined as a temperature difference of ≤ 2^0C which is maintained or improves on two separate consecutive occasions for a period of at least four weeks (8) or at the discretion of the clinical team when temperature difference is not valid; for example in the presence of bilateral foot disease. In the standard care plus arm this will then trigger an MRI. In the intervention arm remission is defined as an absence of sub-chondral bone marrow oedema on MRI, as reported by a radiologist and the absence of clinical signs and symptoms of CN. The clinical team will interpret the results of the MRI report to determine remission.

The final visit will be six months after remission. During these six months we will continue to monitor the foot using the standardised assessment of foot temperature for any clinical signs that the CN has relapsed. We have defined relapse as a temperature difference of >2°C compared to the contralateral foot maintained for two or more occasions or further changes on imaging. The final decision as to whether the CN has relapsed will be at the discretion of the clinical team.

We will explore the feasibility of collecting resource use and quality of life data, to inform the design of the health economics component of a future definitive trial. Data on all

primary care and secondary care visits and admissions to hospital will be collected. Time off work and levels of informal care will also be assessed. We will use the qualitative interviews to gain a deeper, more detailed and rounded contextualised understanding of participants' lived experience of CN and of taking part in this study.

Planned interventions

Standard care plus participants will receive standard care for the assessment and management of CN and any other foot problems; alongside this we will collect study measures (Figure 2). If participants have not had a recent diagnostic X-ray or MRI (within the last three weeks) this will be requested. In this study we have standardised the assessment of foot temperature to monitor CN by using the same device, the Thermofocus 01500A3®. Every 14 days the temperature of both feet will be recorded at intervals of 5 minutes, starting at the removal of the off-loading device and up to 15 minutes. The sites where the temperature will be measured are based on the classification tool developed by Sanders and Frykberg (40). We will classify the stage using the modified (41) Eichenholtz classification tool (42) and location of the CN (40) at baseline using anterior/posterior, oblique and lateral weight bearing X-rays.

Intervention: In addition to standard care plus, participants in the intervention arm will receive serial MRIs at 3, 6, 9 and 12 months. Intervention participants will not undergo further MRIs once remission has been diagnosed, i.e., if remission is diagnosed at 6 months the MRIs at 9 and 12 months will not occur.

317

318 ***Study Procedures (Figure 2)***

319 The schedule of enrolment, interventions and assessments is shown in Figure 2. After giving
320 written informed consent (see Appendix 1) participants will attend for visits every 14 days
321 until remission. All visits will take place in multidisciplinary foot clinics. Wherever possible
322 study measurements and trial interventions will coincide with the participant's existing clinic
323 appointments. This will reduce study burden which is likely to help increase recruitment and
324 retention rates. The study protocol (v1.3, dated 22nd July 2019) is based on the Standard
325 Protocol Items: Recommendations for Interventional Trials (SPIRIT) 2013 Statement for
326 protocols of clinical trials (see Additional file 1).

327

328 Prior to participating in the interviews about the lived experience of CN, participants will
329 receive a further patient information sheet explaining the purpose of the interview and will
330 be asked to complete another consent form (see Appendix 2). All the qualitative interviews
331 will be carried out by the first author (CG), using a semi-structured approach. The topic
332 guide will include a number of probes designed to prompt the participant to increase the
333 level of detail and depth of the information provided from the participants' own viewpoint.
334 Interviews will last approximately 30-40 minutes in a place of the participant's choosing. The
335 interviews will be audiotaped (with the participant's permission) and transcribed in full to
336 capture language and their own expressions.

337

338 ***Analyses***

Quantitative analysis

The feasibility measures including eligibility, recruitment, retention, and withdrawals will be reported as point estimates with 95% confidence intervals. There is no intention to conduct any formal comparative analyses for these measures, though levels of missing data will be explored with respect to certain baseline characteristics, e.g., age and measures of disease severity. Variability in outcomes (e.g. standard deviation) will be estimated with 95% confidence intervals to inform the sample size calculations for a full-scale trial. Any between-group efficacy analyses will only be exploratory. There are no plans for any interim analyses.

We will assess progression of foot deformity by comparing X-rays at baseline, remission and six months post remission. We will measure the change in the Calcaneal Inclination, Talar Declination and Talo-first metatarsal angle between the X-rays. People who have undergone previous minor amputation and/or previous orthopaedic surgical fixation of the foot which alters or removes the anatomical landmarks of the foot will be excluded from this analysis due to the absence of bony landmarks.

The main purpose of the economic analysis is to inform how the data on costs and effects would be collected within a definitive study. Thus, we will estimate completion rates and seek to identify big cost drivers, in order to inform this decision. A preliminary cost-effectiveness analysis will also be performed, although the findings will be treated with

caution. As such, we will estimate the mean incremental cost and mean QALY gain associated with the intervention compared to standard care plus.

Qualitative analysis

The qualitative interviews will be analysed using Inductive Thematic Analysis using the six-step model (43). The first author (CG) will read all the transcribed interviews to record emerging ideas. The interviews will then be subjected to line by line coding using the NVivo data management package. The coding framework will be refined by a second researcher, who will cross-check it against a small sample of transcripts. A modified framework approach will be used to organise the analysis. The coded data will be subjected to a thematic analysis, identifying key categories and themes from the data, ensuring that all participants' responses are adequately captured, and their meaning authentically interpreted. This approach will provide rich descriptions of the data representing accounts of the diverse and personal experiences of people who have taken part in the study and been treated for acute Charcot neuroarthropathy.

Data management and quality assurance

We will set up a Trial Management Group to assist with co-ordination and strategic management of the feasibility study. An initial on-site initiation visit will be completed by CG prior to the sites opening. The primary method of data collection by the research teams will be direct online entry of data onto a purpose-designed secure password-protected electronic case record form. The database complies with data protection requirements (44)

on confidentiality and anonymity. Quality management and monitoring procedures have been discussed and agreed with the sponsor. Central monitoring has been considered appropriate for this study with the option to escalate findings and conduct 'for-cause" on-site triggered monitoring visit if indicated. We will review completed consent forms and selected data points for quality assurance at each site within a week after randomisation of the first participant. Subsequent monitoring will be completed at six monthly intervals to coincide with the Trial Management Group meetings and at the end of data collection.

Safety reporting

Safety monitoring and reporting of adverse events has been discussed and agreed with the sponsor. The study has been assessed as low risk, therefore there will not be a Data Monitoring Committee. The intervention consists of increased frequency of MRI scans without contrast, so a pragmatic approach to safety reporting will be used. MRI scans will be performed in NHS hospitals under routine clinical protocols. Adverse events resulting from MRI scans will be reported by the research teams in line with the Hospital Trust's clinical incident reporting policy. A copy of the anonymised incident form will be forwarded to the Chief Investigator (CG) and reviewed by the Trial Management Group. All other anticipated events, e.g., ulceration, infection, amputation, pain, falls and death will be recorded as secondary outcomes.

Discussion

CN is a poorly understood and under researched complication of diabetes, associated with increased morbidity and mortality compared to people with diabetes without peripheral neuropathy. Evidence is lacking about factors that influence the unexplained variation in treatment times, relapse rates and complications such as ulceration and amputation. We have also identified a lack of evidence to support the efficacy of current monitoring techniques in CN. There is evidence from small studies that MRI may be superior to current methods of monitoring for remission in CN, but this has not been formally evaluated using robust designs. The results of this feasibility study will inform the decision about progressing to a full-sized pragmatic randomised controlled trial: the number of sites required, trial design, the frequency of MRI monitoring, and the choice of process and outcome measures. The embedded qualitative study will provide contextual and meaningful insight into people's experiences of living with CN and what factors they see as contributing to their engagement with the prescribed treatment. Secondly, the qualitative study will advance our understanding of how the condition impacts on participants' quality of life and may contribute to future work on Patient Reported Outcomes Measures in this area (45). Finally, the findings from the qualitative study will provide additional insights into aspects of the trial design and processes that could be improved, in terms of engagement of, and acceptability to participants, based on the participants' experience of involvement in the feasibility study. These aspects could include feedback on the frequency of trial visits, the length of the active and follow-up phases of the trial. and the choice and frequency of completing validated questionnaires. The results of this study will be disseminated to researchers, clinicians, people with diabetes and relevant stakeholders through presentations, publications, and social media press releases.

Trial Status

The CADOM trial originally opened for recruitment in December 2017 and is currently recruiting participants. Recruitment will continue until the end of November 2019.

Abbreviations

ABPI	Ankle brachial pressure index
BMI	Body mass index
CN	Charcot neuroarthropathy
eGFR	Estimated Glomerular Filtration rate, ml/min
EQ-5D-5L	Euroqol 5D
F	Follow up visit
HADS	Hospital Anxiety and Depression Scale
HbA1c	Glycated haemoglobin (A1c), mmol/mol
MRI	Magnetic Resonance Imaging
NHS	National Health Service
R	Remission
SF-12	Medical Outcomes Short-Form Health Questionnaire
VAS	Visual analogue scale

Ethical approval and consent to participate

The trial has been reviewed by East Midlands - Derby Research Ethics Committee, 04/10/2017, ref: 17/EM/0288. The trial is registered on the ISRCTN registry: reference number ISRCTN74101606. All participants will provide written consent to take part in the feasibility trial and will be re-consented by a member of the research team prior to participating in the qualitative interviews. In the future if amendments to the protocol are required the Chief Investigator (CG) will work with the sponsor to apply for approval from Research Ethics Committee and the Health Research Association. Following approval of the amendments this will be cascaded to the research sites. The NHS indemnity scheme will apply to the potential liability of the sponsor for harm to participants arising from the management and conduct of the research.

Consent for Publication

Not applicable

Availability of data and materials

The datasets generated and/or analysed during the current trial will be available from the corresponding author on reasonable request, provided appropriate credit is attributed to the original authors and the data source.

Competing interests

The investigators named on the protocol have no financial or other competing interests that impact on their responsibilities towards the scientific value or potential publishing activities associated with the trial.

Funding

Catherine Gooday is funded by a National Institute for Health Research (NIHR), Clinical Doctoral Fellowship for this research project. This paper presents independent research funded by the National Institute for Health Research (NIHR). The views expressed are those of the authors and not necessarily those of the NHS, the NIHR or the Department of Health and Social Care.

Authors' contributions

CG is the NIHR Clinical Doctoral Fellow and Chief Investigator. CG and FG developed the initial idea for the research. WH, FP, FG, JW, ES and KD all made substantial contributions to the conception and design of the trial. CG drafted the manuscript. All authors critically reviewed and revised the manuscript for important intellectual content. LS provides statistical support. All authors read, amended and approved the final manuscript.

Acknowledgements

This trial is sponsored by the Norfolk & Norwich University Hospitals NHS Trust rdoffice@nnuh.nhs.uk. We are very grateful to all the staff at the Norwich Clinical Trials Unit, all the staff and participants at the trial centres, and Ms Sarah Doyle, Research Administrator.

Inclusion Criteria	Exclusion Criteria
Participants who are willing and have capacity to give informed consent.	People who have received a transplant and others receiving immunosuppressant therapy or using long-term oral glucocorticoids other than in the routine management of glucocorticoid deficiency. Participants on a low dose of oral glucocorticoids (<10mgs for ≤7 days) are eligible to participate in the study.
People with diabetes as diagnosed by the WHO criteria http://www.who.int/diabetes/publications/diagnosis_diabetes2011/en/	Participation in another intervention study on active CN.
Age 18 years or over.	Contra-indication for MRI.
New or suspected diagnosis of acute CN (no previous incidence of acute CN within the last 6 months on the same foot) treated with off-loading.	Treatment for previous suspected CN on the same foot in the last 6 months.
Understand written and verbal instructions in English.	Suspected or confirmed bilateral active CN at presentation.
	Active osteomyelitis at randomisation
	Previous contralateral major amputation.
	Inability to have an MRI scan.
	People receiving palliative care.

Table 1 – Inclusion and exclusion criteria

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497 ***Table 2 – Feasibility, clinical efficacy and patient reported outcomes***

Feasibility outcomes	Clinical efficacy outcomes *Collected – all study visits*	Patient reported outcomes *Collected – baseline, 3 monthly until remission, then at 1 and 6-months post remission*
The proportion of patients who meet the eligibility criteria	Number of new ulcerations on the index foot	Health related quality of life measured: Short Form 12 questionnaire (SF-12) (46) EuroQol-5D-5L questionnaire (EQ-5D-5L) (47)
The number of eligible patients recruited	Number of new ulcerations on the index or contralateral foot	Hospital Anxiety and Depression Scale (HADS) (48)
The number of participants in which an alternative diagnosis is made during the active phase of the trial	Number of new infections on the index or contralateral foot	Pain as assessed by Visual Analogue Scale (VAS)
The proportion of patients that withdraw or are lost to follow up. The term 'withdrawal' encompasses two potential scenarios; withdrawal due to loss of consent or withdrawal due to death	Number of minor and major amputations on the index foot or contralateral at the end of the follow up phase of the study	
Statistical parameters of the key outcome measures to inform a sample size calculation for a definitive trial	Number and severity of falls (Hopkins Fall Grading System)(49)	
Ability to collect quality of life and resource use data	The number of participants in each arm requiring further intervention for CN (e.g. further immobilisation) within 6 months of remission	

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Figure 1 – Patient flow diagram

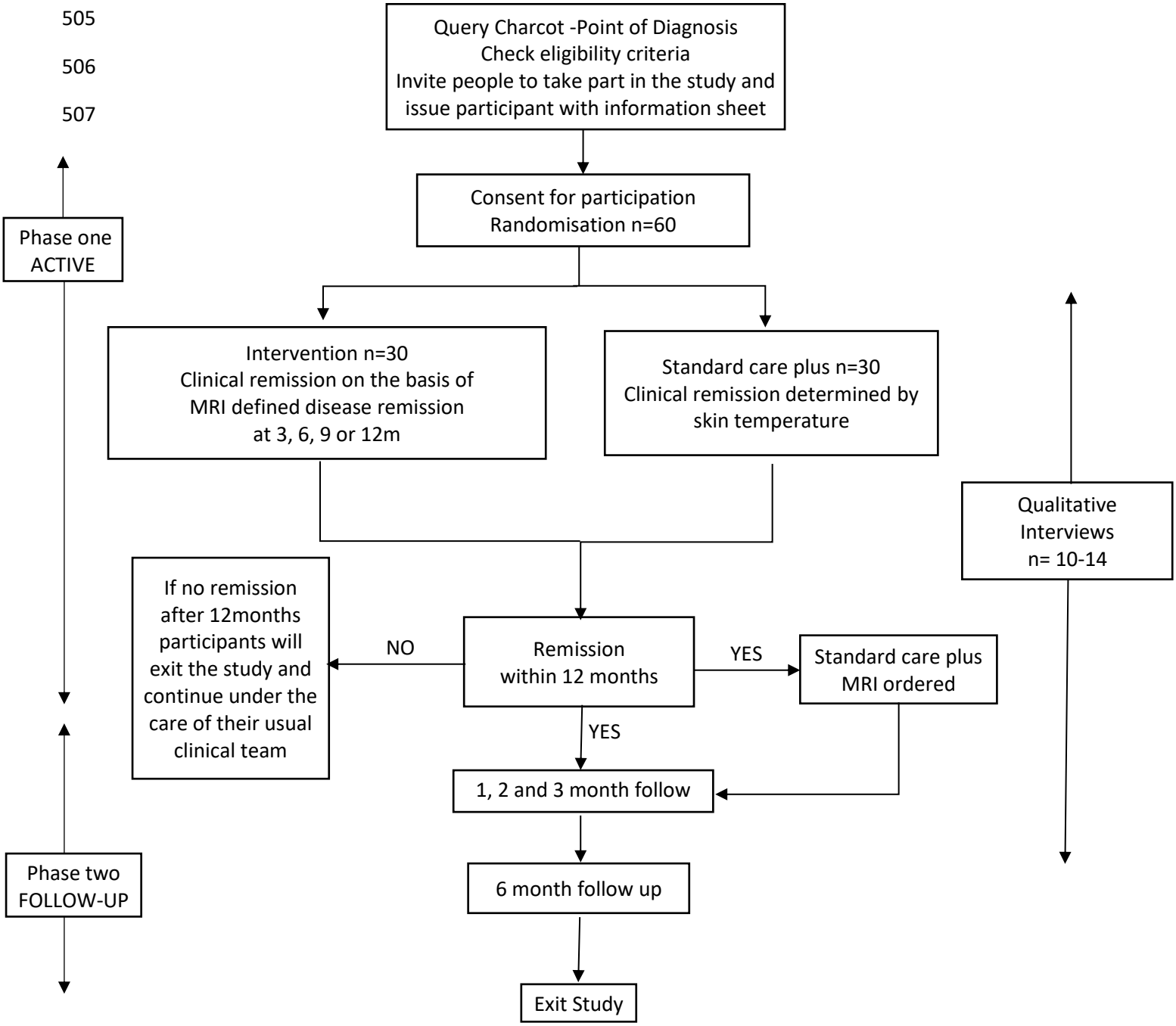

Figure 2 - Schedule of enrolment, interventions and assessments

		Active phase (maximum 12 months)									*R*	*Follow up phase*			
Visit Number		***1***		***6***		***11***		***18***		***26***		***F1***	***F2***	***F3***	***F4***
Month		***0***		***3***		***6***		***9***		***12***		***1***	***2***	***3***	***6***
Enrolment															
Information sheet	*														
Consent		*													
Randomisation		*													
Participant characteristics															
Medical history		*													
HbA1c & eGFR		*													
Foot surgical history		*													
Medications		*													
Classification of CN		*													
Foot assessment															
Foot pulses		*													
ABPI		*													
10g monofilament		*													
Neurotheisometer		*													
Foot temperatures		*	*	*	*	*	*	*	*	*	*	*	*	*	*
Treatment															
Off-loading/footwear		*	*	*	*	*	*	*	*	*	*	*	*	*	*
Interventions															
MRI (standard care plus)											*				
Serial MRI (intervention)				*		*		*		*					
Clinical outcomes															
Ulceration		*	*	*	*	*	*	*	*	*	*	*	*	*	*
Infection		*	*	*	*	*	*	*	*	*	*	*	*	*	*
Amputation		*	*	*	*	*	*	*	*	*	*	*	*	*	*
Falls		*	*	*	*	*	*	*	*	*	*	*	*	*	*
BMI		*		*		*		*		*	*				*
X-ray															*
Patient centred outcomes															
VAS - pain		*		*		*		*		*		*			*
HADS		*		*		*		*		*		*			*
EQ-5D-5L		*		*		*		*		*		*			*
SF-12		*		*		*		*		*		*			*
Health economic outcomes															
Issue patient diary		*	*	*	*	*	*	*	*	*					
Collect patient diary			*	*	*	*	*	*	*	*	*				
Qualitative Study															
Interview			◄	—	—	—	—	—	—	—	—	—	—	—	►

Active phase - while the CN is active participants will attend every 14 days, up to a maximum of 26 visits.
Follow up phase – once CN is in remission participants will transfer into the follow-up phase of the study for six months.
Classification of CN – accordingly to the Sanders and Frykberg and the modified Eichenholtz classification tools

Abbreviations

ABPI – Ankle brachial pressure index
BMI – Body mass index
CN – Charcot neuroarthropathy
eGFR – Estimated Glomerular Filtration rate, ml/min
EQ-5D-5L - Euroqol 5
F – Follow up visit
HADS - Hospital Anxiety and Depression Scale
HbA1c – glycated haemoglobin (A1c), mmol/mol
MRI – Magnetic Resonance Imaging
R – Remission
SF-12 - Medical Outcomes Short-Form Health Questionnaire
VAS – Visual analogue scal

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Appendix 1 – Informed consent form - feasibility trial

University of East Anglia

Insert local header

A study to assess the use of serial MRI to reduce treatment times in Charcot in people with diabetes.

(Short title: CADOM)

Charcot neuroArthropathy Diagnostic Outcome Measures

Patient Consent Form

Principal Investigator:……..

Patient Study ID: ……. Initials: ……

Please initial each box

1. I confirm that I have read and understand the information sheet Version 1.2 10th January 2019 for the above study. I have had the opportunity to ask questions and been given satisfactory answers. ☐

2. I have been given a full explanation of the purpose of the study and what I will be expected to do. ☐

3. I understand that my medical notes and data collected during the Study may be looked at by individuals from the Clinical Trials Unit at the University of East Anglia, from regulatory authorities or from the

YES	NO

NHS Trust, where it is relevant to my taking part in this research, I give permission for these individuals to have access to my records.

4. I understand that my participation is voluntary and that I am free to withdraw at any time without my medical care or legal rights being affected ☐

5. I consent to the storage including electronic, of personal information for the purposes of this study. I understand that any information that could identify me will be kept strictly confidential and that no personal information will be included in the study report or other publication. ☐

6. I understand that even if I withdraw from the above study, the data collected from me up to that point will be used in analysing the results of the study. ☐

7. In the event that the MRI or X-ray shows a previous unknown condition that might need further medical or surgical intervention I agree to the research team referring me on as necessary and informing my GP. ☐

8. I understand that information held by the NHS and records maintained by the NHS Information Centre may be used to keep in touch with me and my health status. I give my permission to register my identifiable details with the NHS Information Centre. ☐

9. I agree to being contacted by the research team when the Charcot has settled, to ask if I would consider taking part in an interview. The interview would involve discussing the experience of being diagnosed and treated for Charcot, and being involved in this study

☐	☐
YES	NO

10. I give permission for a copy of this consent form to be kept confidentially and securely by the Norwich Clinical Trials Unit. ☐

11. I am happy to be contacted to receive updates on how the study is progressing and to be informed about the results of the study at the end ☐

12. I agree to take part in the study. ☐

…….	……..	…….
Name of the patient (Print)	Date	Patient's signature
…….	……..	…….
Name of person taking consent (Print)	Date	Signature

Original to be retained and filed in the site file. 1 copy to patient, 1 copy to be filed in patient's notes

Appendix 2- Informed consent form – qualitative interviews

Insert local header

Interviews

Experiences of being treated for Charcot neuroarthropathy and views about taking part in the clinical trial.

(Short title: CADOM)

Charcot neuroArthropathy Diagnostic Outcome Measures

Patient Consent Form

Principal Investigator: ……..

Patient Study ID: …….. Initials: ……

Please initial each box

1. I confirm that I have read and understand the information sheet Version 1.1 dated 25th August 2017 for the above study. I have had the opportunity to ask questions and been given satisfactory answers. ☐

2. I have been given a full explanation of the purpose of the study and what I will be expected to do. ☐

3. I understand that my medical notes and data collected during the study may be looked at by individuals from the Clinical Trials Unit at the University of East Anglia, from regulatory authorities or from the NHS Trust, where it is relevant to my taking part in this research, I give permission for these individuals to have access to my records. ☐ YES ☐ NO

4. I understand that my participation in the interview is voluntary and that I am free to withdraw from the study at any time, without having to give a reason. ☐

5. I understand that the interview will be recorded on a digital recorder. I give permission for doing this. ☐

6. I understand that the recordings will be saved on a secure computer at the University of East Anglia. The recordings will be destroyed at the end of the study. The transcripts will be kept for 15 years. ☐

7. I consent to the storage including electronic, of personal information for the purposes of this study. I understand that any information that could identify me will be kept strictly confidential and that no personal information will be included in the study report, my thesis, or other publication. ☐

8. I understand that what I say during the interview is confidential, in accordance with the Data Protection Act. However, you must be aware that if you tell the interviewer something which shows that there is a significant risk to you or someone else, they may need to pass this information on.
   If this happens, they will discuss it with you first before anyone else is told ☐

9. I am happy to be contacted to receive updates on how the study is progressing and to be informed about the results of the study at the end.

YES	NO
☐	☐

10. I give permission for a copy of this consent form to be kept confidentially and securely by the Norwich Clinical Trials Unit. ☐

11. I agree to take part in an interview for the above study. ☐

……	……..	……
Name of the patient (Print)	Date	Patient's signature
……	……..	……
Name of person taking consent (Print)	Date	Signature

Original to be retained and filed in the site file. 1 copy to patient, 1 copy to be filed in patient's notes

Supplementary File 1 – SPIRIT Checklist

SPIRIT 2013 Checklist: Recommended items to address in a clinical trial protocol and related documents*

Section/item	Item No	Description	Addressed on page number
Administrative information			
Title	1	Descriptive title identifying the study design, population, interventions, and, if applicable, trial acronym	1
Trial registration	2a	Trial identifier and registry name. If not yet registered, name of intended registry	5
	2b	All items from the World Health Organization Trial Registration Data Set	yes
Protocol version	3	Date and version identifier	16
Funding	4	Sources and types of financial, material, and other support	23
Roles and responsibilities	5a	Names, affiliations, and roles of protocol contributors	1-3
	5b	Name and contact information for the trial sponsor	23
	5c	Role of study sponsor and funders, if any, in study design; collection, management, analysis, and interpretation of data; writing of the report; and the decision to submit the report for publication, including whether they will have ultimate authority over any of these activities	23
	5d	Composition, roles, and responsibilities of the coordinating centre, steering committee, endpoint adjudication committee, data management team, and other individuals or groups overseeing the trial, if applicable (see Item 21a for data monitoring committee)	N/A
Introduction			

Background and rationale	6a	Description of research question and justification for undertaking the trial, including summary of relevant studies (published and unpublished) examining benefits and harms for each intervention	6-10
	6b	Explanation for choice of comparators	8-10
Objectives	7	Specific objectives or hypotheses	10-11
Trial design	8	Description of trial design including type of trial (eg, parallel group, crossover, factorial, single group), allocation ratio, and framework (eg, superiority, equivalence, noninferiority, exploratory)	11
Methods: Participants, interventions, and outcomes			
Study setting	9	Description of study settings (eg, community clinic, academic hospital) and list of countries where data will be collected. Reference to where list of study sites can be obtained	12
Eligibility criteria	10	Inclusion and exclusion criteria for participants. If applicable, eligibility criteria for study centres and individuals who will perform the interventions (eg, surgeons, psychotherapists)	12-13,24
Interventions	11a	Interventions for each group with sufficient detail to allow replication, including how and when they will be administered	15
	11b	Criteria for discontinuing or modifying allocated interventions for a given trial participant (eg, drug dose change in response to harms, participant request, or improving/worsening disease)	N/A
	11c	Strategies to improve adherence to intervention protocols, and any procedures for monitoring adherence (eg, drug tablet return, laboratory tests)	16
	11d	Relevant concomitant care and interventions that are permitted or prohibited during the trial	16,27

Outcomes	12	Primary, secondary, and other outcomes, including the specific measurement variable (eg, systolic blood pressure), analysis metric (eg, change from baseline, final value, time to event), method of aggregation (eg, median, proportion), and time point for each outcome. Explanation of the clinical relevance of chosen efficacy and harm outcomes is strongly recommended	25
Participant timeline	13	Time schedule of enrolment, interventions (including any run-ins and washouts), assessments, and visits for participants. A schematic diagram is highly recommended (see Figure)	27
Sample size	14	Estimated number of participants needed to achieve study objectives and how it was determined, including clinical and statistical assumptions supporting any sample size calculations	12
Recruitment	15	Strategies for achieving adequate participant enrolment to reach target sample size	12
Methods: Assignment of interventions (for controlled trials)			**12**
Allocation:			
Sequence generation	16a	Method of generating the allocation sequence (eg, computer-generated random numbers), and list of any factors for stratification. To reduce predictability of a random sequence, details of any planned restriction (eg, blocking) should be provided in a separate document that is unavailable to those who enrol participants or assign interventions	12
Allocation concealment mechanism	16b	Mechanism of implementing the allocation sequence (eg, central telephone; sequentially numbered, opaque, sealed envelopes), describing any steps to conceal the sequence until interventions are assigned	12
Implementation	16c	Who will generate the allocation sequence, who will enrol participants, and who will assign participants to interventions	12
Blinding (masking)	17a	Who will be blinded after assignment to interventions (eg, trial participants, care providers, outcome assessors, data analysts), and how	11

	17b	If blinded, circumstances under which unblinding is permissible, and procedure for revealing a participant's allocated intervention during the trial	N/A
Methods: Data collection, management, and analysis			
Data collection methods	18a	Plans for assessment and collection of outcome, baseline, and other trial data, including any related processes to promote data quality (eg, duplicate measurements, training of assessors) and a description of study instruments (eg, questionnaires, laboratory tests) along with their reliability and validity, if known. Reference to where data collection forms can be found, if not in the protocol	25
	18b	Plans to promote participant retention and complete follow-up, including list of any outcome data to be collected for participants who discontinue or deviate from intervention protocols	16
Data management	19	Plans for data entry, coding, security, and storage, including any related processes to promote data quality (eg, double data entry; range checks for data values). Reference to where details of data management procedures can be found, if not in the protocol	18
Statistical methods	20a	Statistical methods for analysing primary and secondary outcomes. Reference to where other details of the statistical analysis plan can be found, if not in the protocol	16-18
	20b	Methods for any additional analyses (eg, subgroup and adjusted analyses)	N/A
	20c	Definition of analysis population relating to protocol non-adherence (eg, as randomised analysis), and any statistical methods to handle missing data (eg, multiple imputation)	N/A
Methods: Monitoring			
Data monitoring	21a	Composition of data monitoring committee (DMC); summary of its role and reporting structure; statement of whether it is independent from the sponsor and competing interests; and reference to where further details about its charter can be found, if not in the protocol. Alternatively, an explanation of why a DMC is not needed	19

	21b	Description of any interim analyses and stopping guidelines, including who will have access to these interim results and make the final decision to terminate the trial	N/A
Harms	22	Plans for collecting, assessing, reporting, and managing solicited and spontaneously reported adverse events and other unintended effects of trial interventions or trial conduct	19
Auditing	23	Frequency and procedures for auditing trial conduct, if any, and whether the process will be independent from investigators and the sponsor	18-19
Ethics and dissemination			
Research ethics approval	24	Plans for seeking research ethics committee/institutional review board (REC/IRB) approval	22
Protocol amendments	25	Plans for communicating important protocol modifications (eg, changes to eligibility criteria, outcomes, analyses) to relevant parties (eg, investigators, REC/IRBs, trial participants, trial registries, journals, regulators)	22
Consent or assent	26a	Who will obtain informed consent or assent from potential trial participants or authorised surrogates, and how (see Item 32)	13
	26b	Additional consent provisions for collection and use of participant data and biological specimens in ancillary studies, if applicable	22
Confidentiality	27	How personal information about potential and enrolled participants will be collected, shared, and maintained in order to protect confidentiality before, during, and after the trial	18
Declaration of interests	28	Financial and other competing interests for principal investigators for the overall trial and each study site	23
Access to data	29	Statement of who will have access to the final trial dataset, and disclosure of contractual agreements that limit such access for investigators	22
Ancillary and post-trial care	30	Provisions, if any, for ancillary and post-trial care, and for compensation to those who suffer harm from trial participation	22

Disseminatio n policy	31a	Plans for investigators and sponsor to communicate trial results to participants, healthcare professionals, the public, and other relevant groups (eg, via publication, reporting in results databases, or other data sharing arrangements), including any publication restrictions	20
	31b	Authorship eligibility guidelines and any intended use of professional writers	23
	31c	Plans, if any, for granting public access to the full protocol, participant-level dataset, and statistical code	N/A
Appendices			
Informed consent materials	32	Model consent form and other related documentation given to participants and authorised surrogates	32-37
Biological specimens	33	Plans for collection, laboratory evaluation, and storage of biological specimens for genetic or molecular analysis in the current trial and for future use in ancillary studies, if applicable	N/A

1