

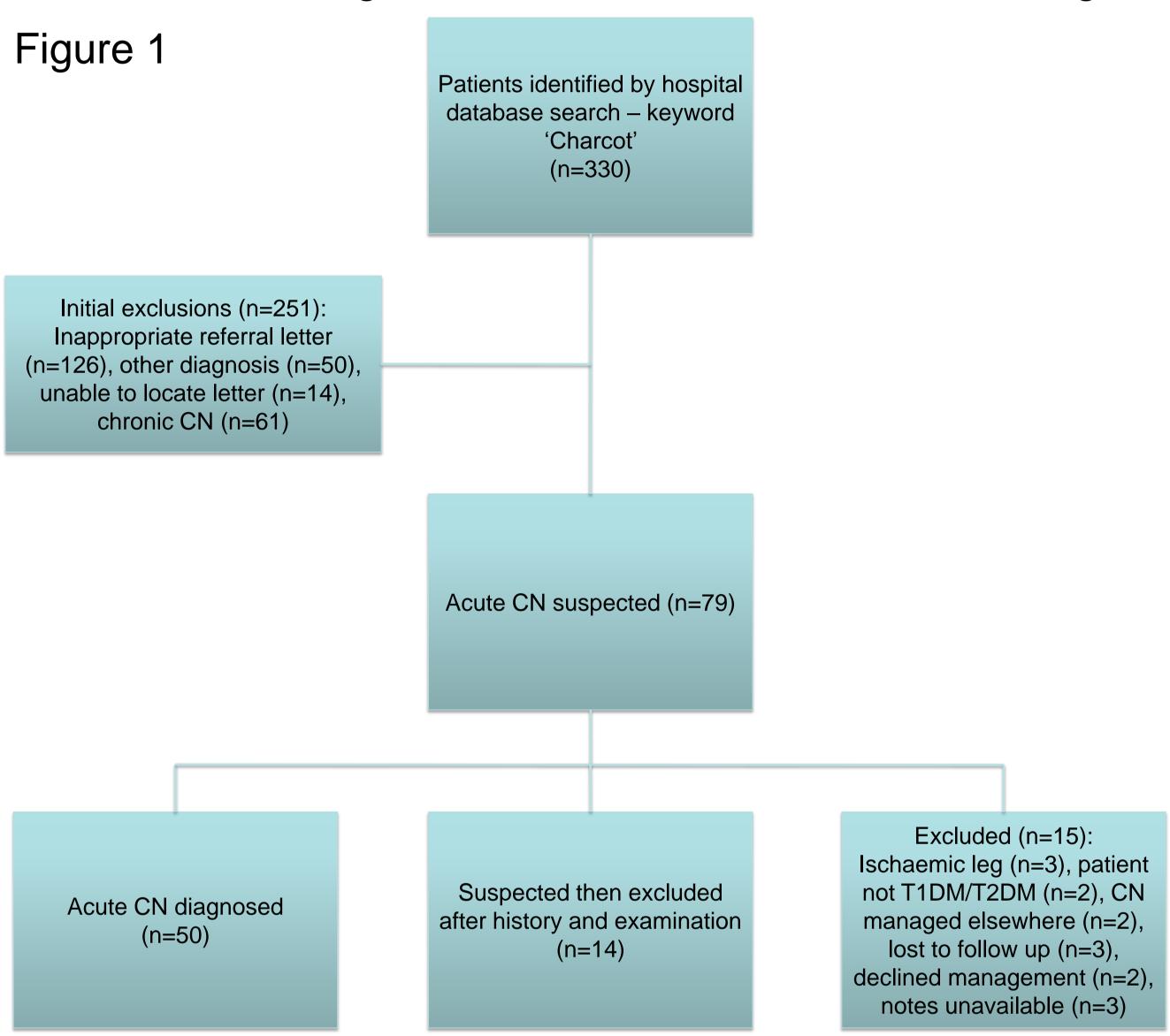
## Outcomes in acute Charcot neuroarthropathy – a single centre experience over 5 years

## \*Coral Stark<sup>1</sup>, \*Tom Murray<sup>1</sup>, Catherine Gooday<sup>2</sup>, Ketan Dhatariya<sup>1,2</sup>

<sup>1</sup>Norwich Medical School, University of East Anglia, Norwich, UK <sup>2</sup>Diabetic Foot Clinic, Elsie Bertram Diabetes Centre, Norfolk and Norwich University Hospital NHS Foundation Trust, Norwich, UK

Background: Charcot neuroarthropathy (CN) is a rare complication of diabetes. A recent, large systematic review suggested that the current gold standard management of acute CN consists of immediate referral to a multidisciplinary foot-care team followed by immobilisation of the foot in a total contact plaster cast (TCC)<sup>1.</sup> However, there are few data describing the natural history and long term results of CN treated with a TCC. Our aim was to look at time taken to achieve clinical resolution, and to see if there was a correlation with location within the foot.

Methods: We performed a retrospective analysis of patients presenting with a new diagnosis of acute CN between October 2007 and October 2012. Figure 1 shows the numbers at each stage of the patient inclusion / exclusion criteria.



**Discussion:** There is general consensus that immobilisation of the foot is necessary to prevent progression in the acute Charcot foot. However there is generally poor quality evidence to differentiate between a TCC and a removable below knee walking boot<sup>1</sup>. Our results are in contrast to those reported by the CDUK group who found that median time to resolution varies greatly between those initially treated in a TCC compared to removable offloading device (12 months and 9 months respectively)<sup>2</sup>. Our study also showed a 38.9% deterioration rate after coming out of TCC – although there were no clinical findings that would help to determine who was at risk of deterioration

**Conclusion:** This study has shown that the mean time to clinical resolution for a newly presenting CN was just over 1 year (53.9±28.0 weeks). Our data shows that the median time to resolution was similar for those initially treated in a non-removable TCC (50 weeks (95% CI: 40 , 66)) and for those initially treated in removable offloading device (53 weeks (95% CI: 35, 68)). These results imply that there is no evidence that wearing a TCC as the initial device is beneficial in reducing time to treatment.

**Results:** In total, 50 patients were included, 34 (68%) were male, whilst 16 (32%) were female. 11 (22%) had T1DM, with the majority having T2DM (78%).

The mean (±SD) duration of DM was 17.9±12.9 years (T1DM) 29.7±12.9; T2DM 14.4±10.7), with the mean age of diagnosis of CN being 62.5±11.7 years (T1DM 55.8; T2DM 64.4). The mean HbA1c was 65.4±19.8 mmol/mol (T1DM 70.0±19.2; T2DM 64.1±20.0).

41 patients (82%) had documented diabetic retinopathy present at diagnosis of CN and 38 (76%) had CKD stages 2-4.

42 of the 50 patients went into remission during the study period, with foot temperatures <2°C for greater than 6 weeks (3 consecutive visits at least 2 weeks apart to the foot clinic). Of these 42 patients, the proportion of CN by location were; forefoot 11.9%, mid-foot 64.3%, hind-foot or ankle 19.1%, and 4.8% were of multiple sites.

36 patients (85.7%) were treated with both TCC and removable offloading device. The remaining 6 patients were treated with one modality only – 5 with a removable offloading device only and 1 with a TCC only. 25 of the 42 patients (59.5%) were initially treated with a TCC, whilst the remaining 17 (40.5%) started in a removable offloading device.

Median time to resolution for patients initially treated with a TCC was 50 weeks (95% CI:40, 66) compared to the median time of 53 weeks (95% CI:35, 68) for those initially treated with removable offloading device.

14 of the 37 patients ever treated with TCC, required re-casting due to clinical deterioration of the acute CN in the removable device. The median time to resolution for these patients was 68 weeks (95% CI: 53, 89) compared to the 33 who had no re-casting, median time of 42.5 weeks (95% CI: 35, 48), (P<0.0001).

For the 42 patients who completed treatment, mean time (±SD) spent in treatment either was 53.9±28.0 weeks. Of this, mean of 30.2±25.0 weeks was spent in TCC, with 23.7±16.2 weeks being spent in a removable offloading device. For those treated solely in a removable offloading device, the mean duration of treatment was 39.0±16.3 weeks. The one patient treated with only a TCC spent 12 weeks in treatment.

8 patients did not complete treatment during the study period. 4 of these had undergone a major amputation. 2 patients died whilst undergoing treatment, and 2 patients had not completed treatment at the end of the study period.

1. Milne TE, Rogers JR, Kinnear EM et al. Developing an evidence-based clinical pathway for the assessment, diagnosis and management of acute Charcot Neuro-Arthropathy: a systematic review. J Foot Ankle Res 2013; 6:30 2. Game FL, Catlow R, Jones GR et al. Audit of acute Charcot's disease in the UK: the CDUK study. Diabetologia 2012; 55(1):32-35