

# The effect of pregabalin or duloxetine on arthritis pain: a clinical and mechanistic study in people with hand osteoarthritis

Nidhi Sofat<sup>1</sup>  
Abiola Harrison<sup>1</sup>  
Mark D Russell<sup>1</sup>  
Salma Ayis<sup>2</sup>  
Patrick D Kiely<sup>3</sup>  
Emma H Baker<sup>1</sup>  
Thomas Richard Barrick<sup>4</sup>  
Franklyn A Howe<sup>4</sup>

<sup>1</sup>Institute for Infection and Immunity, St George's University of London,

<sup>2</sup>Division of Health & Social Care Research, King's Clinical Trials Unit, King's College London, <sup>3</sup>Department of Rheumatology, St George's University Hospitals NHS Foundation Trust, <sup>4</sup>Molecular and Clinical Sciences Research Institute, St George's University of London, London, UK

**Abstract:** Osteoarthritis (OA) is the most prevalent arthritis worldwide and is characterized by chronic pain and impaired physical function. We hypothesized that heightened pain in hand OA could be reduced with duloxetine or pregabalin. In this prospective, randomized clinical study, we recruited 65 participants, aged 40–75 years, with a Numerical Rating Scale (NRS) for pain of at least 5. Participants were randomized to one of the following three groups: duloxetine, pregabalin, and placebo. The primary endpoint was the NRS pain score, and the secondary endpoints included the Australian and Canadian Hand Osteoarthritis Index (AUSCAN) pain, stiffness, and function scores and quantitative sensory testing by pain pressure algometry. After 13 weeks, compared to placebo, ANOVA found significant differences between the three groups ( $P=0.0078$ ). In the intention-to-treat analysis, the pregabalin group showed improvement for NRS pain ( $P=0.023$ ), AUSCAN pain ( $P=0.008$ ), and AUSCAN function ( $P=0.009$ ), but no difference between duloxetine and placebo ( $P>0.05$ ) was observed. In the per protocol analysis, NRS pain was reduced for pregabalin ( $P<0.0001$ ) and duloxetine ( $P=0.029$ ) compared to placebo. We conclude that centrally acting analgesics improve pain outcomes in people with hand arthritis, offering new treatment paradigms for OA pain.

**Keywords:** pain, hand osteoarthritis, sensitization, duloxetine, pregabalin

## Introduction

Osteoarthritis (OA) is the most common form of arthritis worldwide, with hand pain and reduced function causing significant problems for people with hand OA.<sup>1</sup> Pain is a major symptom for people with OA, with 16.7% of US adults aged 45 years and older reporting pain as a predominant problem.<sup>2</sup> Pain and reduced function due to OA place a huge burden on patients and health care services.<sup>2,3</sup> Although several pharmacological agents are available for OA pain management, including acetaminophen, nonsteroidal anti-inflammatory drugs (NSAIDs), and opioids, a large proportion of patients continue to suffer from chronic pain despite using the agents described.<sup>4,5</sup> Recent trials have raised questions about current treatments, suggesting that acetaminophen has poor efficacy in controlling OA pain.<sup>6,7</sup> Pain management in OA is a huge problem, and novel approaches are urgently needed.

Pain is often characterized as having features of inflammatory nociceptive pain and neuropathic components.<sup>8,9</sup> OA is recognized to have features of inflammatory pain and also pain sensitization. Features of pain sensitization can be evaluated using quantitative sensory testing (QST)<sup>10–13</sup> and brain neuroimaging.<sup>14–16</sup> Large studies have shown pain sensitization using QST in knee OA.<sup>10–13</sup> Brain neuroimaging studies in chronic

Correspondence: Nidhi Sofat  
Institute of Infection and Immunity,  
St George's University of London,  
Mailpoint J1A, Cranmer Terrace  
St George's, London SW17 0RE, UK  
Email [nsofat@sgul.ac.uk](mailto:nsofat@sgul.ac.uk)

OA have also demonstrated increased central pain processing in the cingulate cortex, insula, and thalamus compared with normal controls.<sup>14,15</sup> Many clinical trials testing new agents for OA have focused on large joint hip and knee OA, but relatively few trials have been conducted in hand OA. The ideal analgesic drug(s) in OA would achieve sustained pain relief in a dose-dependent manner with few side effects. Centrally acting analgesic drugs such as pregabalin and duloxetine could fulfill these criteria but have not been investigated in hand OA. Duloxetine is a serotonin and noradrenaline reuptake inhibitor (SNRI) that has shown efficacy for improving pain in knee OA.<sup>17</sup> However, no previous studies have evaluated duloxetine in hand OA. Gabapentinoids are three-substituted derivatives of the neurotransmitter gamma-aminobutyric acid that blocks voltage-dependent calcium channels, used to treat epilepsy and neuropathic pain. Interest has grown in gabapentinoids for arthritis, since gabapentin inhibits pain sensitization.<sup>18,19</sup> Arthritic pain can be improved by NSAIDs and pregabalin in OA.<sup>20,21</sup> Ohtori et al<sup>20</sup> found that pregabalin combined with meloxicam was more effective for knee OA pain compared to either drug alone and Arendt-Nielsen et al<sup>21</sup> showed that pain sensitization is improved by NSAIDs in knee OA.

We hypothesized that centrally acting analgesics may alleviate arthritic pain. We conducted a proof-of-concept, randomized, placebo-controlled study comparing duloxetine and pregabalin to placebo for hand OA pain. We used validated primary endpoints for pain, with secondary endpoints for pain sensitization using QST, depression, and anxiety scores. Our report is the first proof-of-concept clinical trial comparing the effect of centrally acting analgesics duloxetine and pregabalin head-to-head vs placebo in hand OA pain with mechanistic secondary endpoints for pain threshold testing.

## Methods

### Study design and participants

All methods were carried out in accordance with relevant guidelines and regulations. All trial protocols were approved by the sponsors, St George's University of London and the Medicines and Healthcare Products Regulatory Agency, UK. Ethical approval was provided by the London-Surrey Borders Ethics Committee, approval number 12/LO/0047. Written informed consent was obtained from all subjects. The clinical trial registration number is NCT02612233. Participants were eligible if they were aged 40–75 years and had hand OA diagnosed by American College of Rheumatology (ACR) criteria<sup>22,23</sup> confirmed by a rheumatologist and experiencing pain of at least  $\geq 5$  on a Numerical Rating Scale (NRS) of 0–10. The trial protocol was followed as published, according to CONSORT guidelines and inclusion–exclusion criteria.

Twenty age-matched subjects without hand OA were enrolled as controls for comparisons for pain testing and brain MRI. The brain MRI data from this study will be reported in a separate publication.

### Inclusion and exclusion criteria

Inclusion criteria were as follows: participants fulfilling the ACR criteria for the diagnosis of hand osteoarthritis, male or female, right or left handed, aged 40–75 years, and on usual care for hand OA including acetaminophen and/or NSAIDs. Exclusion criteria were another rheumatological diagnosis, eg, rheumatoid arthritis, current or planned pregnancy, contraindications to duloxetine or pregabalin such as concomitant use of monoamine oxidase inhibitors, selective serotonin reuptake inhibitors, antidepressants, oral contraceptives, St. John's wort, history of depression, concomitant use of opioids including tramadol and pethidine, use of benzodiazepines, recent surgery, ie, <6 weeks prior to participation in the study, recent insertion of surgical implants, ie, <6 weeks before participation prior to entry, previous use of duloxetine and/or pregabalin, uncontrolled depression, estimated glomerular filtration rate <60 mL/min, hepatic impairment defined as ALT  $>2.5\times$  upper limit of normal within 6 weeks of last clinical assessment, ischemic heart disease, diabetes mellitus, and regular use of alcohol or alcohol abuse (maximum limits are 28 units/week for men and 21 units/week for women, lactose intolerance). The estimated glomerular filtration rate was checked by screening blood tests, and any participants who were outside the stated range were not enrolled. Uncontrolled hypertension was checked by blood pressure in primary care, and any participants with a blood pressure  $>140/90$  were excluded. Baseline laboratory tests of renal function and hepatic function were performed at baseline to screen for any impairment, and participants with levels outside the normal range were excluded. We checked information from all participants about a new diagnosis of diabetes, and any new cases were excluded. For the Hospital Anxiety and Depression Scale (HADS) scoring, a score of  $\geq 12$  for anxiety and/or depression was considered as too high for enrollment to the study and participants with a score of  $>12$  were excluded.

### Randomization and masking

Study drugs were supplied by Sharp Clinical Services (formerly Bilcare GCS, Powys, UK), which overencapsulated pregabalin 150 mg tablets or duloxetine 30 mg tablets and produced visually identical placebo capsules. A mid-ranging dose was selected for each of the trial medications. The random allocation sequence, with a block size of nine, was generated by the manufacturer and implemented through

sequentially numbered containers. Neither participants nor investigators were aware of treatment assignment until after completion of the trial, which was performed after the last patient and last visit were conducted at the end of the trial. Emergency code breaks were administered independently by the staff from the St George's University Hospitals NHS Foundation Trust's Clinical Trials Pharmacy.

## Clinical outcome measures

The primary endpoints were the NRS and the Australian and Canadian Hand Osteoarthritis Index (AUSCAN) rating scale 3.1 for pain,<sup>24,25</sup> which are validated outcome measures for pain. Both NRS and AUSCAN pain endpoints are well-recognized primary endpoints in hand OA clinical trials and have been recommended in international guidelines.<sup>25,28</sup>

Prespecified secondary endpoints included the AUSCAN stiffness and function scales and HADS<sup>26</sup> at baseline and after 12 weeks treatment. All endpoints were specified prospectively.

## Pain algometry

Pain pressure thresholds (PPTs) were used to obtain objective measures of peripheral pain sensitization as we described previously.<sup>27</sup> Briefly, a calibrated digital hand held algometer (FDX 100; Wagner Instruments, Greenwich, CT, USA) was used for all measurements. A standard operating procedure was used, which consisted of testing pain thresholds in all participants in both hands with  $n=30$  regions for each participant, 780 regions in total. Regions tested included dorsal aspects of all distal interphalangeal, proximal interphalangeal and metacarpophalangeal joints of each digit and thumb and the dorsum of each wrist. The 1 cm<sup>2</sup> flat rubber algometer probe was held perpendicular to the dorsal aspect of the skin, and force was applied to provide a constant increase in pressure at a rate of 1 N/cm<sup>2</sup>/s. Therefore, the algometer scores are stated as Newton per centimeter squared in all reported results. The individual was asked to say "stop" when the sensation of pressure became the first sensation of pain. The algometer was applied to each joint being examined three times in succession with an interval between applications. After all three readings were taken, the average from the last two readings was calculated as the PPT. The intervals between each algometer measurement were long enough to prohibit temporal summation.

## Statistical analysis

Our sample size was based on IMMPACT guidelines<sup>25</sup> and OARSI recommendations for RCTs in hand OA<sup>28</sup> using NRS pain for the sample size calculation. For the NRS pain

outcome, we aimed to detect a mean difference of 2.0 (SD 1.9) points between baseline and treatment after 12 weeks. With 16 participants in each group, 80% power with a 0.05 significance level (two sided) is achieved. Recruitment required up to 22 participants per treatment group, allowing a dropout rate of 25%, giving a total intervention study number of 65 participants to achieve desired statistical power.

Planned analyses included initial comparison to detect any significant differences between baseline and 13-week timepoints using primary endpoint NRS and AUSCAN pain difference between all three groups by ANOVA, with a multiple comparisons test,  $\alpha=0.05$ . Following ANOVA, pairwise comparisons were performed for placebo vs pregabalin and placebo vs duloxetine. The intent-to-treat analysis was performed using the last observation at week 13 and carried forward for all participants. We present the NRS pain and AUSCAN pain, stiffness, and function outcomes as mean and confidence interval for all analyses. These are presented after checking the distribution of the data, which followed a normal distribution and were not skewed for the parameters measured. We also show the per protocol analysis for all completers.

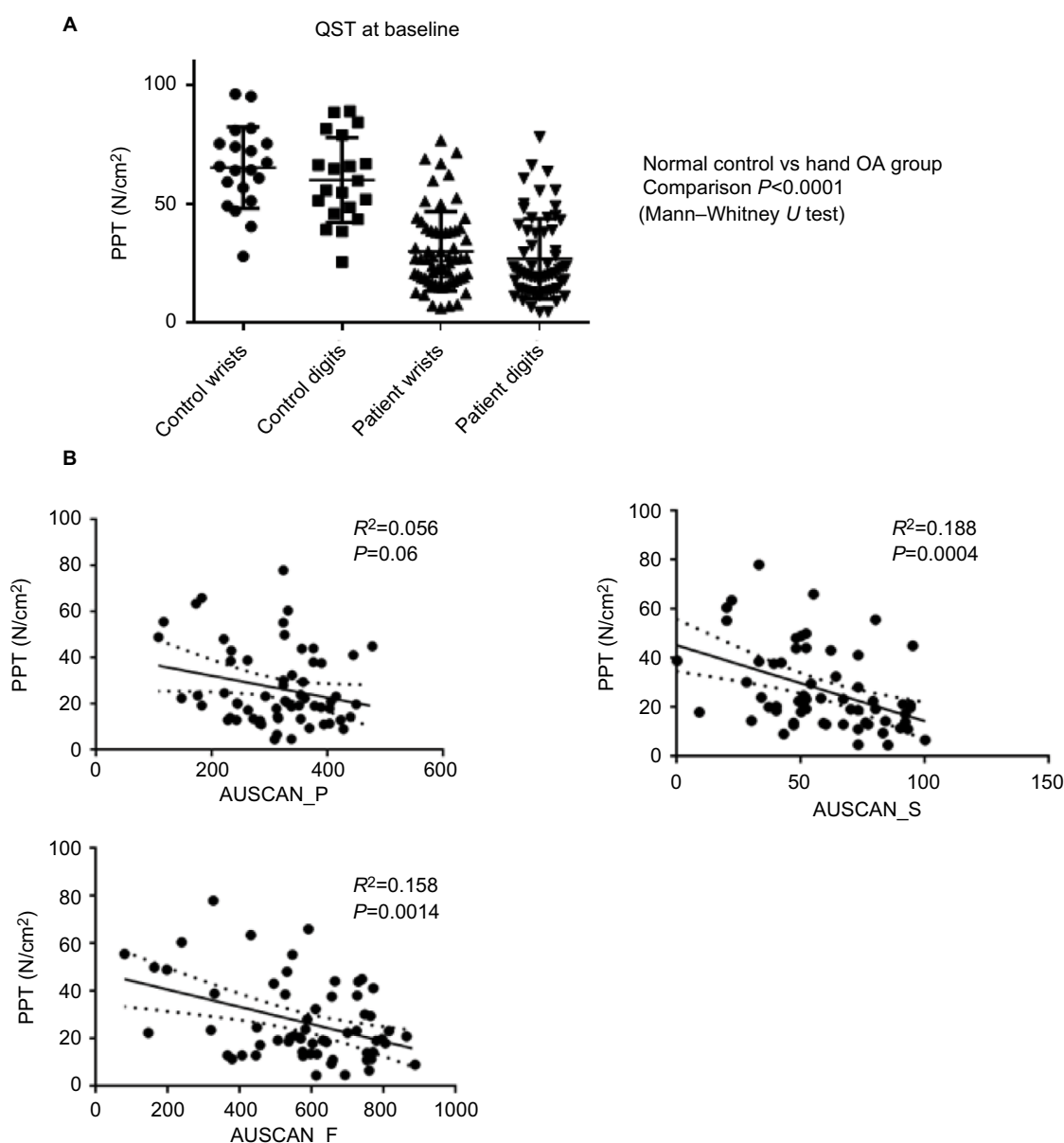
For the comparison of pain pressure algometry (PPT) in non-OA vs OA participants, Mann-Whitney  $U$  was used (Figure 1A). For correlation analyses between AUSCAN scores and PPT, an  $R^2$  correlation and  $P$ -value were calculated using GraphPad Prism (Figure 1B). In the bivariate comparisons of clinical outcome measures and PPT, SPSS was used to calculate an  $R^2$  correlation and  $P$ -value (Table 1).

We used IBM SPSS Statistics 21.0 for all analyses. Graphs were plotted using SPSS or GraphPad Prism Version 7.

## Results

### Characteristics of patient population

Between April 2013 and April 2016, we recruited 65 participants (Figure 2). A total of 21 participants were randomized to duloxetine, a further 22 participants were randomized to pregabalin, and 22 participants were randomized to placebo, respectively. There were 20 age-matched healthy volunteer participants enrolled for the comparison of pain scores using algometry and brain MRI (MRI data from this study will be reported separately). All 65 participants who were randomized to treatment were included in the intention-to-treat (ITT) analysis. A total of 52 participants completed the trial procedures after 13 weeks and were included in the per protocol analyses (Figure 3). Baseline characteristics show that the three treatment groups were well-matched for demographic data (Table 2). The mean disease duration was 3.5 years



**Figure 1** Pain sensitization characteristics of study population.

**Notes:** (A) Data from the DUPRO clinical trial demonstrating reduced pain thresholds globally in the wrist and finger joints in hand OA participants compared to normal age- and sex-matched controls. (B) Graphs demonstrating correlation for PPT in Newton per centimeter squared at baseline with clinical measures for AUSCAN\_P, AUSCAN\_S, and AUSCAN\_F in all groups.

**Abbreviations:** AUSCAN\_P, Australian and Canadian Hand Osteoarthritis Index pain; AUSCAN\_S, Australian and Canadian Hand Osteoarthritis Index stiffness; AUSCAN\_F, Australian and Canadian Hand Osteoarthritis Index function; OA, osteoarthritis; PPT, pain pressure threshold; QST, quantitative sensory testing.

(SD 4.2), which was measured from the time that the participant was first told that they had a diagnosis of hand OA. For prior analgesic use, there was slightly less acetaminophen use at baseline before enrollment in the duloxetine group than in the pregabalin and placebo groups, but for other NSAIDs and opiates, analgesic use was similar in all three groups.

## Patient-reported outcomes

### ITT analysis

Participants in all three groups receiving duloxetine, pregabalin, or placebo reported improvement in pain at the end of the trial.

Comparison of the three groups by ANOVA showed a significant difference at the end of treatment for NRS pain ( $P = 0.035$ ) and AUSCAN pain ( $P = 0.0078$ ) at the end of the trial. Following the primary analysis, pairwise comparisons were performed.

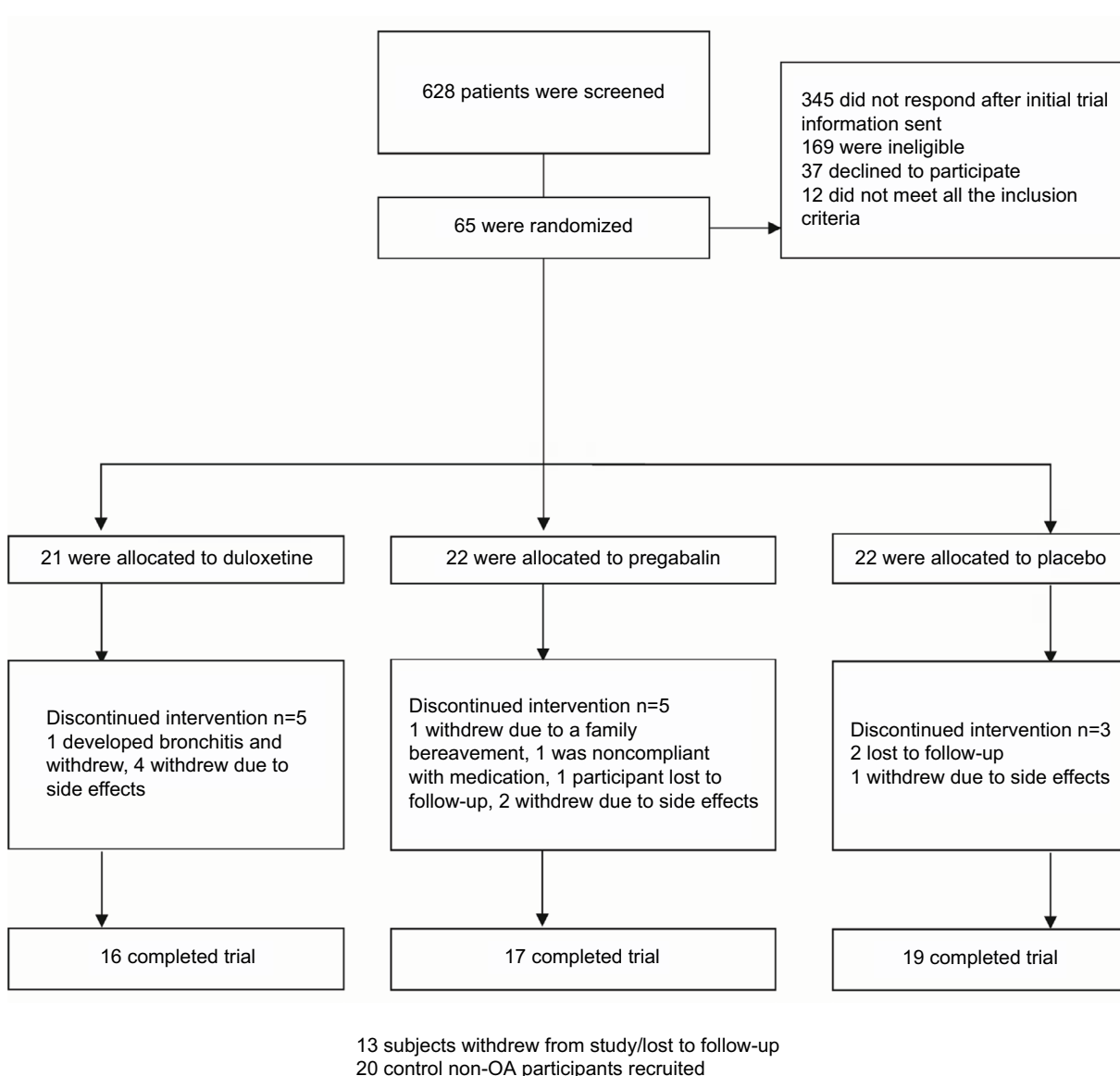
### Pregabalin

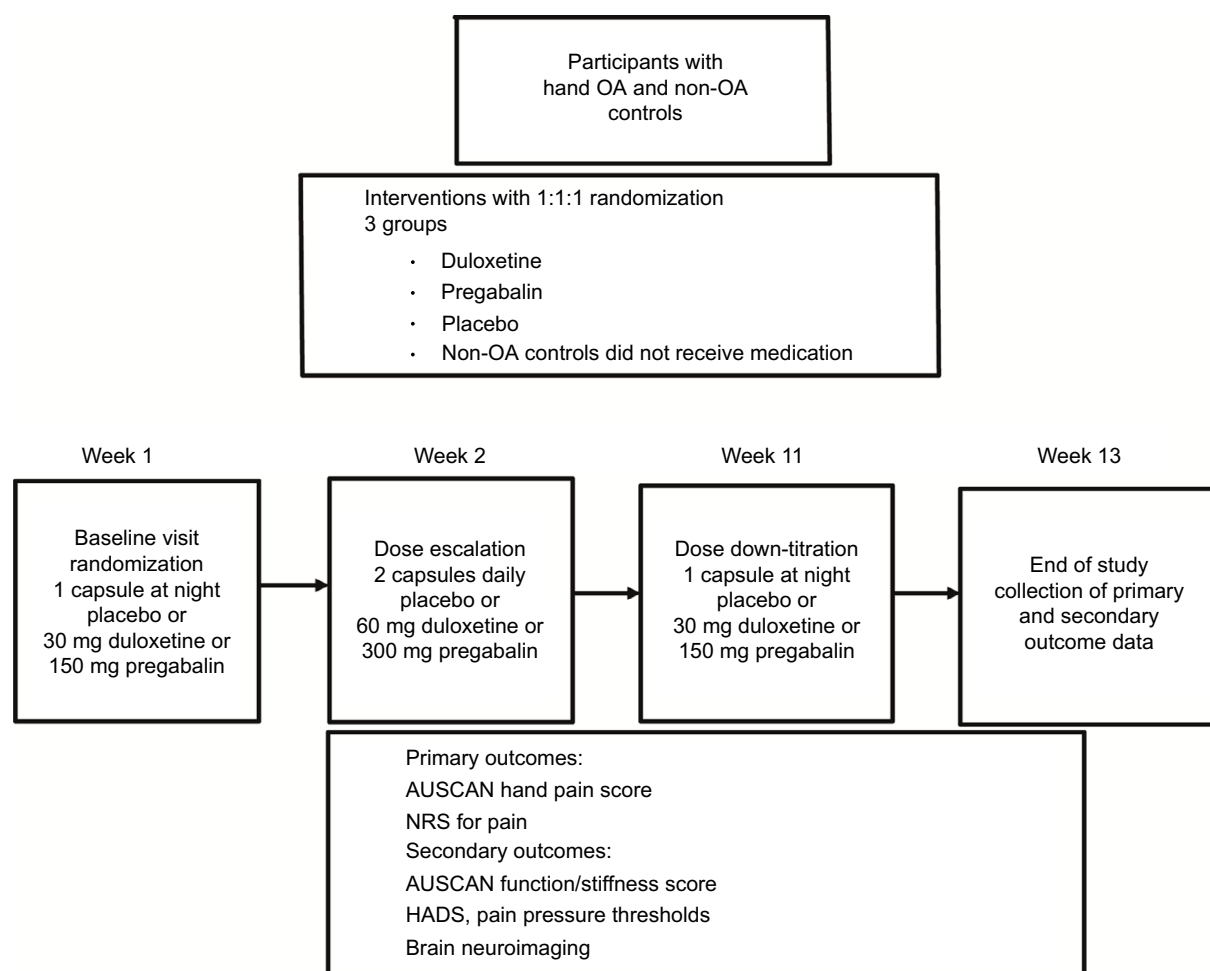
Comparison of pregabalin vs placebo showed a significant improvement in the pregabalin group for primary outcomes of NRS pain ( $P = 0.023$ ), AUSCAN pain ( $P = 0.008$ ), and AUSCAN function ( $P = 0.009$ ) but not AUSCAN stiffness ( $P = 0.22$ ) scores (Table 3 and Figure 4).

**Table 1** Bivariate correlation analysis of clinical scores in study

	AUSCAN_P	AUSCAN_S	AUSCAN_F	QST	HADS_A	HADS_D
NRS	0.606	0.268	0.471	-0.167	-0.010	0.158
	0.000	0.034	0.000	0.203	0.938	0.216
AUSCAN_P		0.251	0.716	-0.234	0.171	0.328
		0.047	0.000	0.072	0.179	0.009
		AUSCAN_S	0.331	-0.429	0.230	0.213
			0.008	0.001	0.070	0.093
			AUSCAN_F	-0.409	0.132	0.294
				0.001	0.304	0.019
				QST	-0.285	-0.187
					0.027	0.152
					HADS_A	0.692
						0.000

**Abbreviations:** AUSCAN\_P, Australian and Canadian Hand Osteoarthritis Index pain; AUSCAN\_S, Australian and Canadian Hand Osteoarthritis Index stiffness; AUSCAN\_F, Australian and Canadian Hand Osteoarthritis Index function; HADS, Hospital Anxiety and Depression Scale; NRS, Numerical Rating Scale; QST, quantitative sensory testing.

**Figure 2** CONSORT flow diagram for the DULoxetine or PRegabalin for Osteoarthritis pain (DUPRO) clinical trial.



**Figure 3** Study flow diagram and outcome measures.

**Abbreviations:** AUSCAN, Australian and Canadian Hand Osteoarthritis Index; HADS, Hospital Anxiety and Depression Scale; NRS, Numerical Rating Scale; OA, osteoarthritis.

**Table 2** Baseline characteristics of study patients (ITT analysis)

Characteristics	Pregabalin (n=22)	Duloxetine (n=21)	Placebo (n=22)
Age (years), mean (SD)	64.0 (5.0)	62.3 (7.3)	62.4 (8.7)
Women	19 (86.4)	14 (66.7)	19 (86.4)
White	20 (90.9)	20 (95.2)	18 (81.8)
Black	2 (9.1)	1 (4.8)	1 (4.6)
Asian			3 (13.6)
Body mass index, mean (SD)	27.1 (6.3)	28.4 (5.9)	27.0 (4.3)
NRS, mean (SD)	6.1 (1.2)	6.4 (1.5)	6.4 (1.4)
AUSCAN pain score, mean (SD)	317.0 (81.4)	296.0 (105.2)	320.3 (66.2)
HADS, mean (SD)	11.6 (7.4)	10.3 (6.1)	12.2 (6.2)
Most common analgesics before inclusion			
Acetaminophen	15	8	15
Other NSAID oral/topical	7	5	5
Codeine-based analgesic	3	4	6

**Note:** Values are numbers (percentages) unless stated otherwise.

**Abbreviations:** AUSCAN, Australian and Canadian Hand Osteoarthritis Index; HADS, Hospital Anxiety and Depression Scale; ITT, intention-to-treat; NRS, Numerical Rating Scale; NSAID, nonsteroidal anti-inflammatory drug.



**Table 3** Primary and secondary outcomes in ITT population

Outcome at 13 weeks	Pregabalin (N=22)	Duloxetine (N=21)	Placebo (N=22)
NRS			
Baseline (95% CI)	6.1 (5.6 to 6.7)	6.4 (5.7 to 7.1)	6.4 (5.7 to 6.9)
13 weeks (95% CI)	3.4 (2.4 to 4.4)	4.3 (2.6 to 5.9)	5.4 (4.1 to 6.8)
Mean difference (95% CI)	-2.7 (-3.5 to -1.9)	-2.3 (-3.8 to -0.9)	-0.9 (-0.2 to 0.2)
P-value	0.023*	0.19	
AUSCAN pain score			
Baseline (95% CI)	317.0 (280.8 to 353.1)	296.0 (248.2 to 343.9)	320.3 (290.9 to 349.6)
13 weeks (95% CI)	176.5 (123.9 to 229.1)	248.1 (162.3 to 333.9)	273.5 (218.0 to 329.0)
Mean difference (95% CI)	-132.1 (-181.1 to -82.9)	-35.8 (-119.7 to 48.2)	-46.6 (-93.9 to 0.75)
P-value	0.008*	0.59	
AUSCAN stiffness			
Baseline (95% CI)	60.18 (51.7 to 68.7)	60.95 (46.98 to 74.9)	55.5 (45.2 to 65.8)
13 weeks (95% CI)	36.5 (23.0 to 49.9)	48.25 (29.87 to 66.6)	50.0 (36.0 to 64.0)
Mean difference (95% CI)	-18.7 (-33.1 to -4.3)	-13.5 (-26.5 to -0.6)	-5.67 (-16.8 to 5.5)
P-value	0.22	0.96	
AUSCAN function			
Baseline (95% CI)	576.2 (499.1 to 653.4)	577.2 (478.0 to 676.4)	582.3 (509.1 to 655.5)
13 weeks (95% CI)	362.2 (281.7 to 442.7)	496.4 (342.4 to 650.5)	508.7 (379.5 to 637.9)
Mean difference (95% CI)	-246.4 (-341.7 to -151.0)	-101.8 (-248.4 to -44.7)	-67.3 (-156.4 to -21.8)
P-value	0.009*	>0.05	
Consumption of rescue medication (total number of days)	9	5	56
HADS			
Anxiety			
Baseline (95% CI)	6.5 (4.6 to 8.4)	5.9 (4.3 to 7.6)	7.2 (5.4 to 9.0)
13 weeks (95% CI)	5.2 (2.9 to 7.5)	4.3 (2.2 to 6.3)	8.2 (6.4 to 9.9)
Mean difference (95% CI)	-0.82 (-2.1 to 0.5)	-1.3 (-3.1 to 0.5)	0.5 (-0.4 to 1.4)
P-value	0.15	0.07	
Depression			
Baseline (95% CI)	5.1 (3.6 to 6.7)	4.4 (2.9 to 5.8)	4.9 (3.6 to 6.2)
13 weeks (95% CI)	4.1 (2.6 to 5.6)	3.8 (1.9 to 5.7)	5.1 (3.9 to 6.3)
Mean difference (95% CI)	-1.1 (-2.1 to -0.02)	-0.3 (-1.9 to 1.2)	0.05 (-1.3 to 1.4)
P-value	0.66	0.99	

**Notes:** \*Indicates significant at <0.05

**Abbreviations:** AUSCAN, Australian and Canadian Hand Osteoarthritis Index; HADS, Hospital Anxiety and Depression Scale; ITT, intention-to-treat; NRS, Numerical Rating Scale.

## Duloxetine

For NRS pain and AUSCAN pain, function, and stiffness outcomes in patients receiving duloxetine compared to placebo, none of these outcomes were significantly different to placebo (Table 3).

## Use of rescue medication

Average use of acetaminophen as rescue medication was much lower in the pregabalin and duloxetine groups than in the placebo group (Table 3). The use of rescue medication in the placebo group was higher, amounting to 56 days.

## Per protocol analysis

There was a reduction in reporting pain in all three groups at the end of the trial. A significant difference between the three groups at the end of treatment for NRS pain score ( $P=0.04$ ) was found by ANOVA. Pairwise comparisons between duloxetine and placebo, pregabalin, and placebo were then performed (Table 4).

## Pregabalin

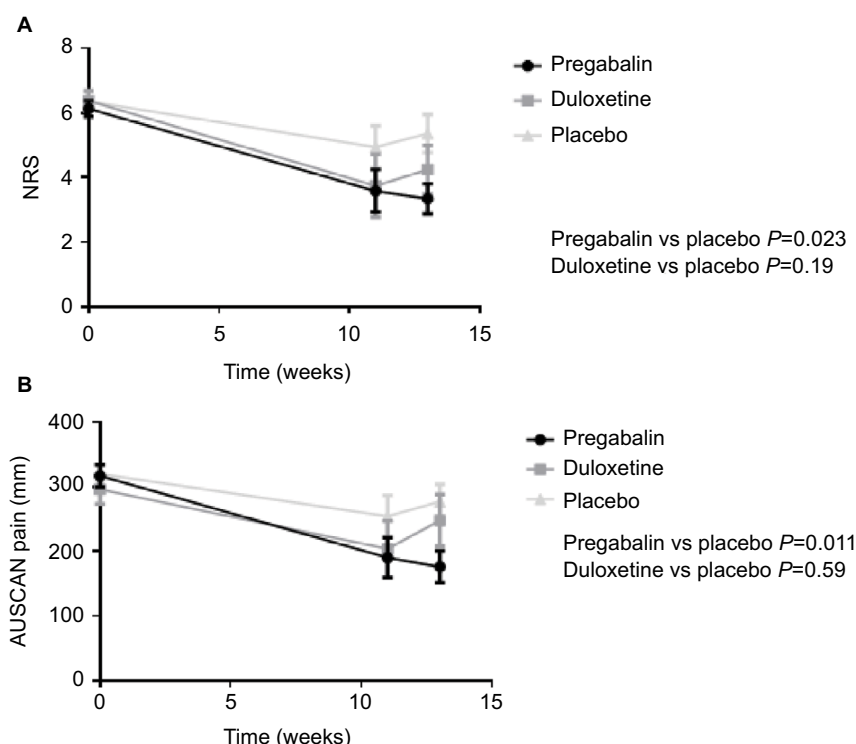
For NRS pain, pregabalin was more effective than placebo ( $P<0.0001$ ). Similarly, compared to placebo, there was a significant improvement in the pregabalin group for AUSCAN pain ( $P=0.013$ ), AUSCAN function ( $P=0.02$ ) but not AUSCAN stiffness ( $P=0.06$ ).

## Duloxetine

For the comparison between placebo and duloxetine treatment, duloxetine was more effective as measured by NRS after 13 weeks ( $P=0.029$ ). For AUSCAN pain, stiffness, and function outcomes in patients receiving duloxetine, these outcomes did not reach statistical significance.

## Adverse events

Side effects were recorded prospectively throughout the study (Table 5). The placebo group showed fewer adverse events with a total of 22 recorded, with no difference in adverse



**Figure 4 (A,B)** Plots for change in primary outcome measures in all treatment groups (ITT analysis).

**Abbreviations:** AUSCAN, Australian and Canadian Hand Osteoarthritis Index; ITT, intention-to-treat; NRS, Numerical Rating Scale.

events between the three groups ( $P=0.73$ ). The highest reporting of adverse events was observed in the pregabalin and duloxetine groups: 55 adverse events were recorded with pregabalin, the most common of which were mental disturbance, headaches, sleepiness, dizziness, and dry mouth. In the duloxetine group, a total of 57 adverse events were recorded; there were a total of four withdrawals due to drug side effects and one participant withdrew due to the development of bronchitis, as shown in the CONSORT flow diagram (Figures 3 and 5). For the pregabalin group, there were two withdrawals due to drug side effects, one withdrawal due to a family bereavement, one withdrawal due to noncompliance, and one withdrawal due to loss to follow-up.

### Pain sensitization by PPT and relation to clinical scores

Using PPT testing as a measure for pain sensitization, compared to non-OA controls, the hand arthritis group had globally reduced pain thresholds ( $P<0.0001$ ) across all finger joints at baseline, even at the metacarpophalangeal joints and wrists where there was little evidence of radiographic OA (Figure 1A). We investigated the correlation between the various clinical scores at baseline with age as a covariate (Table 1). Measurements for the PPT modality of QST showed a significant correlation with AUSCAN stiffness ( $R^2=0.188$ ,  $P=0.0004$ ) and function ( $R^2=0.158$ ,  $P=0.0014$ )

for all patients at baseline (Figure 1B). There was a trend for lower PPT correlated with higher AUSCAN pain scores, although this trend did not reach statistical significance ( $P=0.06$ ). NRS pain correlated strongly with AUSCAN pain and function ( $P<0.0001$ ). PPT measures did not change significantly in any of the three groups after 3 months treatment. We found that all participants with hand OA had lower PPT scores at baseline compared to healthy controls at inclusion and demonstrated a reduction of NRS at follow-up. HADS anxiety and depression scores were significantly correlated after Bonferroni correction. There were weaker correlations (significant without correction) between HADS depression and AUSCAN pain ( $P=0.009$ ) and between AUSCAN function and stiffness scores ( $P=0.008$ ).

## Discussion

### Principal findings

Our clinical study provides the first evidence in chronic painful hand OA that pregabalin and duloxetine are analgesics with potential for use in OA pain, with pregabalin providing the best treatment response and sustained effects beyond the reduction in dose. Second, we observed by QST that hand arthritis subjects have pain sensitization, which may include peripheral and central mechanisms. Third, the central but distinct actions of pregabalin and duloxetine could therefore be exerting an effect on central pain sensitization, which we



**Table 4** Summary table for per protocol analysis

Outcome at 13 weeks (imputed data on per protocol set)	Pregabalin	Duloxetine	Placebo
<b>NRS</b>			
Baseline (95% CI)	6.1 (5.4 to 6.7)	6.6 (5.7 to 7.4)	6.3 (5.6 to 6.9)
13 weeks (95% CI)	3.4 (2.4 to 4.4)	4.3 (2.6 to 5.9)	5.4 (4.1 to 6.6)
Mean difference (95% CI)	-2.7 (-3.5 to -1.9)	-2.3 (-3.8 to -0.9)	-0.9 (-2.3 to 0.2)
P-value	<0.0001*	0.029*	
<b>AUSCAN pain score</b>			
Baseline (95% CI)	308.5 (262.6 to 354.5)	310.6 (254.3 to 367.0)	321.1 (288.7 to 353.4)
13 weeks (95% CI)	176.5 (123.9 to 229.1)	248.1 (162.3 to 333.9)	273.5 (218.0 to 329.0)
Mean difference (95% CI)	-132.0 (-181.1 to -82.9)	-62.5 (-141.6 to 16.6)	-47.1 (-93.8 to 11.7)
P-value	0.013*	0.9	
<b>AUSCAN stiffness</b>			
Baseline (95% CI)	59.9 (51.8 to 67.9)	61.8 (45.8 to 77.8)	56.1 (44.5 to 67.7)
13 weeks (95% CI)	36.5 (23.0 to 49.9)	48.3 (29.9 to 66.6)	50.0 (36.0 to 64.0)
Mean difference (95% CI)	-23.4 (-35.7 to -11.1)	-13.5 (-26.5 to -0.6)	5.7 (-16.8 to 5.5)
P-value	0.06	0.46	
<b>AUSCAN function</b>			
Baseline (95% CI)	608.5 (541.3 to 675.7)	598.3 (481.2 to 715.3)	580.0 (494.6 to 665.4)
13 weeks (95% CI)	362.2 (281.7 to 442.7)	496.4 (342.4 to 650.5)	508.7 (379.5 to 637.9)
Mean difference (95% CI)	-246.3 (-341.7 to -151.0)	-101.9 (-248.4 to -44.8)	-69.7 (-158.3 to -18.9)
P-value	0.02*	0.93	
Consumption of rescue medication (total number of days)	9	5	56
<b>HADS</b>			
<b>Anxiety</b>			
Baseline (95% CI)	6.1 (3.8 to 8.3)	5.6 (3.6 to 7.5)	7.6 (5.8 to 9.5)
13 weeks (95% CI)	5.2 (2.9 to 7.5)	4.3 (2.2 to 6.3)	8.2 (6.4 to 9.9)
Mean difference (95% CI)	-0.82 (-2.1 to 0.5)	-1.3 (-3.1 to 0.5)	0.5 (-0.4 to 1.4)
P-value	0.52	0.21	0.71
<b>Depression</b>			
Baseline (95% CI)	5.1 (3.3 to 6.9)	4.1 (2.3 to 5.9)	5.4 (4.0 to 6.7)
13 weeks (95% CI)	4.1 (2.6 to 5.6)	3.8 (1.9 to 5.7)	5.1 (3.9 to 6.3)
Mean difference (95% CI)	-1.1 (-2.1 to -0.02)	-0.3 (-1.8 to 1.2)	0.05 (-1.3 to 1.4)
P-value	0.41	0.54	0.93

**Notes:** \*Indicates significant at <0.05

**Abbreviations:** AUSCAN, Australian and Canadian Hand Osteoarthritis Index; HADS, Hospital Anxiety and Depression Scale; NRS, Numerical Rating Scale.

and others have demonstrated as a significant component of arthritic pain.<sup>14,15</sup> Finally, we observed improvement for pregabalin in NRS for the ITT analysis, but not for duloxetine, with improvement in NRS for both active drugs only in the per protocol analysis. The results of our trial have strong clinical relevance, since many patients report lack of efficacy or side effects on NSAIDs and other patients have important safety concerns.

## Study strengths and limitations

The lack of new analgesic targets for OA in this most common arthritic disease, coupled with recent data from animal models,<sup>18,19</sup> prompted us to investigate the use of the gabapentinoid pregabalin and the SNRI duloxetine. Pregabalin is licensed for neuropathic pain<sup>29</sup> and duloxetine for depression and diabetic neuropathic pain.<sup>30</sup> Our proof-of-concept trial demonstrated an improvement in pain for

pregabalin and also for duloxetine after 13 weeks treatment. We enrolled subjects who had an NRS pain rating of at least 5 to ensure that clinically meaningful improvements in pain could be detected. There were some differences in our ITT and per protocol analysis: in ITT, pregabalin, but not duloxetine, showed a significant improvement in pain compared to placebo; and in the per protocol analysis, both agents showed an improvement in pain. Although we observed an improvement in pain reporting for both centrally acting agents, pregabalin was more effective after 13 weeks. In our secondary endpoint analyses, we did not see any significant improvement in depression or anxiety scores in any treatment group.

Since this was a proof-of-concept analgesic endpoint study, we did not collect structural outcome data including joint damage progression changes by plain radiograph and synovitis by ultrasound, as described in other studies,<sup>31–34,37</sup>

**Table 5** Side effect profile in all three treatment groups from ITT analysis

System	Pregabalin (N=22)	Duloxetine (N=21)	Placebo (N=22)
Cardiovascular	3	2	1
Digestive	7	18	5
ENT		2	
Endocrine/metabolic	1		
Genitourinary	1		
Hematological			
Mental	9	9	9
Nervous system			
Dry mouth	6	6	4
Headaches	3	8	
Dizziness	7	3	
Sleepiness	5	3	
Loss of balance	7		
Ophthalmological	4	2	1
Respiratory	2	3	
Skin		1	2
Total	55	57	22

**Abbreviations:** ENT, ear, nose and throat; ITT, intention-to-treat.

which could be addressed in future work. We also recognize that our study has been conducted in hand arthritis pain, whereas several large previous datasets have focused on knee OA,<sup>35–37</sup> and there may be some differences in pain characteristics including loading effects due to structural joint differences between the hand and knee.

## Main results in context of other literature

In knee OA, Chappell et al<sup>17</sup> showed that duloxetine was effective for pain, Ohtori et al<sup>20</sup> found that pregabalin with meloxicam was more effective than pregabalin alone, and our data show that both agents have efficacy in chronic hand OA pain with pregabalin showing superiority over duloxetine.

Recent concepts in novel therapeutic agents for OA have included potential therapeutics for structural changes in the joint including synovitis<sup>31</sup> and bone marrow lesions (BML).<sup>32</sup>

	Item number	Checklist item	Page number*
<b>Title and abstract</b>			
	1a	Identification as a randomized trial in the title Yes	Title
	1b	Structured summary of trial design, methods, results, and conclusions Yes	Abstract
<b>Introduction</b>			
Background and objectives	2a	Scientific background and explanation of rationale Yes	Introduction Paragraphs 1–3
	2b	Specific objectives or hypotheses Yes	Introduction Paragraph 4
<b>Methods</b>			
Trial design	3a	Description of trial design (such as parallel and factorial) including a location ratio Yes	Methods, trial design, and participants section
	3b	Important changes to methods after trial commencement (such as eligibility criteria), with reasons Yes	Methods and statistical analysis section
Participants	4a	Eligibility criteria for participants Yes	Methods, study design, and participants section
	4b	Settings and locations where the data were collected Yes	Methods, study design, and participants section
Interventions	5	The interventions for each group with sufficient details to allow replication, including how and when they were actually administered Yes	Methods and study design
Outcomes	6a	Completely defined pre-specified primary and secondary outcome measures, including how and when they were assessed Yes	Methods and outcome section
	6b	Any changes to trial outcomes after the trial commenced, with reasons	Not applicable
Sample size	7a	How sample size was determined Yes	Methods and statistical analysis section
	7b	When applicable, explanation of any interim analyses and stopping guidelines	Not applicable
<b>Randomization</b>			
Sequence generation	8a	Method used to generate the random allocation sequence Yes	Methods, randomization, and masking section
	8b	Type of randomization; details of any restriction (such as blocking and block size) Yes	Methods, randomization, and masking section

**Figure 5** (Continued)

	Item number	Checklist item	Page number*
Allocation concealment mechanism	9	Mechanism used to implement the random allocation sequence (such as sequentially numbered containers), describing any steps taken to conceal the sequence until interventions were assigned Yes	Methods, randomization, and masking section
Implementation	10	Who generated the random allocation sequence, who enrolled participants, and who assigned participants to interventions Yes	Methods, randomization, and masking section
Blinding	11a	If done, who was blinded after assignment to interventions (for example, participants, care providers, those assessing outcomes) and how Yes	Methods, randomization, and masking section
	11b	If relevant, description of the similarity of interventions	Not applicable
Statistical methods	12a	Statistical methods used to compare groups for primary and secondary outcomes Yes	Methods and statistical analysis section
	12b	Methods for additional analyses, such as subgroup analyses and adjusted analyses Yes	Methods and statistical analysis section
<b>Results</b>			
Participant flow (a diagram is strongly recommended)	13a	For each group, the numbers of participants who were randomly assigned, received intended treatment, and were analyzed for the primary outcome Yes	Figure 2 CONSORT diagram
	13b	For each group, losses and exclusions after randomization, together with reasons Yes	Figure 2 CONSORT diagram
Recruitment	14a	Dates defining the periods of recruitment and follow-up Yes	Methods, study design, and participants section
	14b	Why the trial ended or was stopped	Not applicable
Baseline data	15	A table showing baseline demographic and clinical characteristics for each group Yes	Table 2 from paper
Numbers analyzed	16	For each group, number of participants (denominator) included in each analysis and whether the analysis was by original assigned groups Yes	Results, participant section, and CONSORT diagram, Figure 2
Outcomes and estimation	17a	For each primary and secondary outcome, results for each group, and the estimated effect size and its precision (such as 95% confidence interval) Yes	Table 3 from paper
	17b	For binary outcomes, presentation of both absolute and relative effect sizes is recommended	Not a binary outcome
Ancillary analyses	18	Results of any other analyses performed, including subgroup analyses and adjusted analyses, distinguishing prespecified from exploratory Yes	Results and Figure 1
Harms	19	All important harms or unintended effects in each group Yes	Results, adverse events section, and Table 5
<b>Discussion</b>			
Limitations	20	Trial limitations, addressing sources of potential bias, imprecision, and, if relevant, multiplicity of analyses Yes	Discussion, study strengths, and limitations
Generalizability	21	Generalizability (external validity, applicability) of the trial findings Yes	Discussion, study strengths, and limitations, implications for practice and future research
Interpretation	22	Interpretation consistent with results, balancing benefits and harms, and considering other relevant evidence Yes	Discussion, section main results in context of other literature
<b>Other information</b>			
Registration	23	Registration number and name of trial registry Yes	Abstract
Protocol	24	Where the full trial protocol can be accessed, if available Yes	Clinicaltrials.gov
Funding	25	Sources of funding and other support (such as supply of drugs), role of funders Yes	Included in declaration submission

**Figure 5** CONSORT 2010 checklist of information for the DULoxetine or PRegabalin for Osteoarthritis pain (DUPRO) randomized controlled trial.

**Note:** \*Page numbers optional depending on journal requirements. Hopewell S, Clarke M, Moher D, Wager E, Middleton P, Altman DG, et al. CONSORT for reporting randomised trials in journal and conference abstracts. *Lancet*. 2008;371:281–283.<sup>38</sup>

However, such studies have not been without difficulty since recent trials targeting the inflammatory component of OA have not shown improved outcomes<sup>33</sup> and the use of bisphosphonates potentially for reducing BML-related pain need to define significant clinical and structural endpoints.<sup>34</sup> It is possible that patients demonstrating a largely “inflammatory” phenotype are likely to benefit from agents such as NSAIDs, and when there are features of sensitization with ongoing pain, patients may require additional treatment such as centrally acting agents including pregabalin and duloxetine. In the clinic, patients may also require additional treatments if NSAIDs are linked to side effects, lack of efficacy, and ongoing pain.

There is recognition that pain sensitization occurs in people with OA.<sup>11–15,35–37</sup> The main indication from our data of peripheral sensitization in hand OA is that the control subjects had significantly higher PPTs than the hand OA group. We noted that PPTs did not change significantly after treatment, suggesting that pathways which led to sensitization in hand arthritis may continue to exist in the patients even after drug treatment.

## Implications for practice and future research

Pregabalin and duloxetine had efficacy in hand OA pain in our clinical study, with pregabalin showing greater effect than duloxetine for validated pain endpoints. In our study, one or more of the following analgesics had been used by more than half of the participants prior to enrollment in the study: acetaminophen, NSAID, or codeine-based analgesics. When such analgesics had not previously been effective, our trial showed that pregabalin, and to a less significant degree duloxetine, may provide a realistic alternative to pain management in OA. In future, clinical trials that examine the efficacy of centrally acting analgesics over a longer treatment period of >12 weeks in chronic arthritic pain should be conducted. Further studies measuring peripheral and central sensitization will be crucial to understand how pain, loss of function, comorbid conditions, and medication use contribute to the development of arthritic pain.

## Acknowledgments

This work was supported by The Rosetrees' Trust, grant number M11-F1, by the UK National Institute of Health (NIHR) Clinical Research Network and an NIHR Clinical Academic Fellowship to MR. The views expressed are those of the authors and not necessarily those of the NHS, NIHR, or Department of Health.

## Author contributions

All authors contributed toward data analysis, drafting and critically revising the paper and agree to be accountable for all aspects of the work.

## Disclosure

The authors report no conflicts of interest in this work.

## References

1. Dahaghin S, Bierma-Zeinstra SMA, Reijman M, Pols HAP, Hazes JMW, Koes BW. Prevalence and determinants of one month hand pain and hand related disability in the elderly (Rotterdam study). *Ann Rheum Dis*. 2005;64(1):99–104.
2. Felson DT, Lawrence RC, Dieppe PA, et al. Osteoarthritis: new insights. Part 1: the disease and its risk factors. *Ann Intern Med*. 2005;13(8):635–646.
3. Bijlsma JW, Berenbaum F, Lafaber FP. Osteoarthritis: an update with relevance for clinical practice. *Lancet*. 2011;18(377):2115–2126.
4. NICE [webpage on the Internet]. *NICE Guidelines 'Osteoarthritis: Care and Management'*; 2014. Available from: <https://www.nice.org.uk/guidance/cg177>. Accessed May 18, 2017.
5. Zhang W, Doherty M, Leeb BF, et al. EULAR evidence based recommendations for the management of hand osteoarthritis: report of a Task Force of the EULAR Standing Committee for international clinical studies including therapeutics (ESCISIT). *Ann Rheum Dis*. 2007;66(3):377–388.
6. Da Costa BR, Reichenbach S, Keller N, et al. Effectiveness of non-steroidal anti-inflammatory drugs for the treatment of pain in knee and hip osteoarthritis: a network meta-analysis. *Lancet*. 2016;387:2093–2105.
7. Williams CM, Maher CG, Latimer J, et al. Efficacy of paracetamol for acute low-back pain: a double-blind, randomised controlled trial. *Lancet*. 2014;384(9954):1586–1596.
8. Woolf CJ. Pain: Moving from symptom control toward mechanism-specific pharmacologic management. *Ann Intern Med*. 2004;140:441–451.
9. Kidd BL, Urban LA. Mechanisms of inflammatory pain. *Br J Anaesth*. 2001;87:3–11.
10. Suokas AK, Walsh DA, McWilliams DF, et al. Quantitative sensory testing in painful osteoarthritis: a systematic review and meta-analysis. *Osteoarthritis Cartilage*. 2012;20(10):1075–1085.
11. Arendt-Nielsen L, Nie H, Laursen MB, et al. Sensitization in patients with painful knee osteoarthritis. *Pain*. 2010;149(3):573–581.
12. Sofat N, Ejindu V, Kiely P. What makes OA painful? The evidence for peripheral and central pain processing. *Rheumatology*. 2011;50:2157–2165.
13. Neogi T, Frey-Law L, Scholz J, et al; Multicenter Osteoarthritis (MOST) Study. Sensitivity and sensitisation in relation to pain severity in knee osteoarthritis: trait or state? *Ann Rheum Dis*. 2015;74(4):682–688.
14. Sofat N, Smee C, Hermansson M, et al. Functional MRI demonstrates pain perception in hand osteoarthritis has features of central pain processing. *J Biomed Graph Comput*. 2013;3(4):20–26.
15. Gwilym S, Filippini N, Douaud G, Carr A, Tracey I. Thalamic atrophy associated with painful osteoarthritis of the hip is reversible after arthroplasty: a longitudinal voxel-based morphometric study. *Arthritis Rheum*. 2010;62(10):2930–2940.
16. Abaei M, Sagar DR, Stockley EG, et al. Neural correlates of hyperalgesia in the monosodium iodoacetate model of osteoarthritis pain. *Mol Pain*. 2016;12:1–12.
17. Chappell AS, Ossanna MJ, Liu-Seifert H, et al. Duloxetine, a centrally acting analgesic in the treatment of patients with osteoarthritis knee pain: a 13 week, randomized, placebo-controlled trial. *Pain*. 2009;146:253–260.
18. Park J, Yu YP, Zhou CY, et al. Central mechanisms mediating thrombospondin-4-induced pain states. *J Biol Chem*. 2016;291(25):13335–13348.

19. Pan B, Guo Y, Wu HE, et al. Thrombospondin-4 divergently regulates voltage-gated Ca<sup>2+</sup> channel subtypes in sensory neurons after nerve injury. *Pain*. 2016;157(9):2068–2080.
20. Ohtori S, Inoue G, Orita S, et al. Efficacy of combination of meloxicam and pregabalin for pain in knee osteoarthritis. *Yonsei Med J*. 2013;54(5):1253–1258.
21. Arendt-Nielsen L, Lindhardt Egshard L, Petersen KK. Evidence for a central mode of action for COX-2 inhibitor in patients with painful knee osteoarthritis. *Pain*. 2016;157:1634–1644.
22. Altman R, Alarcon G, Appelrouth D, et al. The American College of Rheumatology Criteria for the classification and reporting of osteoarthritis of the hand. *Arthritis Rheum*. 1990;33(11):1601–1610.
23. Kellgren JH, Lawrence JS. Radiological assessment of osteo-arthritis. *Ann Rheum Dis*. 1957;16(4):494–502.
24. Bellamy N, Hochberg M, Tubach F, et al. Development of multinational definitions of minimal clinically important improvement and patient acceptable symptomatic state in osteoarthritis. *Arthritis Care Res (Hoboken)*. 2015;67(7):972–980.
25. Dworkin RH, Turk DC, Farrar JT, Haythornthwaite JA, Jensen MP, Katz NP. Core outcome measures for chronic pain clinical trials: IMMPACT recommendations. *Pain*. 2005;113:9–19.
26. Bjelland I, Dahl AA, Haug TT, Neckelmann D. The validity of the Hospital Anxiety and Depression Scale: an updated literature review. *J Psychosom Res*. 2002;52(2):69–77.
27. Wajed J, Hermansson M, Axford J, Sofat N. Pain threshold testing using algometers is a reliable measure of pain in hand osteoarthritis and correlates with patient-reported pain scores. *Rheumatology*. 2011;50:iii91.
28. Kloppenburg M, Maheu E, Kraus VB, et al. OARSI clinical trials recommendations for RCTs. *Osteoarthritis Cartilage*. 2015;23:772–786.
29. Freynhagen R, Serpell M, Emir B, et al. A comprehensive drug safety evaluation of pregabalin in peripheral neuropathic pain. *Pain Pract*. 2015;15(1):47–57.
30. Gierthmulen J, Baron R. Neuropathic pain. *Semin Neurol*. 2016;36(5):462–468.
31. Kingsbury SR, Tharmanathan P, Arden NK, et al. Pain reduction with oral methotrexate in knee osteoarthritis, a pragmatic Phase III trial of treatment effectiveness (PROMOTE): study protocol for a randomized controlled trial. *Trials*. 2015;16:77.
32. Bruyere O, Cooper C, Arden N, et al. Can we identify patients with high risk of osteoarthritis progression who will respond to treatment? A focus on epidemiology and phenotype of osteoarthritis. *Drugs Aging*. 2015;32:179–187.
33. Basoki N, Lee W, Ruijgrok L, et al. Efficacy of hydroxychloroquine in primary hand osteoarthritis: a randomized, double-blind, placebo controlled trial. *Ann Rheum Dis*. 2015;74:188.
34. Davis AJ, Smith TO, Hing CB, Sofat N. Are bisphosphonates effective in the treatment of osteoarthritis pain? A research synthesis and meta-analysis. *PLoS One*. 2013;8:e72714.
35. Wylde V, Palmer S, Learmonth ID, Dieppe P. Test-retest reliability of quantitative sensory testing in knee osteoarthritis and healthy participants. *Osteoarthritis Cartilage*. 2011;19(6):655–658.
36. Arendt-Nielsen L, Skou ST, Nielsen TA, Petersen KK. Altered central sensitization and pain modulation in the CNS in chronic joint pain. *Curr Osteoporosis Rep*. 2015;13(4):225–234.
37. Felson DT, Niu J, Neogi T, et al; MOST Investigators Group. Synovitis and the risk of knee osteoarthritis: the MOST study. *Osteoarthritis Cartilage*. 2016;24(3):458–464.
38. Hopewell S, Clarke M, Moher D, Wager E, Middleton P, Altman DG, et al. CONSORT for reporting randomised trials in journal and conference abstracts. *Lancet*. 2008;371:281–283.

## Journal of Pain Research

### Publish your work in this journal

The Journal of Pain Research is an international, peer reviewed, open access, online journal that welcomes laboratory and clinical findings in the fields of pain research and the prevention and management of pain. Original research, reviews, symposium reports, hypothesis formation and commentaries are all considered for publication.

Submit your manuscript here: <https://www.dovepress.com/journal-of-pain-research-journal>

Dovepress

The manuscript management system is completely online and includes a very quick and fair peer-review system, which is all easy to use. Visit <http://www.dovepress.com/testimonials.php> to read real quotes from published authors.