Attenuation of early phase inflammation by cannabidiol prevents pain and nerve damage in rat osteoarthritis
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Abstract
Osteoarthritis (OA) is a multifactorial joint disease, which includes joint degeneration, intermittent inflammation, and peripheral neuropathy. Cannabidiol (CBD) is a non-psychoactive constituent of cannabis that has the potential to relieve pain. The aim of this study was to determine whether CBD is anti-nociceptive in OA, and whether inhibition of inflammation by CBD could prevent the development of OA pain and joint neuropathy. Osteoarthritis was induced in male Wistar rats (150-175 g) by intra-articular injection of sodium monolaurate (MIA; 3 mg). On day 14 (end-stage OA), joint afferent mechanosensitivity was assessed using in vivo electrophysiology, whereas pain behaviour was measured by von Frey hair algometers and dynamic incapacitation. To investigate acute joint inflammation, blood flow and leukocyte trafficking were measured on day 1 after MIA. Joint nerve myelination was calculated by G-ratio analysis. The therapeutic and prophylactic effects of peripheral CBD (100-300 μg) were assessed. In end-stage OA, CBD dose-dependently decreased joint afferent firing rate, and increased withdrawal threshold and weight bearing (P ≤ 0.0001; n = 8). Acute, transient joint inflammation was reduced by local CBD treatment (P ≤ 0.0001; n = 6). Prophylactic administration of CBD prevented the development of MIA-induced joint pain at later time points (P ≤ 0.0001; n = 8), and was also found to be neuroprotective (P ≤ 0.05; n = 6-8). The data presented here indicate that local administration of CBD blocked OA pain. Prophylactic CBD treatment prevented the later development of pain and nerve damage in these OA joints. These findings suggest that CBD may be a safe, useful therapeutic for treating OA joint neuropathic pain.

Keywords: Cannabinoids, Osteoarthritis, Pain, Neuropathy, Inflammation

1. Introduction
The most prominent form of synovial joint disease, osteoarthritis (OA), is characterised by joint degeneration, pain, and in some patients, articular neuropathy.1,2 Chronic pain associated with OA is a major concern for which there are few viable treatments. The first-line therapy used to treat OA pain is nonsteroidal anti-inflammatory drugs; however, with long-term use their efficacy declines and they can lead to major adverse gastrointestinal and cardiovascular events. Historically, OA has been classified as noninflammatory arthritis; however, there is now overwhelming evidence that synovitis can occur in response to pro-inflammatory mediators being released into the joint.3,4,5,6,7,8,9,10,11,12,13,14,15,16,17,18,19,20,21,22,23,24,25,26,27,28,29,30

It is believed that this low-level inflammation contributes to degenerative changes that affect the entire joint leading to the development of peripheral sensitisation and nociceptive pain.31,32,33,34 In addition to structural defects, there is growing evidence to suggest that approximately 30% of patients with OA have neuropathic pain.33,34 Thus, a therapeutic which can block inflammation, neuropathy, and pain is sorely needed.

The endocannabinoid system (ECS) plays an important physiological role in the regulation of tissue inflammation and pain.35,36 A functional ECS has been demonstrated in the joints of animals37,38 and humans,39 which acts tonically to maintain joint homeostasis. Immunohistological and pharmacological evidence confirm that cannabinoid 1 (CB1) and cannabinoid 2 (CB2) receptors are expressed on the neurons and microvasculature that supply rat knee joints.39,40 In addition, CB2 receptors are co-localized with pronociceptive transient receptor potential vanilloid-1 (TRPV1) channels where, through common intracellular pathways, they act together to modulate joint pain.39,40,41 This suggests that drugs which target the ECS have the potential to regulate painful arthritis and inflammatory joint disease.

Cannabidiol is the major non-psychoactive component of the cannabis plant.33,34 Pharmacologically, CBD has a complex signalling mechanism whereby it can both activate and silence classical cannabinoid receptors as well as modulate noncanonical cannabinoid receptor pathways. In vitro studies, CBD has been shown to be an inverse agonist at CB2 receptors, and a full agonist at CB1 receptors.40,41 and G protein-coupled receptor-55 (GPR55).33 In vitro, CBD was found to be an agonist at TRPV1 and transient receptor potential ankyrin 1 (TRPA1),42 which play...
a central role in the development of OA,27 in musculoskeletal disease models, systemic administration of CBD suppressed the progression of collagen-induced arthritis by reducing inflammatory cytokine production.28 Although these preliminary findings indicate a possible role for CBD in relieving joint pain, the local effect of articularly applied CBD on OA and joint pain has not been investigated.

The initial aim of this study was to assess the effect of locally administered CBD on joint pain in animals with end-stage OA. Since acute inflammation can contribute to the long-term development of OA joint pain,29 the ability of CBD to reduce acute OA synovitis and prevent the subsequent progression of persistent OA pain was also investigated. Finally, the effect of prophylactic CBD treatment on OA joint neuropathy was assessed.

2. Methods

2.1. Animals

Male Wistar rats (150-175 g; Charles River Laboratories, Senneville, QC, Canada) were housed in ventilated racks at 22°C ± 2°C on a 12:12 hours light-dark cycle (light-on from 7:00 to 19:00). After arrival at the animal care facility, all rats were permitted at least 1 week to acclimatize to their environment. Animals were housed in pairs, cages were lined with woodchip bedding, and animals were provided with environmental enrichment. Standard laboratory chow and water were provided ad libitum. All experimental protocols were approved by the Dalhousie University Committee on the Use of Laboratory Animals, which acts in accordance with Animal Research: Reporting of In Vivo Experiments (ARRIVE) and the standards put forth by the Canadian Council for Animal Care.

2.2. Sodium monoiodoacetate model of osteoarthritis

Animals were deeply anaesthetised (2%-4% isoflurane; 100% oxygen at 1 L/min) until cessation of all sensory reflexes. The right knee joint was shaved, swabbed with 100% ethanol and 50 μL of sodium monoiodoacetate (MIA) (3 mg in saline) was injected into the joint space (intra-articular; Laric.). The knee was then manually extended and flexed for 30 seconds to disperse the solution throughout the joint.

2.3. Electrophysiological recording of joint afferents

After OA development (14-19 days after MIA), animals were deeply anaesthetised using urethane (25% solution; 2 g/kg i.p.). Core body temperature was measured by a rectally inserted thermometer and maintained at 37°C ± 1°C by a thermostatically controlled heating blanket (CWE Inc, Archmore, PA). After loss of the pedal withdrawal reflex, the trachea was cannulated to allow for artificial ventilation with a Harvard rodent respiratory pump (Harvard Apparatus, Holliston, MA) with 100% O2 (stroke volume: 2.5 ml; breath frequency: 52 breaths/min). The left carotid artery was cannulated to allow for continuous measurement of the mean arterial blood pressure. The cannula was attached to an in-line pressure transducer (Kent Scientific Corp, Torrington, CT) attached to a differentially amplified blood pressure monitor (World Precision Instruments, Sarasota, FL). The jugular vein was cannulated for administration of the muscle relaxant gallamine triethiodide (50 mg/kg), which eliminated neural interference from hind limb musculature, and the distal saphenous artery was cannulated for close intra-arterial (i.a.) administration of CBD or vehicle to the knee joint (100 μL injection volume). A specialised clamp was fixed to the mid-shaft of the isolated right femur and attached to a stereotaxic frame to prevent movement of the proximal aspect of the rat hind limb. The right hind paw was then placed in a shoe-like holder that was connected to a force transducer and torque meter (Data Track 244-1-RT, Intertechnology, ON, Canada) to standardise the amount of rotational force being applied to the knee joint. A longitudinal skin incision was made along the medial aspect of the hind limb and the reflected skin was sutured to a metal “O” ring to create a pool which was filled with warm mineral oil to prevent tissue desiccation. The medial articular branch of the saphenous nerve was isolated and transected in the inguinal region to prevent spinal reflexes. The epineurium was removed and the nerve teased to isolate fine neurofilaments which were then placed on a platinum recording electrode to measure single-unit activity. To identify a joint afferent fibre and its receptive field, the knee joint was gently probed with a blunt glass rod. The mechanical threshold of each recorded joint afferent was determined by gradually increasing the torque applied to the joint until the fibre elicited an action potential. The conduction velocity of the fibres were determined by electrically stimulating the receptive field with a pair of silver bipolar stimulating electrodes (0.6 Hz, 2 ms pulse width, 1-15 V). The mechano-sensitivity of the joint fibre was assessed by applying noxious outward rotations to the knee and counting the number of action potentials elicited during the rotation. Noxious rotation refers to torque occurring outside the normal range but not severe enough to cause soft tissue injury.

2.3.1. Experimental timeline

On day 14 post-MIA induction, 3 sets of noxious rotations, each lasting 5 seconds, were applied 5 minutes apart as a baseline measurement of afferent activity. After close i.a. infusion of CBD (100, 200, or 300 μg in 100 μL) or vehicle (100 μL), joint mechano-sensitivity was assessed for an additional 15 minutes. To minimise the use of animals, multiple doses of CBD or vehicle were assessed in each fibre. A washout period of at least 50 minutes was observed between the administration of varying doses of CBD or vehicle to allow afferent firing to return to baseline levels. The percentage change in afferent activity before and after administration of CBD or vehicle was calculated offline using Spike2 software (Cambridge Electronic Design, Cambridge, United Kingdom). All recorded fibres fired in response to close i.a. administration of potassium chloride (KCl; 1 mM, 0.1 μL) at the conclusion of the experiment, confirming that administered drugs had reached the mechanosensory nerve endings and that the recorded fibre was still viable.

2.4. Behavioural pain measurements

2.4.1. Von Frey hair mechanosensitivity

Von Frey hair mechanosensitivity was used as a measure of secondary allodynia. Alert, unanaesthetised animals were placed in a Plexiglas chamber with a metal mesh flooring which allowed access to the plantar surface of each hind paw. After allowing the animal to acclimatize until exploratory behaviour ceased (approximately 10 minutes), ipsilateral hind paw mechanosensitivity was assessed using a modification of the Dixon up-down method.9 A von Frey hair was applied perpendicular to the plantar surface of the ipsilateral hind paw (avoiding the toe pads) until the hair flexed; the filament was then held in place for 3 seconds. If there was a positive response (ie, withdrawal, shaking, or licking of the hind paw), the next lower strength hair was applied; if there was a lack of response, the next higher strength hair was applied up to a cutoff of 15 g bending force. The 50% withdrawal threshold was
determined using the following formula: $10^{(y_{f} + k_{s})}/10,000$; where $y_{f}$ = value (in log units) of the final von Frey hair used, $k$ = tabular value for the pattern of the last 6 positive and/or negative responses, and $s$ = mean difference (in log units) between stimuli.

### 2.4.2. Hind limb incapacitation

To perform dynamic weight bearing (DWB) measurements, animals were placed in a Perspex chamber (model BIO-DWB-AUTO-R; Bioseb, Boulogne, France) with a pressure-sensitive floor and allowed to move freely. Hind limb weight bearing was tracked and recorded over a 3-minute period. Weight borne on the ipsilateral hind paw was calculated as a percentage of the total weight borne on the hind limbs.

### 2.4.3. Experimental timeline

Animals underwent baseline von Frey hair mechanosensitivity and DWB testing. Separate cohorts were treated on day 14 post-MIA with an L. lactis injection of either vehicle (50 μL) or CBD (100-300 μg/50 μL). In other experimental cohorts, day 14 OA rats were treated with the highest dose of CBD (300 μg/50 μL) and either the CB2 receptor antagonist, AM281 (75 μg/50 μL), the CB2 receptor antagonist, AM630 (75 μg/50 μL), or the TRPV1 receptor antagonist, SB-366791 (30 μg/50 μL) administered locally (subcutaneously; s.c.) over the joint 10 minutes before L. lactis CBD administration. Behavioural pain measurements for these experiments were conducted at 0, 1, 3, 7, 10, 14.

### 2.5. Inflammation measures

Animals were deeply anaesthetised by an intraperitoneal injection of urethane (25% solution; 2g/kg i.p.). A longitudinal incision was made along the ventral skin of the neck to expose the trachea which was cannulated with PE-200 tubing to permit unrestricted breathing. The right carotid artery was also cannulated with PE-30 tubing filled with heparinised saline (1 U/mL) to allow for continuous monitoring of the mean arterial pressure (MAP).

#### 2.5.1. Intravital microscopy

Both hind limbs were immobilised and the capsule of the ipsilateral knee was exposed by surgically removing a small ellipse of the overlying skin and superficial fascia. Physiological buffer (37°C ± 1°C) was immediately and continuously perfused over the exposed joint. Intravital microscopy was used to assess leukocyte-endothelial interactions within the micromicrocirculation of the knee joint, as described previously. The synovial microcirculation was visualised under incident fluorescent light using a Leica DM2500 microscope with a HXC APO L 20X objective and an HC Plan 10X eyepiece giving a final magnification of $\times 200$. In vivo leukocyte staining was achieved by intravenous administration of 0.05% rhodamine 6G (in saline). Straight, unbranched postcapillary venules (15-50 μm in diameter) were chosen for visualisation and 3 fluorescent videos (per time point) were captured for 1 minute each by a Leica DFC 3000 camera (Leica Microsystems Canada Inc, Richmond Hill, ON, Canada). Two measures of leukocyte-endothelial interactions were used to assess articular inflammation: (1) the number of rolling leukocytes to pass an arbitrary line perpendicular to the venule in 1 minute were counted and (2) the number of adherent leukocytes within a 100-μm portion of the venule. Rolling leukocytes were defined as positively stained blood cells travelling slower than the surrounding blood flow, and adherent leukocytes were defined as positively stained cells that remained stationary for a minimum of 30 seconds.

#### 2.5.2. Laser speckle contrast analysis

In the same animals, knee joint blood flow was measured by laser speckle contrast analysis (LASCIA) using a PeriCam PSI System (Perimed Inc, Ardmore, PA). At each time point, 1-minute recordings of the exposed knee joint were taken at a working distance of 10 cm with a frame capture rate of 25 images per second. Using dedicated software (PIMSOFT, Version 15.4.8078), images were averaged to generate 1 perfusion image per second. At the end of the experiment, rats were euthanised and a dead scan of the knee was taken. This "biological zero" value was subtracted from all measurements to account for any Brownian motion in the tissue. Images were analysed offline where mean blood perfusion (perfusion units) in a defined region of interest approximating the knee joint was calculated.

#### 2.5.3. Experimental timeline

Inflammation measures were conducted on day 1 post-MIA induction, which corresponds to the peak of inflammation in this OA model. After baseline intravital microscopy and LASCIA recordings (1 minute) a 50-μL bolus of CBD (300 μg) or vehicle (separate cohort) was applied topically over the exposed knee joint. Subsequent recordings were taken at 5, 15, 30, 60, 120, and 180 minutes after drug administration. In separate cohorts, day 1 MIA rats were treated with the highest dose of CBD (300 μg/50 μL) and either the CB2 receptor antagonist, AM281 (75 μg/50 μL), the CB2 receptor antagonist, AM630 (75 μg/50 μL), or the TRPV1 receptor antagonist, SB-366791 (30 μg/50 μL) administered topically over the joint 10 minutes before CBD administration.

#### 2.6. G-ratio analysis of the saphenous nerve

A segment of the saphenous nerve was isolated proximal to the ipsilateral knee joint and placed in 2.5% glutaraldehyde (diluted with 0.1 M sodium cacodylate buffer), and stored at 4°C for at least 1 week. The nerve samples were then removed from the fixative and rinsed 3 times with 0.1 M sodium cacodylate buffer. The samples were fixed in 1% osmium tetroxide for 2 hours, rinsed with distilled water, and then placed in 0.25% uranyl acetate (4°C) overnight. The samples were then dehydrated in a graduated series of acetone (50%, 70%, 85%, and finally 100%). The samples were then dried in 100% acetone for 10 minutes. Epon-araldite resin was used to mount the samples. The samples were placed in a 3:1 ratio of dried 100% acetone to resin for 3 hours, followed by a 1:3 ratio of dried 100% acetone to resin overnight. Next the samples were placed in 100% Epon-araldite resin for 3 hours and cured in an oven at 60°C for 48 hours. Finally, using an LKB Huxley ultramicrotome with a diamond knife, the samples were sectioned into 100 nm thick slices. Cross-sectional slices of nerves were placed onto a copper wire grid consisting of 300 individual squares per inch (each square measuring 63 ± 58 μm) and then stained with 2% aqueous uranyl acetate for 10 minutes and finally lead citrate for 4 minutes.
Characterisation of the recorded fibres in the electrophysiology experiments.

<table>
<thead>
<tr>
<th>Fibre type</th>
<th>Mechanical Threshold, mNm</th>
<th>Noxious rotation, mNm</th>
<th>Electrical threshold, V</th>
<th>Conduction velocity, m/s</th>
<th>n</th>
</tr>
</thead>
<tbody>
<tr>
<td>II</td>
<td>14 ± 3 (8-28)</td>
<td>23 ± 3 (20-35)</td>
<td>4.3 ± 0.6 (3-6)</td>
<td>2.96 ± 0.3</td>
<td>5</td>
</tr>
<tr>
<td>IV</td>
<td>13 ± 3 (10-15)</td>
<td>25 ± 6 (20-30)</td>
<td>5.2 ± 0.3 (5-6.5)</td>
<td>1.88 ± 0.07</td>
<td>2</td>
</tr>
<tr>
<td>ND</td>
<td>18 ± 3 (2-25)</td>
<td>27 ± 2 (15-35)</td>
<td>ND</td>
<td>ND</td>
<td>9</td>
</tr>
</tbody>
</table>

Fibres are classified as thin-unmyelinated type IV (>2 mNm) or unmyelinated type II (<2 mNm) units. For some fibres, it was not possible to determine conduction velocity and are classified as ND, not determined. Data are mean ± SEM (range).

The copper wire grids containing the saphenous nerve sections were inserted into a JEOL JEM 1230 transmission electron microscope (JEOL Corp Ltd, Tokyo, Japan). The microscope was set at a voltage of 80.0 kV, and images were captured at ×2500 using a Hamamatsu ORCA-HR digital camera (Hamamatsu Photonics, Hamamatsu City, Japan). One nerve cross-section image was visually partitioned into 9 quadrants and 3 images were captured from quadrants 1, 5, and 9. All fibres were assessed using the G-ratio plugin in ImageJ processing software. The G-ratio was calculated using the equation $G = \sqrt{a/A}$, where $a$ is the internal axonal area and $A$ is the total axonal area of the fibre. The higher the G-ratio, the higher the degree of demyelination.

2.7. Drugs and reagents

Cannabidiol (2-[[1R,6R]-3-methyl-6-(1-methylthiethyl)-2-cyclohexen-1-yl]-5-pentyl-1,3-benzenediol) was obtained from Tocris Bioscience (Bio-Technne, Abingdon, United Kingdom), AM281 (CB1 receptor antagonist; 1-(2,4-dichlorophenyl)-5-(4-iodophenyl)-4-methyl-N-(4-morpholinyl)-1H-pyrazolo[3,4-b]pyridine) and AM630 (CB2 receptor antagonist; 6-iodo-2-methyl-1-(2-morpholin-4-ylethyl)[1,3]indol-3-yl)-[4-methoxyphenyl][methanone) were obtained from Cayman Chemicals (Ann Arbor, MI). SB-366791 (N-(3-methoxyphenyl)-4-chlorocinnamidic acid, rhodamine B, cremophor, dimethyl sulfoxide (DMSO), urathane, and MIA were obtained from Sigma-Aldrich (St. Louis, MO). Solutions of CBD, AM281, AM630, and SB-366791 were prepared in vehicle (1:1:18; DMSO:cremophor:saline) on the day of use. Rhodamine B (0.05%) and MIA were dissolved in saline. Physiological buffer (135 mM NaCl, 20 mM NaHCO3, 5 mM KCl, 1 mM MgSO4·7H2O, pH = 7.4) was prepared in the laboratory.

2.8. Statistical analysis

All data were expressed as mean ± SEM. Data were tested for Gaussian distribution by the Kolmogorov–Smirnov test. All data were normally distributed and were therefore analysed using parametric statistics (2-way analysis of variance (ANOVA), 1-way ANOVA, unpaired 2-tailed Student t test, and paired 2-tailed Student t test). A $p$ value less than 0.05 was considered statistically significant.

3. Results

3.1. Effect of acute administration of cannabidiol on joint afferent mechanosensitivity

A total of 17 afferent fibres were recorded in this study. Fibres were characterised based on mechanical and electrical thresholds, and conduction velocity (summarised in Table 1).

On days 14 to 19 post-MIA induction, close i.a. administration of CBD rapidly reduced noxious movement-evoked firing of knee afferent fibres (Fig. 1A) in a dose-dependent manner ($p < 0.0001$; $n = 8$, Fig. 1B). The desensitising effect of 300 μg CBD during noxious joint rotation was significant at 3 minutes after drug application and reached a maximum anti-nociceptive

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**Figure 1.** Dose-dependent effect of CBD on joint afferent firing in established OA. Example of a single-unit recording whereby CBD attenuated firing evoked by noxious rotation (A). Cannabidiol (100, 200, or 300 μg i.a.) decreased afferent firing relative to baseline (B). (*$p < 0.05$ 2-way ANOVA with Bonferroni post hoc test; $n = 8$). The dose-dependent effect of CBD treatment on afferent firing rate was averaged over the 15 minutes after administration (C). (**$p < 0.001$, ****$p < 0.0001$ 1-way ANOVA with Bonferroni post hoc test; $n = 8$). Data are mean values ± SEM. ANOVA, analysis of variance; CBD, cannabidiol; i.a., intra-articular; OA, osteoarthritis.
effect at 7 minutes (29.3% ± 7.4% change compared with baseline). Although all doses of CBD significantly decreased the mean afferent firing over the course of the 15 minutes assessed, the 300 μg dose was the most effective, decreasing firing by 22.9% ± 1.2% overall (P < 0.0001, n = 8, Fig. 1C).

3.2. Effect of acute administration of cannabidiol on sodium monoiodoacetate–induced pain

Intra-articular injection of MIA produced secondary allodynia and weight-bearing deficits in the ipsilateral hind paw and hind limb, respectively, 14 days after injection (P < 0.0001; n = 24; Figs. 2A and B). When compared with vehicle control, low dose CBD (100, 200 μg) had no effect on withdrawal thresholds or hind limb weight-bearing (P > 0.05; n = 8; Figs. 2A and B). The 300 μg dose of CBD, however, significantly increased hind paw withdrawal threshold and hind limb weight bearing over the 3 hours after injection (P < 0.0001; n = 8; Figs. 2A and B). All subsequent experiments used the 300 μg dose of CBD.

To determine whether CBD was acting locally, 300 μg of the drug was injected into the contralateral knee and the withdrawal threshold was assessed in the ipsilateral joint 1 hour later and hind limb weight bearing was assessed in the ipsilateral joint 3 hour later. It was found that the high dose of CBD administered to the contralateral knee had no effect on ipsilateral hind paw withdrawal thresholds indicating that CBD was not acting centrally in this pain test (P < 0.01; n = 8-10; Fig. 3A). However, in the hind limb weight-bearing test, contralateral CBD was not statistically different from the ipsilateral CBD group (P > 0.05; n = 6-22; Fig. 3B).

The cannabinoid receptor antagonists AM281 and AM630 had no effect on CBD-induced analgesia (P > 0.05; n = 6-8; Figs. 4A and B). Conversely, the TRPV1 antagonist, SB-366791, significantly inhibited the analgesic effect of CBD (P < 0.05; n = 6-8; Fig. 4A) with respect to the hind paw withdrawal threshold, but did not have a significant effect on hind limb weight bearing at 3 hours after injection (P > 0.05; n = 6-22; Fig. 4B).

3.3. Effect of acute administration of cannabidiol on sodium monoiodoacetate–induced inflammation

One day after i.t. injection of MIA, rolling leukocytes (P < 0.0001; n = 6-12; Fig. 5A), adherent leukocytes (P < 0.0001; n = 6-12; Fig. 5B), and knee joint perfusion were all significantly increased compared with naïve animals (P > 0.05; n = 6-12; Fig. 5C). After baseline recordings were completed on day 1 post-MIA injection, topical administration of CBD (300 μg) significantly decreased rolling and adherent leukocytes when compared with vehicle over the 3-hour time course (P < 0.0001; n = 6; Figs. 5A and B). Cannabidiol had a moderate inhibitory effect on synovial hyperemia (P < 0.05; n = 6; Fig. 5C). The MAP was unaffected by CBD treatment over the 3-hour time course.
In end-stage OA, intra-articular injection of 300 μg of CBD improved unrestrained hind limb weight bearing and hind paw withdrawal threshold (Fig. 2). These observations, along with our electrophysiology data, assert that CBD acts locally in the joint to reduce joint mechanical pain as revealed by improved weight bearing as well as a reduction in centrally mediated secondary allodynia as determined by hind paw withdrawal threshold. Contralateral injection of CBD had no discernible effect on ipsilateral secondary allodynia confirming that the analgesic effect of intra-articular CBD was localised to the site of administration for this pain test. The anti-nociceptive effect of low dose CBD (100 and 200 μg) observed with electrophysiology was not seen in the behavioural pain assessments. This may be because electrophysiology is a highly sensitive technique that detects subtle response to test agents in the periphery, whereas pain behaviours are more complex and encompass the entire pain pathway. The rationale for using two pain behavioural tests in this study was to interrogate different aspects of the pain pathway. Dynamic incapacitation is a measure of spontaneous pain that is associated with joint degeneration or inflammation arising from peripheral sensitisation. In contrast, von Frey hairs were used to investigate evoked, reflexive response (i.e., paw withdrawal, shake, and lick) at a site distal to the injured joint. This secondary allodynia is a consequence of central sensitisation in late stages of the MIA model, and can be indicative of nerve injury. Thus, it seems that local injection of CBD is effective at reducing direct nociceptive and inflammatory pain in the joint as well as ameliorating neuropathic features of OA pain.

Both CB1 and CB2 receptor antagonists failed to block the CBD-mediated improvements in hind paw withdrawal threshold and weight bearing. Although CBD has been shown to act as an inverse agonist at CB2 receptors and a full agonist at CB1 receptors, it has also been shown to act through GPR55, serotonin receptors (5-HT1A), and various transient receptor potential ion channels. Transient receptor potential vanilloid-1 is known to be involved in MIA-induced peripheral sensitisation, therefore, antagonist experiments were performed to test the involvement of this ion channel in CBD-mediated analgesia. Here, the TRPV1 antagonist SB-366791 attenuated the secondary allodynia imparted by CBD in established OA. This mechanism of action has been previously reported in in vitro studies using human embryonic kidney (HEK 293) cells and using cell membranes from mouse and rat brains. In vivo, TRPV1 antagonism has also been shown to block the pain-relieving effect of CBD in a model of carrageenan-induced paw oedema.
and a moderate increase in joint blood flow. Local application of CBD significantly reduced these acute, inflammatory changes corroborating what has previously been reported in other inflammatory models.\textsuperscript{8,12,20} Oral administration of CBD, for example, has been shown to be anti-inflammatory and anti-hyperalgesic in the carrageenan model of plantar oedema.\textsuperscript{5} Melfait et al., showed that systemic administration of CBD, both intraperitoneally and orally, suppressed disease severity and decreased serum inflammatory cytokine levels in the collagen model of rheumatoid arthritis.\textsuperscript{20} Moreover, CBD administered by a transdermal gel reduced joint swelling, immune cell infiltration, synovial membrane thickening, and the synthesis of pro-inflammatory biomarkers in the Freund complete adjuvant model of inflammatory arthritis.\textsuperscript{12} The data presented here demonstrate for the first time that CBD has the capacity to reduce the inflammatory flares associated with OA.

The inhibitory effect of CBD on leucocyte trafficking was blocked by the TRPV1 antagonist SB-366791. Opening of TRPV1 ion channels causes the peripheral release of inflammatory neuro peptides which promote neurogenic inflammation and enhanced leucocyte trafficking in joints.\textsuperscript{18,41} Thus, the anti-inflammatory effects of CBD observed here could be due to desensitisation of TRPV1 ion channels as has been shown elsewhere.\textsuperscript{14} The anti-rolling effect of CBD on joint leucocytes was also blocked by AM630 suggesting that CB\textsubscript{2} receptors may be involved in opposing leucocyte capture in day 1 MIA joints. Zhao et al. showed that activation of CB\textsubscript{2} receptors can inhibit the expression of P-selectin which is the adhesion molecule responsible for leucocyte rolling.\textsuperscript{43} Whether CBD inhibits joint P-selectin activity by a CB\textsubscript{2} receptor mechanism requires further investigation.

A central hypothesis of this study was that early inhibition of OA-related inflammation with CBD would reduce the development of persistent joint pain. Prophylactic treatment of OA joints with CBD on days 1 to 3 after MIA induction prevented secondary allodynia at day 14, but had no effect on hind limb weight bearing. Inflammation associated with MIA diminishes by day 7,\textsuperscript{4} therefore the pain associated with end-stage OA in this model is largely due to joint degeneration and peripheral neuropathy. Thus, by abolishing early inflammation with prophylactic treatment, CBD attenuates central sensitisation and neuropathic pain development in OA.

Previous studies have shown that MIA-induced OA causes peripheral nerve damage.\textsuperscript{25,38} Demyelination of the ipsilateral saphenous nerve was confirmed by an increase in G-ratio, purporting MIA-induced peripheral neuropathy compared with saline control animals.\textsuperscript{25} This study showed that prophylactic treatment with CBD during the early inflammatory phase of MIA prevented this loss of nerve myelin 14 days later, suggesting that blockade of inflammatory flares during OA could protect against joint nerve damage. The G-ratio data would benefit from future studies examining the expression of a biomarker for peripheral nerve damage to further support this finding.

The findings presented here and elsewhere support the concept that MIA recapitulates the neuropathic aspect of OA pain, which is found in approximately 30% of patients.\textsuperscript{1,34} CBD treatment may be a beneficial therapeutic for the population of patients who experience neuropathic arthritis, and are refractory to currently used first- and second-line analgesics. Several cannabinoid compounds, including CBD, have been shown to be neuroprotective in other musculoskeletal disorders. In a preclinical model of multiple sclerosis, CBD was shown to improve clinical recovery and rotarod scores in animals, correlating with and indicative of a neuroprotective effect.\textsuperscript{43} In addition, CBD and Δ\textsubscript{9}-tetrahydrocannabinol have both been implicated in slowing the progression and promoting the survival of neurones in a preclinical model of
Figure 6. Contribution of cannabinoid and noncannabinoid receptors to the anti-inflammatory effects of CBD. The anti-rolling effect of CBD at 30 minutes was blocked (A) by CB1 receptor antagonist AM630 (75 μg) and TRPV1 antagonist SB366791 (30 μg), but not CB2 receptor antagonist AM281 (75 μg). The anti-adherence effect of CBD in day 1 MIA joints was blocked by SB366791 (B). **P < 0.0001, ***P < 0.001 1-way ANOVA with Fischer LSD post hoc test; n = 60). Data are mean values ± SEM. ANOVA, analysis of variance; CBD, cannabidiol; MIA, sodium monooiodoacetate; VEH, vehicle.

Amyotrophic lateral sclerosis. These studies, in addition to the results presented here, highlight the potential utility of CBD as an analgesic and neuroprotective agent in OA.

Cannabidiol is a non-euphoria producing compound and has a more desirable side effect profile compared with other cannabinoid compounds and commonly prescribed analgesics. Animal studies where CBD was administered systemically showed that the animals had no signs of adverse side effects. For example, exploratory behaviour in rats was not altered by systemic CBD, indicating limited central effects of treatment. Our study shows for the first time that CBD is an effective antinociceptive and anti-inflammatory agent when administered locally around the joint. Successful relief of OA symptoms by peripherally administered CBD suggests a therapeutic option that has a low chance of adverse effects which is more desirable for patients.

Figure 7. Effect of prophylactic CBD administration on the development of pain over 14 days post-MIA injection. Treating MIA knee joints with CBD (300 μg; s.c.; days 0–3) significantly improved von Frey hair withdrawal threshold over the 14-day development of OA when compared with vehicle (A). Pretreatment of MIA knee joints with CBD had no significant effect on hindlimb weight bearing (B). **P < 0.0001, ***P < 0.001, **P < 0.01, *P < 0.05 2-way ANOVA with Bonferroni post hoc test; n = 8). Data are mean values ± SEM. ANOVA, analysis of variance; CBD, cannabidiol; MIA, sodium monooiodoacetate; OA, osteoarthritis; s.c., subcutaneous; VEH, vehicle.
5. Conclusions

This study showed for the first time that local CBD administration inhibited pain and peripheral sensitisation in established OA. Topical treatment with CBD reduced leukocyte trafficking and joint hyperaemia during the early stages of MIA. By attenuating this initial inflammatory response with CBD, end-stage OA pain and peripheral neuropathy were abrogated. Thus, CBD may be a safer therapeutic to treat OA pain locally as well as block the acute inflammatory fases that drive disease progression and joint neuropathy.

Conflict of interest statement

The authors have no conflicts of interest to declare.

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