The Cannabinoid CB1/CB2 Agonist WIN55212.2 Promotes Oligodendrocyte Differentiation In Vitro and Neuroprotection During the Cuprizone-Induced Central Nervous System Demyelination

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Introduction

Understanding how different populations of neurons develop, integrate, and function as a whole has direct impact for the etiology of neurological disorders. Neurons represent a major cell type in the central nervous system (CNS) and are critically associated with glial cells in the formation of effective synapses which provide communication in a timely manner [1]. Oligodendrocytes are a type of glial cells that wrapped axons in the CNS to produce large amounts of myelin sheath [1]. Myelin sheath isolates the axon and promotes the action potentials propagation [2]. Some types of injuries to the CNS drive axonal demyelination and cause severe disorders such as multiple sclerosis, neuropathies, myelopathies, and leukodystrophies. One of the most crucial steps in remyelination is the stimulation of oligodendrocyte precursor cells (OPCs), which progress across several stages from immature to mature oligodendrocytes [1]. The cuprizone (CPZ) mouse model allows the investigation of the complex molecular mechanisms behind non-autoimmune-mediated demyelination. Matsushima and Morell [3] determined that C57BL/6 mice fed with 0.2% CPZ-supplemented diet for 4–6 weeks displayed acute demyelinating lesions followed by spontaneous remyelination.

By activating cannabinoid receptors, Gomez et al. [4] reported a markedly oligodendrocyte differentiation in vitro. The effects of cannabinoids are classically associated with CB1 and CB2 receptors. These receptors are key components of the endocannabinoid system as well as the endogenous cannabinoids ligands, their carriers, and synthetic/degrading enzymes [5]. CB1 receptor is highly expressed in the brain [6] and it is essential for several behaviors including anxiety and fear responses (for review, see [7]). The most important endogenous ligands for cannabinoid receptors are N-arachidonylethanolamine (AEA) [8] and 2-arachidonoylglycerol (2-AG) [9]. N-Arachidonoyl phosphatidylethanolamine phospholipase D (Naplopld) is the central enzyme responsible for the synthesis of AEA, while 2-AG is mainly produced by diacylglycerol lipase (Daqla) in the adult brain as described [10]. After the
activation of cannabinoid receptors, AEA and 2-AG are removed from the synaptic cleft and hydrolyzed by the action of fatty acid amide hydrolase (Faah) and monoacylglycerol lipase (Magl), respectively [11].

With the rationale that CB1 receptor is expressed in oligodendrocytes [4] and marijuana consumption alters CNS myelination [12,13], we directed the present study to evaluate the effects of the cannabinoid CB1/CB2 agonist WIN55212.2 in (1) an in vitro model of oligodendrocyte differentiation and (2) the murine model of demyelination by cuprizone.

Materials and Methods

A total of 90 C57Bl6/J male mice at the age of 7–8 weeks were obtained from Charles River Laboratories (Germany). After arrival, the animals were housed 5 mice per cage and maintained under standard conditions (12-h light/dark cycle with 6:00/18:00 lights on/off, room temperature of 21 ± 2°C, and food and water ad libitum). Procedures performed on mice were in accordance with NIH guidelines for the use of animals in research and the European Communities Council Directive (86/609/EEC).

HOG16 Oligodendrocyte In Vitro Model

The cell line HOG16 was obtained from Eucellbank (University of Barcelona, Spain) in agreement with Dr. G. Dawson (University of Chicago, USA). Oligodendrocytes were cultured in standard medium containing high-glucose Dulbecco’s modified Eagle medium (Invitrogen Ltd., Warrington, UK) complemented with 10% fetal bovine serum (HyClone, Logan, UT, USA), the antibiotics penicillin (50 U/mL) and streptomycin (50 µg/mL) (Invitrogen Ltd., UK) at fully humidified atmosphere incubator at 37°C and 5% CO2. To induce differentiation, cells were exposed to differentiation medium that contained 0.05% FBS, 30 nM triiodothyronine (T3), 30 nM selenium, 0.5 µg/mL insulin (all from Sigma–Aldrich, Munich, Germany), 50 µg/mL transferrin (US Biological, Salem, MA, USA), and the antibiotics used above [14]. The effect of 1 µM WIN55212.2 (WIN) (Sigma–Aldrich) was tested under both conditions according to Bologov et al., [15]. Cells grew with standard medium and when they reached 50% confluency were divided into two groups. One of them remained with standard medium, and in the other one, the medium was replaced by differentiation medium. After 24 h, the cells were subjected to lysis buffer or fixed. Quantitative RT-PCR was performed in triplicate wells and in three independent experiments. Immunohistochemistry was carried out in a Lab-Tek II Chamber Slide System (Nunc, Roskilde, Denmark). Figure 1(A) shows a schematic drawing of the experiment.

Immunocytochemistry on Cell Culture

The cells were washed in phosphate-buffered saline (PBS), fixed in 2% paraformaldehyde (PFA) for 30 min, permeabilized with 0.05% Triton X-100 for 30 min, washed in PBS, and incubated overnight at 4°C in PBS with anti-CB1 receptor antibody (ab23703, dilution 1:300; Abcam, Bristol, UK) and anti-MBP antibody (250746, dilution 1:250; Abbiotec, Aachen, Germany). Next day, after washing the cells with PBS, they were incubated for

30 min with goat anti-rabbit DyLight488-labeled secondary antibody (46402, dilution 1:300; Thermo Scientific, Karlsruhe, Germany), and finally, the excess of secondary antibody was eluted washing with PBS [14]. Cells were analyzed under a fluorescence microscope (Olympus light microscope BX51, Hamburg, Germany) equipped with a camera and filters that allow excitation at 495/10 nm, and labeled structures were digitally photographed using ×100 magnification.

The Cuprizone Murine Model

Mice received regular diet for 0, 3, and 12 weeks (controls) or were fed with 0.2% CPZ for 3 and 6 weeks (treated). In the recovery group, animals were fed with CPZ for 6 weeks, followed by a recovery period of 6 weeks with regular diet. In parallel to CPZ feeding, mice were intraperitoneally injected with WIN or phosphate-buffered saline (vehicle) once per day. WIN was dissolved in 10% DMSO (Sigma–Aldrich), 0.1% Tween 80 (Sigma–Aldrich) in 0.9% saline and prepared freshly every day. The same volume of WIN (drug) or phosphate-buffered saline (vehicle) (200 µL) was applied i.p. The drug was administered at 0.5 or 1 mg/kg. At certain time points (at 3 and 12 weeks), mice were evaluated by
testing the baseline startle response and prepulse inhibition (PPI). After behavioral testing, mice were deeply anesthetized by intraperitoneal injection of 2,2,2-tribromoethanol (Sigma-Aldrich, Germany) and transcardially perfused with cold PBS. From one half of the brain, the corpus callosum was dissected and collected for RNA analysis. The other half of the brain was fixed with 4% PFA and processed later for EPON embedding. We selected this brain region because it is mainly affected in this model [3]. Although this approach is not ideal for the electron microscopy sample preparation, we have chosen this experimental design to be able to perform RNA and morphological analysis on the same brain tissue. Figure 1(B) shows a schematic drawing of the experiment.

Startle response and Prepulse Inhibition (PPI)

An experimental session consisted of a 3-min habituation to 65 dB background white noise (continuously during the session), followed by a baseline recording for 2 min. Startle reflexes were resulted from an acoustic stimuli. The startle reaction to an acoustic stimulus evoked a movement of the platform which was recorded during a frame time of 100 ms (the onset of the acoustic stimulus) and stored for further analysis. For prepulse inhibition tests, the 120-dB startle pulse of 40-ms duration was applied either alone or preceded by a nonstartling prepulse stimulus of 70-, 75-, or 80-dB intensity and 20-ms duration. An interval of 100 ms with background white noise was used between each prepulse and pulse stimulus. Ten trials of each type were displayed in a randomly order with intertrial intervals ranging from 8 to 22 seconds. The amplitude of the startle response (AUs) during the recording frame time was determined as the maximum force displayed during a reaction to 120-dB acoustic stimulus. Maximum amplitudes were averaged individually, separately for all types of trials as described [16].

RNA Isolation

The corpus callosum from each of the CPZ-exposed and controls was sonicated with a blender in RNase-free lysis buffer (Applied Biosystems, Carlsbad, CA, USA). Cell cultures were stopped by removing the medium and adding RNase-free lysis buffer (Applied Biosystems, USA). Samples were kept for 1 h at 4°C. Total RNA was isolated using the TRizol protocol (Invitrogen Ltd., Darmstadt, Germany). The RNA was digested with RNase-free DNase (Qia-gen, Valencia, CA, USA) and checked for integrity by electrophoresis (Bioanalyzer; Agilent Technologies, Santa Clara, CA, USA). N = 5 mice/group.

Electron Microscopy

As described above, mice were deeply anesthetized and transcardially perfused with PBS; half of the brain was dissected and immersion-fixed in 4% PFA and stored at 4°C. Later, the brains were postfixed with 4% formaldehyde (Serva, Heidelberg, Germany), 2.5% glutaraldehyde (Science Services, Munich, Germany), and 0.5% NaCl in phosphate buffer pH 7.4 [17]. Finally, the corpus callosum was postfix with 2% OsO4 (Science Services) in 0.1 M phosphate buffer pH 7.3 and embedded in EPON (Serva) after dehydration with ethanol and propylene oxide. EPON blocks with embedded tissue were then trimmed, using a Leica EM TRIM (Leica, Vienna, Austria), to the size of the corpus callosum. In the following, ultrathin sections were stained with an aqueous solution of 4% uranyl acetate followed by lead citrate [18]. The pictures were taken in an unbiased random fashion with a Zeiss EM900 electron microscope (Zeiss, Oberkochen, Germany) using a side-mounted 2k CCD camera (TRS, Waakirchen, Germany). Counting of myelinated and demyelinated axons was performed using NIH ImageJ software (National Institutes of Health, Bethesda, MD, USA); a total area of \(10^4 \text{ \mu m}^2 = 7840 \text{ \mu m}^2\) was analyzed per mouse and every axon counted. The g ratio was calculated as the ratio between the number of demyelinated axons and the number of myelinated axons. Counting was performed on groups of m = 5 mice.

Focused Gene Signature Profiling

The prognostic 35-gene signature was implemented in the digital transcript counting (nCounter) assay (NanoString) [19]. Total RNA (200–400 ng) was assayed according to the manufacturer’s instructions. Data were standardized by scaling with geometric mean of built-in control gene probes after log transformation (base 2) for each sample. Genes for testing were chosen according to the following criteria: myelin, cannabinoid, corpus callosum connectivity, and psychiatric disorders-related genes. Probe sets for each gene were designed and synthesized by NanoString nCounter technologies (Seattle, WA, USA) (Table S1). N = 5 mice/group.

Statistical Analysis

Statistical significance was evaluated by two-way ANOVA, and a Bonferroni post hoc test was used when appropriate. Significance values were set to \(P < 0.05\). In all figures and text, data are represented as mean ± SEM. Significant effects were identified using Statistica (StatSoft Software, Tulsa, OK, USA).

Results

HOG16 Oligodendrocyte Model

Cells were differentiated from immature oligodendrocytes into a more mature phenotype, that is, cells that were able to synthesize myelin proteins. These cells were grown in standard medium (SM) and differentiation medium (DM) supplemented with 1 μM of WIN, respectively (Figure 1A). Fluorescent immunocytochemistry showed a morphological change associated with differentiation. Cells changed from a round shape (Figure 2A, i-ii, v-vi) to a stellated shape (Figure 2A, iii-iv, vii-viii). The drug also increased the amount of myelin basic protein (MBP) under both SM and DM shown by immunofluorescence staining (Figure 2A, v, vii). The amount of CB1 receptor increased in those cells cultured with standard medium supplemented with WIN (Figure 2A, vi). Mbp and the cannabinoid CB1 receptor (Cnr1) gene expression were further analyzed by quantitative RT-PCR (Figure 2B). In brief, the cannabinoid agonist WIN promoted the expression of Mbp under either SM or DM (\(P < 0.05; P < 0.001\), respectively) (Figure 2B).
Cells cultured in differentiation medium supplemented with WIN increased MBP expression when compared with those cells grown in standard medium combined with WIN (P < 0.01) (Figure 2B). Moreover, CB1 receptor mRNA expression increased under SM with WIN (P < 0.05) as depicted in Figure 2B.

The Cuprizone Murine Model

Startle Response and Prepulse Inhibition (PPI)

Mice exposed to CPZ for 3 weeks displayed no changes in the startle response (Figure 3i). However, we measured an increase in PPI 70 dB in CPZ-fed mice treated with 0.5 mg/kg of WIN when compared with both control and CPZ alone (P < 0.05) (Figure 3ii). In addition, CPZ-fed mice treated with 0.5 mg/kg of WIN displayed higher PPI 75 dB than CPZ alone (P < 0.05) (Figure 3iii). Mice exposed to CPZ alone showed a lower score of PPI 80 dB in contrast to control and CPZ-fed mice subjected to either 0.5 or 1 mg/kg of WIN (P < 0.05) (Figure 3iv). In contrast, no significant differences were found in the recovery group (data not shown).

Methylene Blue/Azure II Staining and Electron Microscopy

Ultrastructural analysis of the corpus callosum revealed a clear demyelination after 3 weeks of CPZ alone (P < 0.001) (Figure 4A, ii; B, ii; C), while CPZ-fed mice subjected to 0.5 mg/kg of WIN showed a lower g ratio than those fed with CPZ and acutely treated with vehicle buffer (P < 0.001) (Figure 4B, iii; C). In contrast, the administration of 1 mg/kg of WIN in CPZ-fed mice induced the maximum demyelination as indicated the g ratio when compared with either controls (P < 0.001), CPZ alone (P < 0.001), or CPZ treated with 0.5 mg/kg of WIN (P < 0.001) (Figure 4B, iv; C). However, no significant differences concerning the g ratio were reported in the recovery group (Figure 4C).

Focused Gene Signature Profiling

The results are summarized in Table 1 according to the gene expression profile data depicted in Figure S1 (3 weeks) and Figure S2 (12 weeks).
Figure 3  Startle response and prepulse inhibition (PPI) during the acute demyelination. We measured an increase in PPI 70 dB in CPZ-fed mice treated with 0.5 mg/kg of WIN when compared with either control or CPZ alone ($P < 0.05$) (ii). In addition, CPZ-fed mice treated with 0.5 mg/kg of WIN displayed higher PPI 75 dB than CPZ alone ($P < 0.05$) (iii). Mice exposed to CPZ alone declined PPI 80 dB in contrast to control and CPZ-fed mice subjected to either 0.5 or 1 mg/kg of WIN ($P < 0.05$) (iv). Data are expressed as mean ± SEM. $P$ values were set as follows: * $P < 0.05$, ** $P < 0.01$, and *** $P < 0.001$. N = 10. CTR, control animals fed with standard diet and treated with phosphate-buffered saline; CPZ, cuprizone-fed animals; WIN, WIN55212.2.

Figure 4  The corpus callosum cytoarchitecture. Ultrastructural analysis of the corpus callosum revealed a clear demyelination after 3 weeks of CPZ alone ($P < 0.001$) (A-ii, B-ii, C), while CPZ-fed mice subjected to 0.5 mg/kg of WIN showed a lower g ratio than those fed with CPZ and acutely treated with vehicle buffer ($P < 0.001$) (B-iii, C). In contrast, the administration of 1 mg/kg of WIN in CPZ-fed mice induced the maximum demyelination as indicated the g ratio when compared with either controls ($P < 0.001$), CPZ alone ($P < 0.001$), or CPZ treated with 0.5 mg/kg of WIN ($P < 0.001$) (B-iv, C). However, no significant differences concerning the g ratio are reported in the recovery group (C). Data are expressed as mean ± SEM. An § indicates significant differences between CPZ-fed groups and their respective control group. CPZ intragroup comparisons between vehicle and drug-treated mice are indicated by an +. Otherwise underlined * indicated comparisons between CPZ-fed treated with 0.5 and those treated with 1 mg/kg. $P$ values were set at $P < 0.001$. N = 5. CTR, control animals fed with standard diet and treated with phosphate-buffered saline; CPZ, cuprizone-fed animals; WIN, WIN55212.2.
The administration of cuprizone increased the expression of apolipoprotein E (ApoE) following cuprizone alone ($P < 0.01$). Moreover, the expression of myelin oligodendrocyte glycoprotein (Mog), myelin basic protein (Mbp), and myelin-associated glycoprotein (Mag), and N-acyl phosphatidylethanolamine phospholipase D (Napepld) declined following cuprizone alone ($P < 0.05$) when compared with their respective controls as depicted in Table 1. The administration of cuprizone increased the expression of retinoid X receptor alpha (Rxra) in mice exposed to vehicle (CPZ alone) as well as in combination with 0.5 mg/kg of WIN ($P < 0.01$) (Table 1). Moreover, the expression of myelin oligodendrocyte glycoprotein (Mog), myelin basic protein (Mbp), myelin-associated glycoprotein (Mag), and N-acyl phosphatidylethanolamine phospholipase D (Napepld) declined following cuprizone alone ($P < 0.05$) or in combination with either 0.5 or 1 mg/kg of WIN ($P < 0.01$) (Table 1). Animals exposed to CPZ diet and treated with either 0.5 or 1 mg/kg of WIN exhibited lower expression of nogo receptor (Ngr1) ($P < 0.001$) than controls (Table 1). Daily injection with 0.5 mg/kg of WIN in CPZ-fed mice decreased the expression of diacylglycerol lipase alpha (Dagla) in contrast to controls ($P < 0.05$) (Table 1). In addition, the administration of 1 mg/kg WIN in CPZ-fed animals decreased specifically the expression of 2',3'-cyclic nucleotide 3' phosphodiesterase (Cnp) and also does calreticulin (Calr) ($P < 0.05$). The recovery group exposed to 1 mg/kg WIN showed lower Olig2, Sox10, Mag, Calr, and discoidin domain receptor 1 (Ddr1) expression in comparison with CPZ alone ($P < 0.001$) (Table 1). Moreover, the expression of myelin oligodendrocyte glycoprotein (Mog), myelin basic protein (Mbp), and myelin-associated glycoprotein (Mag) declined following cuprizone alone ($P < 0.01$) (Table 1). Animals exposed to CPZ diet and treated with either 0.5 or 1 mg/kg of WIN exhibited lower expression of nogo receptor (Ngr1) ($P < 0.001$), special AT-rich sequence-binding protein 2 (Satb2) ($P < 0.05$), and nerve growth factor inducible (Vgf) ($P < 0.05$). The remaining comparisons are depicted in the table. $P$ values were set as follows: NS ($P > 0.05$), *$P < 0.05$, **$P < 0.01$, and ***$P < 0.001$. N = 5. CTR, control animals fed with standard diet and treated with phosphate-buffered saline; CPZ, cuprizone-fed animals.

During the acute demyelination, apolipoprotein E (ApoE) expression increased following CPZ alone ($P < 0.01$) and in combination with either 0.5 or 1 mg/kg of WIN ($P < 0.001$). Moreover, the expression of myelin oligodendrocyte glycoprotein (Mog), myelin basic protein (Mbp), and myelin-associated glycoprotein (Mag) declined following cuprizone alone ($P < 0.05$) or in combination with either 0.5 or 1 mg/kg of WIN ($P < 0.01$). Animals exposed to CPZ diet and treated with either 0.5 or 1 mg/kg of WIN exhibited lower expression of nogo receptor (Ngr1) ($P < 0.001$), special AT-rich sequence-binding protein 2 (Satb2) ($P < 0.05$), and nerve growth factor inducible (Vgf) ($P < 0.05$) than controls (Table 1). Daily injection with 0.5 mg/kg of WIN in CPZ-fed mice decreased the expression of diacylglycerol lipase alpha (Dagla) in contrast to controls ($P < 0.05$) (Table 1). In addition, the administration of 1 mg/kg WIN in CPZ-fed animals reduced the expression of 2',3'-cyclic nucleotide 3' phosphodiesterase (Cnp) and also does calreticulin (Calr) ($P < 0.05$) (Table 1). CPZ-fed mice subjected to 1 mg/kg of WIN displayed higher Ski protooncogene (Ski) expression than those mice treated with 0.5 mg/kg of WIN ($P < 0.01$) (Table 1).

### Table 1  Summary of gene expression analysis

<table>
<thead>
<tr>
<th>Time (weeks)</th>
<th>Comparison</th>
<th>Gene</th>
<th>$P$ value</th>
<th>Gene</th>
<th>$P$ value</th>
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<td>Mbp</td>
<td>$&lt;0.01^{**}$</td>
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<td></td>
<td></td>
<td>Rxra</td>
<td>$&lt;0.05^{*}$</td>
<td>Mog</td>
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<td></td>
<td></td>
<td></td>
<td>Mag</td>
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<td></td>
<td></td>
<td></td>
<td>Napepld</td>
<td>$&lt;0.05^{*}$</td>
</tr>
<tr>
<td></td>
<td>CPZ+0.5 versus CTR</td>
<td>ApoE</td>
<td>$&lt;0.001^{***}$</td>
<td>NogoR</td>
<td>$&lt;0.001^{***}$</td>
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<tr>
<td></td>
<td></td>
<td>Rxra</td>
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<td>Mog</td>
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<td>Mbp</td>
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<td></td>
<td></td>
<td></td>
<td>Mag</td>
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<td></td>
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<td>Vgf</td>
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<td>Rxra</td>
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<td>Calr</td>
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<td>Mag</td>
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<td>$&lt;0.05^{*}$</td>
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*Time = 3: apolipoprotein E (ApoE) expression increased following CPZ alone ($P < 0.01$) and in combination with either 0.5 or 1 mg/kg of WIN ($P < 0.001$). Moreover, the expression of myelin oligodendrocyte glycoprotein (Mog), myelin basic protein (Mbp), and myelin-associated glycoprotein (Mag) declined following cuprizone alone ($P < 0.05$) or in combination with either 0.5 or 1 mg/kg of WIN ($P < 0.01$). Animals exposed to CPZ diet and treated with either 0.5 or 1 mg/kg of WIN exhibited lower expression of Nogo receptor (Ngr1) ($P < 0.001$), special AT-rich sequence-binding protein 2 (Satb2) ($P < 0.01$; $P < 0.05$, respectively), and nerve growth factor inducible (Vgf) ($P < 0.05$). In addition, the administration of 1 mg/kg WIN in CPZ-fed animals decreased specifically the expression of 2',3'-cyclic nucleotide 3' phosphodiesterase (Cnp) and also does calreticulin (Calr) ($P < 0.05$).

*Time = 12: the expression of Mag and Olig2 decreased following CPZ feeding and 0.5 mg/kg of WIN in contrast to CPZ alone ($P < 0.05$). The recovery group exposed to 1 mg/kg WIN showed lower Olig2, Sox10, Mag, Calr, and discoidin domain receptor 1 (Ddr1) expression in comparison with CPZ alone ($P < 0.05$). Furthermore, the expression of Olig2 decreased in CPZ-fed mice treated with 1 mg/kg of WIN when compared with those mice treated with 0.5 mg/kg of WIN ($P < 0.05$). The remaining comparisons are depicted in the table. $P$ values were set as follows: NS ($P > 0.05$), *$P < 0.05$, **$P < 0.01$, and ***$P < 0.001$. N = 5. CTR, control animals fed with standard diet and treated with phosphate-buffered saline; CPZ, cuprizone-fed animals.*
After a recovery period of 6 weeks following cuprizone dietary, dystrobrevin binding protein 1 (Dtnbp1) expression was higher in mice treated with 1 mg/kg of WIN when compared with either CPZ alone or CPZ combined with 0.5 mg/kg of WIN (P < 0.05) (Table 1). Furthermore, Mag and oligodendrocyte transcription factor (Olig2) expression were downregulated following CPZ feeding and 0.5 mg/kg of WIN in contrast to CPZ alone (P < 0.05) (Table 1). The recovery group exposed to 1 mg/kg of WIN showed lower Olig2, Mag, SRY-Box 10 (Sox10), Calr, and discoidin domain receptor 1 (Ddr1) expression in comparison with CPZ alone (P < 0.05) (Table 1). In addition, the expression of Olig2 decreased in CPZ-fed mice treated with 1 mg/kg of WIN when compared with those animals treated with 0.5 mg/kg of WIN (P < 0.05) (Table 1).

### Discussion

After confirming the presence of CB1 receptor in a cultured oligodendrocyte cell line, we also proved that the synthetic cannabinoid agonist WIN exerted a strong effect on immature oligodendrocytes resulting in increased MBP levels 24 h once the differentiation process started, in agreement with Gomez et al. [4]. This fact encouraged us to investigate the effect of WIN in a well-described mouse model of demyelination, the cuprizone model [3]. Here, we focused our study on the acute demyelination and the recovery period.

### Acute Demyelination

CPZ-fed mice exhibited PPI deficits that were reversed by 0.5 mg/kg of WIN. Despite this, Xu et al. [20] reported that mice exposed to CPZ for 3 days exhibited lower prepulse inhibition as well as higher dopamine levels [20]. Overactivation of dopaminergic system in the forebrain correlates with PPI deficits in humans [21] and also in animals [20]. The cannabinoids are actively involved in dopamine release in the prefrontal cortex [22]. Furthermore, ultrastructural analysis of the corpus callosum revealed more myelinated axons following 3 weeks of CPZ diet combined with 0.5 mg/kg, but not in those mice treated with 1 mg/kg. In fact, many studies have demonstrated either therapeutic or deleterious effects of cannabinoids on the viability of various cell types (for review, see [23]) that might be attributable to blockage or stimulation of calcium entry into the cell, which in turn depends on cannabinoid concentration [24].

In neurodegenerative processes, CNS cholesterol synthesis is compromised. ApoE plays an important role in CNS cholesterol and lipid metabolism [25]. Therefore, the dysregulation of the endocannabinoid system has been associated with ApoE activity [26]. Recent reports suggest that demyelination in experimental allergic encephalomyelitis (EAE) is related to an increase in ApoE [25], which supports the present findings. In addition, retinoid acids (rxrs) are essential for cholesterol metabolism [27,28] and stimulate oligodendrocyte differentiation [29]. Interestingly, we found that 0.5 mg/kg of WIN during CPZ exposure increased the expression of Rxra which might protect myelinated axons against cuprizone. In contrast, the drug at 1 mg/kg aggravates CNS demyelination as indicated ultrastructural and gene expression analysis. Particularly, CPZ-fed animals subjected to 1 mg/kg displayed a decrease in the expression of myelin-related genes and also Cnp. Cnpase is involved in RNA trafficking, splicing, and metabolism in the myelinating oligodendrocyte [30–33]. The severity of demyelination in EAE model is modulated by CB1 receptor [34,35], which underpins the present findings. Furthermore, we reported lower Napepld expression in all groups fed with CPZ and this fact might indicate less AEA content. However, previous studies revealed decreased AEA levels [36], whereas others observed normal [37] or higher [38] AEA levels in demyelinated brains. In addition, the gene involved in 2-AG synthesis (Dagla) was declined following CPZ diet and 0.5 mg/kg of WIN although others have found conflicting results [39–42]. This discrepancy might be attributed to the animal model (rats vs. mice), the methods, the sampling time applied, or the target brain structure analyzed.

Administration of CPZ in combination with 0.5 mg/kg of WIN decreased the expression of NgR1 and might confer neuroprotection against CPZ. In fact, NgR1 participates in oligodendrocyte differentiation, myelinization, and cell cytoskeleton reorganization [44], while the blockade of NgR1 signaling prevents axonal degeneration in EAE model [44]. Accordingly, it is noteworthy to speculate that the loss of NgR1 might protect the axon.

Animals fed with CPZ and subjected to either 0.5 or 1 mg/kg of WIN might suffer a dysfunction in the corticocortical connectivity resulting from a downregulation of Satb2 expression. DNA-binding protein Satb2 is essential for the normal elaboration of corticocortical network [45]. During the acute demyelination, the expression of Ski protooncogene increased following CPZ diet and 1 mg/kg of WIN when compared with those animals subjected to 0.5 mg/kg. Such increase might contribute to counteract the exacerbated demyelination observed after CPZ dietary combined with 1 mg/kg of WIN. In line with these findings, it is described that neurons for Ski-deficient mice lose their identity and cannot culminate to form the corpus callosum [46].

The present findings demonstrated lower Vgf expression following the administration of CPZ combined with either 0.5 or 1 mg/kg of WIN, but not after CPZ alone. Vgf is a component of the chromogranin/secretogranin family [47,48] actively involved in amyotrophic lateral sclerosis (ALS) neuroprotection [49]. Because it is described that declined levels of this bioactive peptide (Vgf) could promote neurodegeneration [50], further investigation of these aspects is required.

Proteins associated with endoplasmic reticulum (ER) stress have recently been shown in demyelinating disorders [51,52]. Among these, the chaperone Calr has an important role in serum myelin clearance [53]. Indeed, we assume that CPZ-fed mice treated with 1 mg/kg of WIN showed an impairment in serum myelin clearance as indicated the loss of Calr expression [53].

### The Recovery Period

Animals fed with normal chow for a recovery period of 6 weeks following cuprizone diet displayed no differences regarding sensory motor gaiting, while the corpus callosum was apparently normal [3]. However, certain genes persisted deregulated. Briefly, animals exposed to CPZ and treated with either 0.5 or 1 mg/kg of WIN declined the expression of Mag in comparison with CPZ alone and this fact might indicate a failure in the trophic support for oligodendrocytes [54]. After a recovery period of 6 weeks,
once CPZ was removed from diet, 1 mg/kg of WIN declined the expression of myelin-related genes such as Olig2, Sox10, and Ddr1 in comparison with CPZ alone. The proliferation of OPCs at the demyelinated region is tightly coordinated by a complex interplay of intrinsic, extrinsic, and epigenetic mechanisms [55]. Among the intrinsic factors, Olig2 and Sox10 are critical for oligodendrocytes [56,57]. An overexpression of Ddr1 has been reported in demyelinating oligodendrocytes [58], during oligodendrocyte differentiation as well as during CPZ-induced remyelination [14]. Accordingly, we demonstrated that myelin-related processes are affected by 1 mg/kg WIN during both demyelination and recovery period. Additionally, we found that Olig2 remained downregulated in CPZ-fed animals treated with 1 mg/kg in comparison with those treated with 0.5 mg/kg, which in turn supports that myelination is not fully achieved.

Following CPZ and 1 mg/kg of WIN, Dtnbp1 expression increased in comparison with the remaining CPZ-fed animals, which could promote neurite outgrowth and cytoskeleton rearrangements in order to restore corpus callosum connectivity. In fact, Dtnbp1 regulates neurotransmitter release, signal transduction, and recovery period. Additionally, we found that Olig2 remained downregulated in CPZ-fed animals treated with 1 mg/kg in comparison with those treated with 0.5 mg/kg, which in turn supports that myelination is not fully achieved.

In conclusion, we proved that the drug WIN promotes oligodendrocyte differentiation in vitro. During the CPZ-induced CNS demyelination, we assessed a neuroprotective effect of 0.5 mg/kg of the drug as shown by (1) better sensory motor gating, (2) more myelinated axons/area, and (3) overexpression of Rxra in combination with lower NgR1 expression. In contrast, CPZ-fed mice treated with 1 mg/kg of WIN exhibited the maximum demyelination as indicated by (1) more demyelination axons/area and (2) downregulation of Vgl, Calr, and myelin-related gene expression coupling specifically with lower Cnp expression. However, the detailed mechanisms of action that underlie the cannabinoid neuroprotection warrant further investigation.

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Conflict of Interest

The authors declare no conflict of interest.

References

Supporting Information

The following supplementary material is available for this article:

Table S1. Nanostring nCounter codestex. Column titles from left to right: official gene symbol; target sequence; accession.

Figure S1. Nanostring nCounter gene expression profile at the third week. During the acute demyelination, ApoE expression increased following CPZ alone (P < 0.01) and in combination with either 0.5 or 1 mg/kg of WIN (P < 0.001). The administration of cuprizone increased the expression of Nr2a in mice exposed to vehicle (CPZ alone) as well as in combination with 0.5 mg/kg of WIN (P < 0.01). Moreover, the expression of Mog, Mbp, Mag, and Napepld were declined following CPZ alone (P < 0.05) or in combination with either 0.5 or 1 mg/kg of WIN (P < 0.01). Animals exposed to CPZ diet and treated with either 0.5 or 1 mg/kg of WIN exhibited lower expression of Ngr1 (P < 0.001), Satb2 (P < 0.01; P < 0.05, respectively), and Vgf (P < 0.05). Daily injection with 0.5 mg/kg of WIN in CPZ-fed mice decreased the expression of Dagla in contrast to controls (P < 0.05). In addition, the administration of 1 mg/kg of WIN in CPZ-fed animals decreased specifically the expression of Cnp and also does Calr (P < 0.05). CPZ-fed mice subjected to 1 mg/kg of WIN displayed higher Ski expression than those mice treated with 0.5 mg/kg (P < 0.01). The remaining comparisons were statistically nonsignificant and not included in this figure. Data were normalized by scaling with geometric mean of built-in control gene probes after log transformation (base 2) for each sample. Level of significance (-log10(p)) was set to *1.30103, **2, and ***3. –Log2FC values below 0 indicate downregulation, whereas upper 0 indicate upregulation. N = 5, CTR, control animals fed with standard diet and treated with phosphate-buffered saline; CPZ, cuprizone-fed animals.

Figure S2. Nanostring nCounter gene expression profile at the twelfth week. After a recovery period of 6 weeks following cuprizone diet, Dtnbp1 expression increased in CPZ-fed mice treated with 1 mg/kg of WIN when compared with either CPZ alone or in combination with 0.5 mg/kg of WIN (P < 0.05). Furthermore, Mag and Olig2 expression underwent a downregulation following CPZ feeding and 0.5 mg/kg of WIN in contrast to CPZ alone (P < 0.05). The recovery group exposed to 1 mg/kg of WIN showed lower Olig2, Sox10, Mag, Calr, and Ddr1 expression in comparison with CPZ alone (P < 0.05). In addition, the expression of Olig2 decreased in CPZ-fed mice treated with 1 mg/kg of WIN when compared with those mice treated with 0.5 mg/kg of WIN (P < 0.05). The remaining comparisons were statistically nonsignificant and not included in this figure. Data were normalized by scaling with geometric mean of built-in control gene probes after log transformation (base 2) for each sample. Level of significance (-log10(p)) was set to *1.30103, **2, and ***3. –Log2FC values below 0 indicate downregulation, whereas upper 0 indicate upregulation. N = 5, CTR, control animals fed with standard diet and treated with phosphate-buffered saline; CPZ, cuprizone-fed animals.