



Cannabidiol Acts at 5-HT_{1A} Receptors in the Human Brain: Relevance for Treating Temporal Lobe Epilepsy

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Experimental evidence indicates that cannabidiol (CBD) induces anxiolytic and antiepileptic effects through the activation of 5-HT_{1A} receptors. These receptors are coupled to G_{i/o} proteins and induce inhibitory effects. At present, the interaction of CBD with 5-HT_{1A} receptors in the human brain is unknown. The aim of this study focused on evaluating the interaction between CBD and 5-HT_{1A} receptors in cell membranes obtained from the hippocampus and temporal neocortex of autopsies and patients with drug-resistant mesial temporal lobe epilepsy (DR-MTLE). Cell membranes were isolated from the hippocampus and temporal neocortex of a group of patients with DR-MTLE who were submitted to epilepsy surgery ($n = 11$) and from a group of autopsies ($n = 11$). The [³H]-8-OH-DPAT binding assay was used to determine the pharmacological interaction of CBD with 5-HT_{1A} receptors. The [³⁵S]-GTP γ S assay was used to investigate the CBD-induced activation of G_{i/o} proteins through its action on 5-HT_{1A} receptors. The CBD affinity (pK_i) for 5-HT_{1A} receptors was similar for autopsies and patients with DR-MTLE (hippocampus: 4.29 and 4.47, respectively; temporal neocortex: 4.67 and 4.74, respectively). Concerning the [³⁵S]-GTP γ S assay, no statistically significant changes were observed for both hippocampal and neocortical tissue ($p > 0.05$) at low CBD concentrations (1 pM to 10 μ M). In contrast, at high concentrations (100 μ M), CBD reduced the constitutive activity of G_{i/o} proteins of autopsies and DR-MTLE patients (hippocampus: 39.2% and 39.6%, respectively; temporal neocortex: 35.2% and 24.4%, respectively). These changes were partially reversed in the presence of WAY-100635, an antagonist of 5-HT_{1A} receptors, in the autopsy group (hippocampus, 59.8%, $p < 0.0001$; temporal neocortex, 71.5%, $p < 0.0001$) and the group of patients with DR-MTLE (hippocampus, 53.7%, $p < 0.0001$; temporal neocortex, 68.5%, $p < 0.001$). Our results show that CBD interacts with human 5-HT_{1A} receptors of the hippocampus and temporal neocortex. At low concentrations, the effect of CBD upon G_{i/o} protein activation is limited. However, at high concentrations, CBD acts as an inverse agonist of 5-HT_{1A} receptors. This effect could modify neuronal excitation and epileptic seizures in patients with DR-MTLE.

Keywords: serotonin, 5-HT_{1A} receptor, hippocampus, mesial temporal lobe epilepsy, drug-resistant epilepsy, cannabidiol, temporal neocortex

INTRODUCTION

Cannabidiol (CBD), the main non-psychoactive component of *Cannabis* plants (Russo, 2017), has a terpenophenolic structure hydroxylated at carbons 1 and 3 (Jones et al., 1977). This structure gives CBD lipophilic properties, which allow its passage across the blood–brain barrier (Calapai et al., 2020).

CBD induces antiepileptic effects in humans and experimental models (Silvestro et al., 2019). In patients with Dravet- or Lennox-Gastaut syndromes, the administration of CBD reduces the frequency and severity of the seizures (Maa and Figi, 2014; Thiele et al., 2018; Lazaridis et al., 2019). In patients with drug-resistant temporal lobe epilepsy (DR-TLE), the coadministration of CBD with antiseizure drugs reduces the number and intensity of epileptic seizures (Cunha et al., 1980). CBD administration also decreases seizure activity and neuronal hyperexcitability in experimental models of temporal lobe epilepsy (Khan et al., 2018; Patra et al., 2019). Moreover, CBD produces anxiolytic effects in humans and experimental models (Shannon et al., 2019). CBD induces antidepressant effects that are more evident when tissue serotonin levels are high (Sales et al., 2018). These effects are partially explained because CBD interacts with 5-hydroxytryptamine_{1A} (5-HT_{1A}) receptors (Magen et al., 2010). In cultured Chinese hamster ovary (CHO) cells expressing 5-HT_{1A} receptors, CBD showed micromolar affinity in displacing [³H]-8-OH-DPAT from 5-HT_{1A} receptors, increased [³⁵S]-GTPγS binding in this G_{i/o} protein-coupled receptor, and reduced cAMP concentration. According to these results, the authors concluded that CBD behaves as an agonist of 5-HT_{1A} receptors (Russo et al., 2005). These effects were not reproduced when cell membranes obtained from rat brainstem were exposed to CBD. However, this phytocannabinoid enhanced the ability of 8-OH-DPAT, an agonist of 5-HT_{1A} receptors, to stimulate [³⁵S]-GTPγS binding. These results suggest that CBD shows an allosteric interaction with 5-HT_{1A} receptors (Rock et al., 2012).

At present, the interaction of CBD with 5-HT_{1A} receptors in the human brain is not known. Indeed, changes of this interaction induced by drug-resistant epilepsy may represent a condition that augments or reduces the efficacy of CBD to control the seizure activity. The aim of this study focused on evaluating the interaction of CBD with 5-HT_{1A} receptors in cell membranes obtained from the hippocampus and temporal neocortex of patients with drug-resistant mesial TLE (DR-MTLE). The results were compared with brain tissue with no neurological disorders obtained from autopsies.

MATERIALS AND METHODS

Patients with DR-MTLE

Patients with DR-MTLE underwent an extensive presurgical evaluation that included video electroencephalogram (EEG) and magnetic resonance imaging (MRI) in the Epilepsy Clinic of the General Hospital from Mexico Dr. Eduardo Liceaga. Four serial EEGs were conducted to determine the presence and location of epileptiform activity. T1–T2-weighted MRI served

to identify mesial sclerosis. Patients with cortical dysplasia or neocortical TLE were excluded from the study. After the presurgical evaluation, 11 patients with DR-MTLE (five females and six males) were included in the present study. The scientific and ethics committees from all the institutions involved approved this experimental protocol (authorization number DI/15/403/03/32). Written informed consent was obtained from all participants.

A standard anterior temporal lobectomy ipsilateral to the epileptogenic zone was performed in every patient 48 h after the last seizure occurrence. During the surgical procedure, samples from the hippocampus and temporal neocortex were collected immediately after resection and stored at -70°C (Table 1).

Autopsies

Samples from the hippocampus and temporal neocortex were obtained from 11 autopsies (three females and eight males, 37.1 ± 17.8 years of age). Death was not associated with neurological disorders, for which brain tissues were analyzed and considered as controls. Autopsy samples were collected with a postmortem interval (PMI) of 15.91 ± 3.11 h. The samples were frozen immediately after resection and stored at -70°C . Autopsies were performed at the Institute of Forensic Sciences in Mexico City (Table 2).

Radioligand Displacement Assay

To assess the interaction of CBD with 5-HT_{1A} receptors, we conducted radioligand displacement assays and evaluated the ability of CBD to displace a radioactively labeled ligand bound to these receptors. Cell membranes were obtained as previously described, with some modifications (Benyhe et al., 1997). Briefly, brain tissue (~ 500 mg) was homogenized in ice-cold 50 mM Tris–HCl buffer (pH 7.4). Subsequently, it was centrifuged at 15,000 rpm for 25 min at 4°C . The resulting pellet was resuspended in 50 mM Tris–HCl buffer (pH 7.4) and incubated for 30 min at 35°C . At the end of the incubation, the preparation was centrifuged under the conditions previously indicated. The resultant pellet was resuspended in buffer (50 mM Tris–HCl, 1 mM EGTA, and 5 mM MgCl₂•6H₂O, pH 7.4), and protein concentration was determined by the Bradford method (Bradford, 1976).

The radioligand displacement assay was performed in a final reaction volume of 500 μl that contained 250 μg of protein of the membrane suspension and increasing concentrations of CBD (10 nM to 10 mM). The assay was carried out in the presence of [³H]-8-OH-DPAT at 0.7 nM. This ligand has a high affinity for 5-HT_{1A} ($K_i = 0.56$) and low affinity for other receptors (K_i ranging from 41.9 to $>10,000$; Middlemiss and Fozard, 1983; Schlegel and Peroutka, 1986; Brown et al., 1990; Pauwels et al., 1996; Kleven et al., 1997). According to the K_i of [³H]-8-OH-DPAT for the different receptors, we expected that the results obtained represent the interaction of this ligand on 5-HT_{1A} receptors.

The mixture was incubated for 45 min at 35°C . Non-specific binding was determined in the presence of the 5-HT_{1A} receptor antagonist WAY-100635 (10 μM). The reaction was terminated by rapid filtration on a Brandel M-48 multifilter through a Whatman GF/C glass fiber filter followed by three washes

TABLE 1 | Clinical data of patients with drug-resistant mesial temporal lobe epilepsy (DR-MTLE).

ID	Gender	Age (years)	Age of seizure onset (years)	Duration of epilepsy (years)	Frequency of seizures (per month)	Side of focus	Precipitating factors	ASD before surgery
HUM-138	F	37	4	33	75	Left	Febrile seizures in childhood	PHT/CBZ/VA/OXC/CLN
HUM-154	F	9	3	6	7	Right	Febrile seizures in childhood	VA/TOP/CBZ/PHB/LEV/LAM
HUM-155	M	31	12	19	20	Left	Febrile seizures in childhood	PHT/VA/CBZ/LAM
HUM-161	M	19	0.16	19	2	Left	Febrile seizures in childhood	PHT/VA/CBZ/TOP
HUM-162	M	56	18	38	30	Right	Temporal-occipital AVM	CBZ/DZP/PHT/PHB/LEV/VA
HUM-168	M	28	15	13	15	Bilateral	TBI and mother with epilepsy	CBZ/CLN/LEV
HUM-169	M	46	12	34	15	Right	TBI	PHT/CBZ
HUM-173	F	46	8	38	2	Right	Unknown	PHT/OXC
HUM-184	F	28	13	15	16	Bilateral	Unknown	LAM/VA/CBZ/LEV
HUM-192	M	41	32	9	10	Left	Father with epilepsy	LAM/VA/CBZ/LEV
HUM-193	F	22	11	11	4	Right	Febrile seizures in childhood	LAM/VA/LEV/TOP

ASD, antiseizure drugs; AVM, arteriovenous malformation; CBZ, carbamazepine; CLN, clonazepam; F, female; LAM, lamotrigine; LEV, levetiracetam; M, male; OXC, oxcarbazepine; PHB, phenobarbital; PHT, phenytoin; TBI, traumatic brain injury; TOP, topiramate; VA, valproic acid.

with ice-cold buffer (50 mM Tris-HCl, pH 7.4). Radioligand binding was determined as disintegrations per minute (DPM) values (Beckman LS6000SC scintillation counter), which were normalized with respect to the maximum binding. Data were fitted to a non-linear regression to determine the inhibitory concentration 50 (IC₅₀) with the model $Y = \text{Bottom} + (\text{Top} - \text{Bottom}) / (1 + 10^{(\text{LogIC}_{50} - X) * \text{HillSlope}})$ using Prism software (GraphPad Software, Inc.). The same equation was used to determine the Hill coefficient, which gives information on the number of interacting sites and possible allosteric interactions (Prinz, 2010). The Cheng-Prusoff equation was used to determine the inhibition constant (K_i) considering the dissociation constant (K_d) of [³H]-8-OH-DPAT equal to 0.46 nM (Cheng and Prusoff, 1973; Cusack et al., 1994). Data are expressed as the mean ± standard error of the mean (SEM).

[³⁵S]-GTPγS Binding Assay

5-HT_{1A} receptors are highly expressed G_{i/o} protein-coupled receptors that activate inhibitory pathways. Considering that

CBD acts on several G_{i/o} protein-coupled receptors, such as CB₁, CB₂, opioid, and 5-HT_{1A}, among others (Alves et al., 2020), we initially evaluated if CBD was able to activate the G_{i/o} protein and if a specific antagonist to 5-HT_{1A} receptors was able to prevent such effect. The [³⁵S]-GTPγS binding assay was used for this purpose.

Brain tissue was homogenized in buffer (50 mM Tris-HCl, 1 mM EGTA, and 3 mM MgCl₂•6H₂O, pH 7.4). The homogenate was centrifuged at 20,000 rpm for 45 min at 4°C. The resulting pellet was resuspended in buffer (50 mM Tris-HCl, 0.2 mM EGTA, 9 mM MgCl₂•6H₂O, and 150 mM NaCl, pH 7.4) and centrifuged under the same conditions. The resulting pellet was resuspended in buffer once more, and protein concentration was determined as described above (see “Radioligand Displacement Assay” section).

The [³⁵S]-GTPγS binding assay was carried out as previously described (Spetea et al., 1998; Cuellar-Herrera et al., 2014), with minor variations. Briefly, cell membranes (10 μg of protein) were incubated at 30°C for 60 min in a reaction tube that contained Tris-EGTA buffer [Tris-50 mM HCl, 1 mM EGTA, 3 mM MgCl₂•6H₂O, 100 mM NaCl, and 0.1% (w/v) albumin; pH 7.4], GDP (100 μM), [³⁵S]-GTPγS (100 μM), and increasing concentrations of CBD (1 pM to 100 μM). Total binding was measured in the absence of CBD. Non-specific binding was estimated in the presence of unlabeled GTPγS (100 μM). Data were analyzed as specific binding that resulted from subtracting non-specific binding from total binding. If any effect was obtained, WAY-100635 (100 μM) was included in the assay to determine the participation of 5-HT_{1A} receptors. The reaction was terminated by rapid filtration on a Brandel M-48 multifilter through a Whatman GF/B glass fiber filters followed by three washes with ice-cold buffer (50 mM Tris-HCl and 5 mM MgCl₂•6H₂O, pH 7.4). A concentration-effect curve was built

TABLE 2 | Clinical data of autopsies.

ID	Gender	Age (years)	Cause of death	PMI (h)
A2	M	29	Polycontusion	18
A3	M	30	Ballistic trauma	14
A7	M	45	Suffocation	18
A8	M	73	Complications associated with diabetes	15
A10	M	36	Ballistic trauma	12
A11	F	12	Suffocation	14
A12	F	40	Ballistic trauma	20
A14	F	45	Unknown	10
A16	M	57	Myocardial infarction	18
A17	M	25	Penetrating wound in thorax	18
A21	M	16	Suffocation	18

F, female; h, hours; M, male; PMI, postmortem interval.

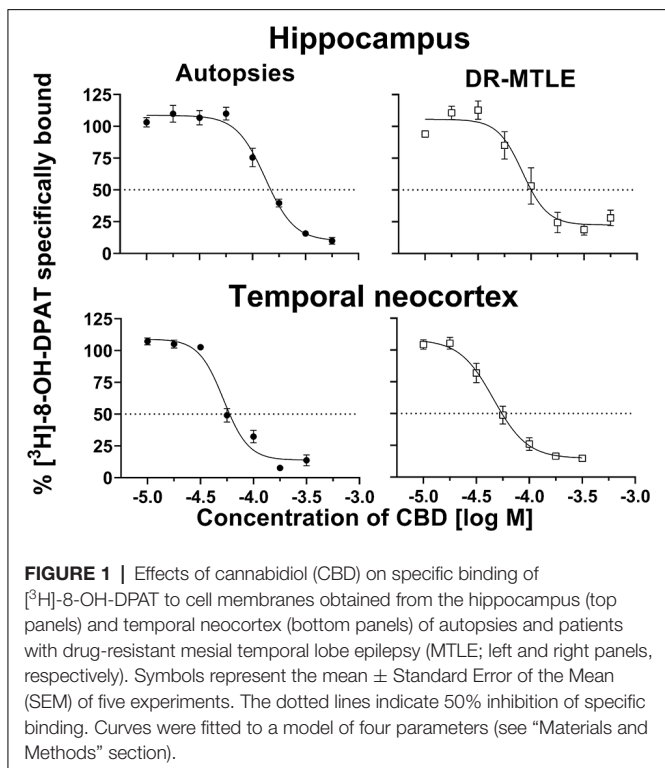
with the percentage of activation calculated considering the basal binding (in the absence of stimulation) as zero. Data were analyzed by a two-way ANOVA, taking group and CBD concentration as factors (concentration–effect curve), or a one-way ANOVA (effect of 100 μ M in presence or absence of WAY-100635). p values < 0.05 were considered statistically significant.

RESULTS

CBD Acts on 5-HT_{1A} Receptors in Human Brain Tissue

In the hippocampal tissue obtained from autopsies, CBD displaced [³H]-8-OH-DPAT from its binding sites in a concentration-dependent manner ($IC_{50} = 129.40 \pm 9.40 \mu$ M; $pK_i = 4.29 \pm 0.03$). In this tissue, the Hill coefficient was 3.37 ± 0.27 . The radioligand displacement assay on the hippocampal tissue of patients with DR-MTLE revealed similar values ($IC_{50} = 93.61 \pm 18.25 \mu$ M, $p = 0.1194$; $pK_i = 4.47 \pm 0.09$, $p = 0.0748$) as those observed in the autopsies. However, in the group of DR-MTLE patients, the Hill coefficient was higher (4.84 ± 0.35 , $p < 0.05$) than in the group of autopsies.

In the temporal neocortex, we observed similar values in both DR-MTLE and autopsy groups (Figure 1). In the autopsy group, CBD displacement values ($IC_{50} = 54.34 \pm 3.66 \mu$ M; $pK_i = 4.67 \pm 0.03$; Hill coefficient = 4.67 ± 0.93) were found within the same range of concentration as in the group of patients with DR-MTLE ($IC_{50} = 46.84 \pm 6.30 \mu$ M, $p = 0.334$; $pK_i = 4.74 \pm 0.05$, $p = 0.2643$; Hill coefficient = 3.18 ± 0.57 , $p = 0.2091$; Table 3).



CBD Modifies the Activity of G_{i/o} Protein-Coupled Receptors

In the autopsies, CBD did not produce significant changes in the binding of [³⁵S]-GTP γ S at low concentrations (1 pM to 10 μ M, $p > 0.05$), neither in the hippocampus nor in the temporal neocortex. However, at high concentrations (100 μ M), CBD reduced [³⁵S]-GTP γ S binding, which was 39.2% and 35.2% lower in the hippocampus and temporal neocortex, respectively, than the basal levels ($p < 0.05$ and $p < 0.01$, respectively; Figure 2).

When evaluating the brain tissue of patients with DR-MTLE, the [³⁵S]-GTP γ S assay revealed similar results as those observed in the group of autopsies. No changes were detected at low concentrations (1 pM to 10 μ M, $p > 0.05$), whereas 100 μ M CBD induced a significant reduction of [³⁵S]-GTP γ S binding in the hippocampus and temporal neocortex (39.6% and 24.4% lower, respectively, when compared to basal binding levels, $p < 0.05$; Figure 2). In comparison with the autopsy group, a less evident [³⁵S]-GTP γ S binding decrease induced by CBD was observed in the temporal neocortex of the DR-MTLE group ($p < 0.05$).

The decrease in the constitutive activity of G_{i/o} proteins induced by a high concentration of CBD was partially reversed in the presence of WAY-100635 at 100 μ M in both the autopsy group (hippocampus, 59.8%; $p < 0.0001$; temporal neocortex, 71.5%; $p < 0.0001$) and the DR-MTLE group (hippocampus, 53.7%; $p < 0.0001$; temporal neocortex, 68.5%; $p < 0.001$; Figure 3).

DISCUSSION

The radioligand displacement assay results support CBD affinity for 5-HT_{1A} receptors in the brain tissue obtained from autopsies and patients with DR-MTLE. Furthermore, the experiments using [³⁵S]-GTP γ S revealed that CBD decreased the constitutive activity of G_{i/o} protein-coupled receptors at high concentrations and an antagonist of 5-HT_{1A} receptors significantly reversed this effect.

Our experiments showed that CBD displaces [³H]-8-OH-DPAT from its binding site on 5-HT_{1A} receptors in a concentration-dependent manner in human hippocampal and neocortical samples. These findings support the affinity of CBD for 5-HT_{1A} receptors in the human brain. Indeed, they are in agreement with previous studies indicating that the exposure to CBD at micromolar concentrations displaced [³H]-8-OH-DPAT from its binding site in CHO cells expressing the human 5-HT_{1A} receptor (Russo et al., 2005).

The present results indicate that CBD interacts with 5-HT_{1A} receptors at high concentrations regardless of its lower affinity ($pK_i \approx 4.5$) in comparison with other ligands such as serotonin ($pK_i = 9.2$), 8-OH-DPAT ($pK_i = 8.0$), or WAY-100635 ($pK_i = 8.7$; U.S. National Library of Medicine). However, the Hill coefficients obtained suggest that CBD is an allosteric modulator of 5-HT_{1A} receptors (Hill, 1910; Goutelle et al., 2008). This condition may facilitate the binding of endogenous ligands and agonists to 5-HT_{1A} receptors (May et al., 2007; Saleh et al., 2018). Indeed, the allosteric condition of CBD may represent a novel

TABLE 3 | Data obtained from the radioligand displacement assay in the hippocampus and temporal neocortex of autopsies and patients with drug resistance mesial lobe epilepsy.

Brain structure	Group	IC ₅₀ (μM)	K _i (μM)	pK _i	Hill coefficient
Hippocampus	Autopsies	129.40 ± 9.40	51.31 ± 3.73	4.29 ± 0.03	3.37 ± 0.27
	Patients	93.61 ± 18.25	37.12 ± 7.24	4.47 ± 0.09	4.84 ± 0.35*
Temporal neocortex	Autopsies	54.34 ± 3.66	21.55 ± 1.45	4.67 ± 0.03	4.67 ± 0.93
	Patients	46.84 ± 6.30	18.57 ± 2.50	4.74 ± 0.05	3.18 ± 0.57

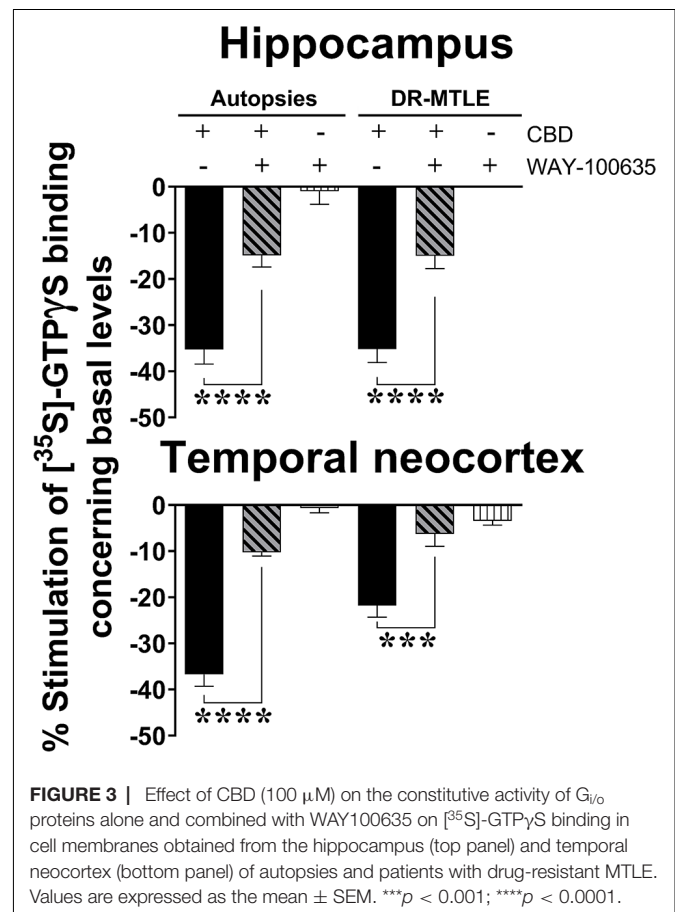
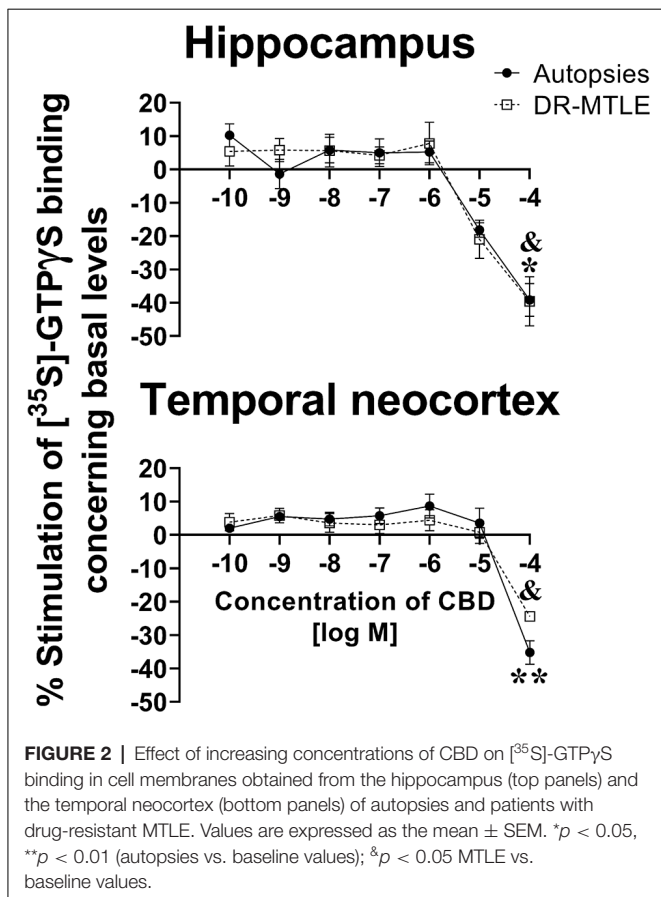
IC₅₀, inhibitory concentration 50; K_i, inhibition constant; pK_i, inverse logarithm of K_i. K_i was determined from IC₅₀ according to the Cheng-Prusoff equation (1973): $K_i = IC_{50}/(1 + [radioligand]/K_D)$. Values are expressed as the mean ± SEM of five experiments. **p* < 0.05 vs. autopsies.

therapeutic strategy to influence the effects of 5-HT_{1A} receptors (Azam et al., 2020).

Radioligand binding assay revealed differences in IC₅₀ of CBD between hippocampus and temporal neocortex. It also showed the inversion of the Hill coefficient profile between autopsies and patients when temporal neocortex and hippocampus were compared. The results obtained suggest that activation of 5-HT_{1A} receptors by CBD induces different functional consequences within these brain regions and depending on the pathological condition. It is known that variations of Hill coefficients are associated with conformational changes of the receptor and the number of binding sites (Colquhoun, 1998; Prinz, 2010). We found that the group of DR-MTLE patients showed a higher Hill coefficient in the hippocampus, the brain structure that mainly develops aberrant changes due to epilepsy

(Bartolomei et al., 2008). This result indicates that epilepsy might be producing changes in the binding sites for CBD on the 5-HT_{1A} receptors in the hippocampus (Lolkema and Slotboom, 2015). This finding is consistent with a previous study in which the Hill coefficient increase was associated with neuronal hyperexcitability and seizure activity due to a conformational change in the γ2 subunit of the GABA_A receptor (Migita et al., 2013). Further binding kinetics studies are necessary to determine changes mediating the enhanced Hill coefficient in 5-HT_{1A} receptors in the hippocampus of patients with DR-MTLE.

Regarding the [³⁵S]-GTPγS assay, low concentrations of CBD did not modify the activity of G_{i/o} proteins in cell membranes obtained from the human brain, neither in autopsies nor in patients with DR-MTLE. These results



are similar to those previously reported by Rock et al. (2012), who described that G_{i/o} protein-coupled receptors in rat brainstem membranes were not activated at low concentrations of CBD. This finding could be explained because the receptor–transducer selectivity was not evaluated. This is an important limitation of the experimental procedure used in the present study since the [³⁵S]-GTPγS assay evaluates all the interactions of G_{i/o} protein-coupled receptors. Another possibility is that, at low concentrations of CBD, an equilibrium state between the activation and inhibition of G_{i/o} proteins is achieved, which is a common condition for the G_{i/o} protein-coupled receptors (Seyedabadi et al., 2019).

The constitutive activity of 5-HT_{1A} receptors is susceptible to the effect of inverse agonists (Newman-Tancredi et al., 1997; Milligan, 2003). We found that high concentrations of CBD (100 μM) decreased the binding of [³⁵S]-GTPγS to G_{i/o} proteins below the baseline values. Therefore, CBD reduced the constitutive activity of receptors coupled to G_{i/o} proteins. These changes were partially reversed when cell membranes were exposed to CBD in the presence of WAY100635, an antagonist of 5-HT_{1A} receptors. The results obtained indicated that at high concentrations, CBD modifies the constitutive activity of 5-HT_{1A} receptors acting as an inverse agonist.

Inverse agonists could indirectly increase the signalling of targeted receptors through the increase in the proportion of receptors on the cellular surface (Abbas et al., 2007; Kumar et al., 2019). Therefore, the continuous administration of inverse agonists of 5-HT_{1A} receptors induces antiallosteric effects due to inverse tolerance (Deseure et al., 2003). According to this information, the continuous administration of CBD as an inverse agonist of 5-HT_{1A} receptors may represent a therapeutic strategy to augment the signaling of these and other G_{i/o} protein-coupled receptors involved in neuroprotection. Additional experiments are essential to support this notion.

It is known that CBD is an inverse agonist of other G_{i/o} protein-coupled receptors, such as CB2 receptors (Thomas et al., 2007) and GPR3 and GPR6 orphan receptors (Laun and Song, 2017). In the present study, WAY-100635 partially blocked the CBD-induced decreased binding of [³⁵S]-GTPγS to G_{i/o} proteins. WAY-100635 is considered the quintessential antagonist of the 5-HT_{1A} receptors ($K_i = 2.2$ nM). However, although with lower affinity values, it is also an antagonist of other receptors such as 5-HT_{2A} ($K_i = 6260$ nM), 5-HT_{2B} ($K_i = 24$ nM), and D2-like ($K_i = 16.4$ – 940 nM; Chemel et al., 2006). Therefore, WAY-100635 could be blocking the constitutive activity of 5-HT_{1A}, 5-HT_{2A}, 5-HT_{2B}, and D2-like receptors as well. On the other hand, [³H]-8-OH-DPAT has low affinity for 5-HT₇ (Thomas et al., 1998) and α1 receptors (Yoshio et al., 2001). Then, some of the results obtained can involve the action of CBD on these receptors. Future experiments should be conducted to investigate the effects of CBD on binding and constitutive activity of different type of receptors in the brain of patients with epilepsy. These experiments should include kinetic binding studies and displacement experiments (pseudo-competition experiments) in the presence and absence of CBD.

5-HT_{1A} receptors play an important role in cerebral functions, and they are considered targets to develop novel therapeutic strategies. They show heterogeneous distribution, including pre- and postsynaptic localization, as well as cross-talk with different types of 5-HT and other neurotransmitter receptors. Dysfunction of 5-HT_{1A} receptors is associated with psychiatric disorders such as anxiety and depression (Popova and Naumenko, 2013). Experimental evidence indicates decline of 5-HT_{1A} receptor binding in the brain of patients with epilepsy (Toczek et al., 2003; Theodore et al., 2012). It is suggested that 5-HT_{1A} receptors on astrocytes represent a potential therapeutic target for the treatment of neurodegenerative disorders (Miyazaki and Asanuma, 2016). An important limitation of the present study is that the current methodology does not allow one to identify in which type of cells the 5-HT_{1A} receptors were evaluated. More experiments are needed to investigate the effects of CBD on specific brain cells and its therapeutic relevance.

DATA AVAILABILITY STATEMENT

The raw data supporting the conclusions of this article will be made available by the authors, without undue reservation.

ETHICS STATEMENT

The studies involving human participants were reviewed and approved by Ethics committee of the Hospital General de Mexico and Centro de Investigacion y de Estudios Avanzados. The patients/participants provided their written informed consent to participate in this study.

AUTHOR CONTRIBUTIONS

LR conceived and designed the study and wrote the manuscript. CM-A carried out the experiments and analyzed the results. FC-C carried out the experiments. FV performed the neurosurgery of patients. AV and GA-C identified and evaluated the patients with epilepsy. MC-H collected samples, participated in the analysis of the results, and wrote the manuscript. All authors contributed to manuscript revision, read, and approved the submitted version.

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Conflict of Interest: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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