Research Paper: Role of Nitric Oxide in the Antipruritic Effect of WIN 55,212-2, a Cannabinoid Agonist

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ABSTRACT

Introduction: For centuries, cannabinoids are known to be effective in pain relief. Itch is an unpleasant sensation that provokes a desire to scratch. Since itch and pain are two sensations sharing a lot in common, we aimed to investigate whether the cannabinoid agonist WIN 55,212-2 reduces serotonin-induced scratching behavior and also observe whether modulation of Nitric Oxide (NO) production mediates the antipruritic effect of WIN 55,212-2.

Methods: Scratching behavior is induced by intradermal injection of serotonin (50 µg/50 µL/mouse) to BALB/c mice. The cannabinoid agonist WIN 55,212-2 (1, 3, 10 mg/kg, IP) was given 30 min before serotonin injection. To observe the effect of NO modulation on the antipruritic effect of cannabinoids, the endothelial nitric oxide synthase (NOS) inhibitor L-NAME (3 mg/kg, IP), the neuronal NOS inhibitor 7-nitroindazole (3 mg/kg, IP), and the NO precursor L-arginine (100 mg/kg, IP) were administered together with WIN 55,212-2.

Results: WIN 55,212-2 reduced serotonin-induced scratches at higher doses (3, 10 mg/kg; P<0.0001). The endothelial NOS inhibitor L-NAME, the neuronal NOS inhibitor 7-nitroindazole, and the NO precursor L-arginine did not influence the antipruritic action of WIN 55,212-2. When NO modulators were used alone, only the neuronal NOS inhibitor 7-nitroindazole attenuated serotonin-induced scratches (P<0.0001).

Conclusion: Our findings indicate that exogenous cannabinoids may attenuate serotonin-induced scratches and NO does not mediate the antipruritic effect of WIN 55,212-2. On the other hand, neuronal NOS inhibition may play a role in the production of serotonin-induced scratches.
1. Introduction

Cannabinoids are chemicals that produce their effects mostly via activating cannabinoid receptors (CB1, CB2); they include phytocannabinoids, synthetic cannabinoids, and endocannabinoids (Maccarrone et al., 2015; Olah, Szekanecz, & Biro, 2017; Ulugol, 2014). The analgesic activity of cannabinoids has been known for centuries; however, these drugs could not be used efficiently in the clinics due to their potential drug abuse and unwanted central side effects, including the development of drug tolerance and addiction. Following many successful clinical trials, cannabinoids have been approved for indications such as neuropathic pain, multiple sclerosis, and so on first in the USA and Canada, and then in many European countries in recent years (Grotenhermen & Muller-Vahl, 2012; Lucas, 2012; Ulugol, 2014). The number of approved indications and countries cannabinoids are used therapeutically is expected to increase in the years ahead.

Pruritus (itch), a common unpleasant symptom, is seen not only in skin diseases but also in systemic disorders. Pain and itch share a lot in common in terms of pathophysiology, which suggests why pain mechanisms should also be evaluated when itch is being investigated (Ross, 2011; Schmelz, 2010). Recently, similar to pain, not only peripheral mechanisms but also central mechanisms have been suggested to play important roles in itch. These new notions have started to change the approaches to the therapy of pruritus (Cevikbas, Steinhoff, & Ikoma, 2011). Understanding the detailed mechanisms of itch will contribute to the development of more efficient antipruritic medications with fewer side effects.

The number of studies regarding the analgesic effects of cannabinoids is quite remarkable. Nevertheless, the effects of cannabinoids on itch have not been investigated sufficiently despite the similarities between pain and itch mechanisms. Cannabinoid receptor agonists have been shown to reduce itch behavior, whereas cannabinoid receptor antagonists like rimonabant increased itch behavior dose-dependently in mice (Darmani & Pandya, 2000). Other research studies also pointed out that the brain penetrating CB1 inverse agonist rimonabant creates itch sensation, and this effect is reduced by cannabinoid receptor agonists (Janoyan, Crim, & Darmani, 2002; Schlosburg, O’Neal, Conrad, & Lichtman, 2011). Furthermore, it is determined that the peripheral application of the synthetic cannabinoid HU210 suppresses histamine-induced responses in human skin (Dvorak, Watkinson, McGlone, & Rukwied, 2003). In addition to these studies, few studies have carried out on the role of the endocannabinoid system in itch behavior (Schlosburg, Boger, Cravatt, & Lichtman, 2009; Spradley, Davoodi, Gee, Carstens, & Carstens, 2012; Tosun, Gunduz, & Ulugol, 2015).

Nitric Oxide (NO) is produced from L-arginine by Nitric Oxide Synthase (NOS) and regarded as a neuronal messenger and a modulator in the central nervous system (Snyder & Breit, 1991). NOS has at least three isoforms: endothelial

### Highlights

- Exogenous cannabinoids reduce serotonin-induced scratching behavior.
- Nitric oxide does not mediate the antipruritic action of cannabinoids.
- Cannabinoids have the potential to be used as antipruritic drugs.

### Plain Language Summary

Cannabinoid drugs are not used effectively due to their potential drug abuse and side effects. For centuries, cannabinoids are known to exert analgesic actions, but they also produce antipruritic effects. There are numerous studies on the mechanisms of the analgesic effects of cannabinoids; however, there are only a few research on their antipruritic mechanism of action. In this study, we observed the modulatory role of nitric oxide in the effect of cannabinoids on serotonin-induced scratches which nitric oxide did not play role in this action. We induced scratching behavior by administering serotonin intradermally. Then we injected the cannabinoid agonist WIN 55,212-2 and observed the reduction of the scratching behavior. Afterwards, we administered an endothelial nitric oxide synthase inhibitor, a neuronal nitric oxide synthase inhibitor, and a nitric oxide precursor and showed that nitric oxide does not mediate the antipruritic effects of WIN 55,212-2. Whatever the mechanism of action, cannabinoids have the potential to be used as antipruritic drugs, especially if their side effects are reduced.

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similar to our earlier studies was re-played to calculate the number of bouts of scratching, the manner was considered as one bout of scratching. The video taped for 30 min under quiet circumstances. In general, serotonin injection, the animals were put into a transparent shaved rostral part of the back of the mice. Immediately after min before WIN 55,212-2, respectively.

In groups 5-7, to evaluate the effect of NO modulation on the antipruritic effect of cannabinoids, the non-selective NOS inhibitor L-NAME (3 mg/kg, IP), the nNOS inhibitor 7-nitroindazole (3 mg/kg, IP), and the NO precursor L-arginine (100 mg/kg, IP) were administered 10 min before WIN 55,212-2, respectively.

In groups 8-10, NOS inhibitors L-NAME (3 mg/kg, IP) and 7-nitroindazole (3 mg/kg, IP) and the NO precursor L-arginine (100 mg/kg, IP) were given, respectively. Thus, we attempted to confirm the cannabinoid action on serotonin-induced scratching behavior and investigate the mediatory effect of NO in this antipruritic effect.

2.5. Drugs

Serotonin hydrochloride was dissolved in 0.9% saline, while WIN 55,212-2, L-NAME, 7-nitroindazole, and L-arginine were dissolved in 20% DMSO, 1% Tween 80, 1% ethanol and 78% saline. This composition is also used in the vehicle group. All of the drugs were purchased from Sigma-Aldrich. Drug doses and treatment times were determined from our previous research studies (Gunduz, Topuz, Karadag, & Ulugol, 2016; Ulugol et al., 2002; Ulugol, Topuz, Gunduz, Kizilay, & Karadag, 2016).

2.6. Statistical analysis

Bartlett’s test was used to test Gaussian distribution. To assess the significance of any difference, one-way ANOVA test followed by the Bonferroni post hoc test was performed. Results were expressed as Mean±Standard Error of the Mean. A P<0.05 was considered significant. GraphPad Prism 6.0 was used for statistical analysis and to plot the graphs.
3. Results

3.1. Effect of WIN 55,212-2 on serotonin-induced scratching behavior

Although WIN 55,212-2 reduced the number of scratches at its lowest dose (1 mg/kg), this effect was not statistically significant. WIN 55,212-2 elicited significant antipruritic activity at higher doses (3, 10 mg/kg, P<0.0001, Figure 1).

3.2. Effect of NO modulators on the antipruritic activity of WIN 55,212-2

NOS inhibitors L-NAME (3 mg/kg, IP) and 7-nitroindazole (3 mg/kg, IP) and the NO precursor L-arginine (100 mg/kg, IP) did not change the antipruritic activity of WIN 55,212-2 (Figure 2). When NO modulators are used alone, only nNOS inhibitor 7-nitroindazole (3 mg/kg, IP) significantly reduced the number of scratches produced by serotonin (P<0.0001, Figure 2).

3.3. Effects of WIN 55,212-2 and NO modulators on locomotor performance

WIN 55,212-2 disrupted motor function at its highest dose (10 mg/kg; P <0.0001, compared with the control, Figure 3). NOS inhibitors L-NAME (3 mg/kg, IP) and 7-nitroindazole (3 mg/kg, IP) and the NO precursor L-arginine (100 mg/kg, IP) did not exert any effect on locomotor activity (Figure 3).

4. Discussion

As mentioned earlier in the introduction, research on the analgesic effects of cannabinoids has been carried out many times. However, despite many established similarities between the mechanisms of pain and itch (Cevikbas, Steinhoff, & Ikoma, 2011; Ross, 2011; Schmelz, 2010), the effects of cannabinoids on itch has remained under-researched. The few existing studies on the matter demonstrated that cannabinoid receptor agonists decrease scratching behavior. Also, cannabinoid receptor antagonists like rimonabant, and a brain penetrating CB1 inverse agonist, dose-dependently increase drug-induced scratches (Darmani & Pandya, 2000; Janoyan, Crim, & Darmani, 2002; Schlossburg et al., 2011; Todurga et al., 2016). Peripheral administration of the potent synthetic cannabinoid HU210 has also been reported to suppress histamine-induced scratches (Dvorak et al., 2003). In accordance with this research and our previous report (Todurga et al., 2016), this study indicates that the cannabinoid receptor agonist WIN 55,212-2 diminishes serotonin-induced scratches dose-dependently. WIN 55,212-2 caused an insignificant decrease in the number of scratches at the dose of 1 mg/kg, whereas it almost ceased scratching behavior at higher doses (3 and 10 mg/kg).

NO modulates neuronal function and contribute to various biological tasks in the central nervous system. Moreover, the influence of NOS system on cannabinoid activity has been shown; activation of cannabinoid receptors by anandamide led to NO production (Prevot et al., 1998), and CB1 receptors stimulated cyclic GMP production in neuronal cells (Carney et al., 2009; Jones, Carney, Vrana, Norford, & Howlett, 2008). NO has also been suggested to play important roles in the development of tolerance to some effects of cannabinoids. It has also been indicated that NO is involved in the development of tolerance to the hypothermic and catalepsy effects of cannabinoids (Azad et al., 2001; Spina, Trovati, Parolaro, & Giagnoni, 1998), whereas a negligible contribution to cannabinoid

![Figure 1](image-url). Effects of systemic administration of the cannabinoid agonist WIN 55,212-2 (1, 3, 10 mg/kg, IP) on serotonin-induced scratches. * P<0.0001, compared with the control.
analgesic action has been proposed (Azad et al., 2001; Spina et al., 1998; Thorat & Bhargava, 1994). In contrast, it has been suggested that L-arginine/NO pathway is involved in the development of tolerance to the analgesic action of the cannabinoid agonist WIN 55,212-2 (Banafshe, Ghazi-Khansari, & Dehpour, 2005). Given the similarities between the mechanisms of pain and itch, we observed that NO modulation did not influence the antipruritic effect of cannabinoids. Neither endothelial and neuronal NOS inhibitors nor the NO precursor had any effect on the antipruritic action of the cannabinoid receptor agonist WIN 55,212-2. Thus, NO does not play any role in the antipruritic effect of cannabinoids.

Administration of NOS inhibitors and the NO precursor L-arginine on their own revealed different results on serotonin-induced scratches. Neither the non-selective NOS inhibitor L-NAME nor the NO precursor L-arginine had any effect on serotonin-induced scratches. However, the nNOS inhibitor 7-nitroindazole significantly reduced serotonin-induced itch behavior. In another research, in line with our findings, L-NAME had no effect on serotonin-induced scratches, whereas the iNOS inhibitor aminoguanidine prevented itching behavior. Researchers pointed to the involvement of NO in serotonin-induced itching and highlighted the importance of iNOS in the process, yet they had not used

**Figure 2.** Effects of the non-selective NOS inhibitor L-NAME (3 mg/kg, IP), selective nNOS inhibitor 7-nitroindazole (3 mg/kg, IP) and the NO precursor L-arginine (100 mg/kg, IP) on serotonin-induced scratches and the antipruritic effect of WIN 55,212-2 (3 mg/kg, IP). * P<0.0001, compared with the control.

**Figure 3.** Effects of WIN 55,212-2 (1, 3, 10 mg/kg, IP), L-NAME (3 mg/kg, IP) and L-arginine (100 mg/kg, IP) on locomotor performances. * P<0.0001, compared with the control.
nNOS inhibitors (Ostadhadi, Haj-Mirzaian, Azimi, Mansouri, & Dehpour, 2015). Another report concluded that NO/cGMP pathway mediated chloroquine-induced itching and that specifically nNOS inhibition had a role in the process (Foroutan, Haddadi, Ostadhadi, Sistany, & Dehpour, 2015). Conclusions from these studies are consistent with our results, all suggesting the importance of nNOS inhibition in experimental itch models. It should also be known that the involvement of eNOS cannot be excluded since 7-nitroindazole also inhibits eNOS at higher doses (Ayajiki, Fujioka, Okamura, & Toda, 2001).

Our findings show that intraperitoneal injection of the cannabinoid receptor agonist dose-dependently decreased serotonin-induced scratching. Mechanism of action of WIN 55,212-2 may include both cannabinoid CB1 and or CB2 receptors since it is a non-selective CB1/CB2 agonist. Based on the currently presented data, it cannot be decided whether the actions are realized at the level of the central nervous system or already at the periphery. Peripherally restricted CB1 agonists could have been used to minimize central side effects, but in this case, peripheral side effects should be taken into consideration and this was not the purpose of this study. Another possible mechanism for cannabinoids is that they may also inhibit Transient Receptor Potential (TRP) channels, especially TRPV1, which plays an important role in sensing itch as well as pain (Morita et al., 2015; Ross, 2011; Schmelz, 2010). In our study, neither the endothelial and the neuronal NOS inhibitors nor the NO precursor influenced the antipruritic action of cannabinoids. On the other hand, the neuronal NOS inhibitor decreased scratching when used alone. We concluded that NO does not mediate the antipruritic action of cannabinoids and neuronal NOS inhibition plays a role in serotonin-induced itching. Whatever the mechanism of action, we believe that cannabinoids have the potential to be used as antipruritic drugs, if their side effects can be reduced.

Ethical Considerations

Compliance with ethical guidelines

Adequate measures were taken to minimize pain or discomfort, and "Animal Care Ethics Committee" of Trakya University approved all experimental protocols of this study (TUHADYEK-2015/43).

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Authors' contributions

Conceptualization: Ozgur Gunduz, Ahmet Ulugol; Methodology: Investigation, Funding Acquisition: Resources: All authors; Writing – original draft, writing – review & editing: Ozgur Gunduz, Ahmet Ulugol; Supervision: Ahmet Ulugol; All authors have read and approved the manuscript before submission.

Conflict of interest

The authors declare no conflict of interest.

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