Honey and Black Seed Synergistically Promote Regeneration of Oligodendrocytes in Cuprizone Intoxicated Quail Brain

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ABSTRACT

Multiple sclerosis (MS) is a complex disorder, characterized by demyelination and loss of axonal parts of neurons in the central nervous system involving multiple genetic and environmental factors. Although, the demyelinated lesions develop throughout the brain, but more frequently are extensive in white matter. Currently, three different approaches are being utilized to treat MS, where synthetic drugs are the most frequently used but they do not cure the disease. Secondly, the stem cell therapy, but this too has limited success in treating MS in humans. The third technique involving administering hormones has been found to be most effective method but this too have some significant side effects. Alternatively, natural products can potentially serve as an affordable and effective substitute for the treatment of MS with minimum or no side effects. Blackseeds (Nigella sativa) and honey possessing potent neuroprotective, antioxidant, and anti-inflammatory properties with no reported side effects can be a prospective candidate for an alternate remedial treatment of MS in animal model as well as in humans. In this study we established a new animal model (quail), to assess the synergistic efficacy of honey and black seed against demyelination within brain. A total of 35 male quails were used, among 10 were non treated and 25 were treated with 200 mg/kg/day cuprizone (CPZ) demyelination for six months to induce demyelination. After that they were divided into seven groups of five animal each where 3 CPZ treated groups received either honey, black seed oil or mixture of both for 6 weeks after demyelination. Behavioral tests were performed at the end of treatment. Afterwards, oligodendrocyte population was estimated in cerebellar white matter after histology. It was found that all three treatments efficiently induce remyelination. Interestingly, the mixture of honey and black seed was significantly more efficient than honey and black seed alone. Our data support the need of clinical trials for administration of N. sativa and honey in MS patients.

INTRODUCTION

Within the central nervous system, oligodendrocytes are specialized glial cells that form myelin sheath around axons facilitating the saltatory conduction, ultimately enhance nerve transmission efficiency (Mitew et al., 2014). Alterations in the structure or function of these cells lead to numerous disabilities, more frequently multiple sclerosis (MS). It is a heterogeneous disease with unknown autoantigens which are responsible for induction of disease. MS develops due to an imbalance between demyelination and remyelination resulting into permanent demyelination of axons leading to cerebral atrophy at earlier stages and breakdown of blood brain barrier at later stages of the disease (Coles et al., 2008; Jakimovski et al., 2019). It is well established that MS is a disease induced by both genetic and environmental factors (Dendrou et al., 2015) and often more prevalent in females as compared to males (Hollenbach and Oksenberg, 2015). Recent therapeutic approaches for MS involves prolonged use of medications merely to reduce the symptoms and slowdown progression of the disease.
(Ghaiad et al., 2017). Hormonal therapy and stem cell therapies like embryonic stem cell, and mesenchymal stem cells, on the other hand, have been tested in animal models with considerable success but have limitations for humans use (Rivera et al., 2019; Hohlfeld, 2021). In general, the existing drugs merely focus on the prevention of the symptoms while their long-term usage has further adverse side effects in addition to high costs (Alfaqih, 2020). To reduce side effects, and increase cost effectiveness of existing drugs, herbal medicines can be a potential candidate for effective treatment against MS (Mojaverrostami et al., 2018). Honey is a natural compound mainly comprised of carbohydrates, water and a small quantity of nitrogen, organic acids, minerals and vitamins which enhance its biological potential for therapeutic usage (Machado et al., 2018). Honey exhibits antimicrobial, antioxidant, anti-inflammatory, anticancer, anti- hyperlipidemic, and cardioprotective properties (Vazhacharickal, 2021) due to bioactive compounds such as polyphenols with reported protective roles in cardiovascular diseases (CVD), diabetes, cancer, gastrointestinal tract diseases and neurodegenerative diseases (Hossen et al., 2017).

*Nigella sativa* L. (Ranunculaceae) (black seed) is an annual flowering plant (Mojaverrostami et al., 2018) found in Southern Europe, Southwest Asia and North Africa (Tavakkoli et al., 2017). A number of studies suggest that *N. Sativa* is beneficial for the treatment of diseases associated with cardiovascular, nervous system, skin, infectious, reproductive system, respiratory system, skeletal system, and gastrointestinal system (Ahmad et al., 2013). Being herbal medicating containing biologically active compounds, such as thymoquinone, nigellone and melatin, black seeds are being tested in clinical trials against neurological disorders (Ahmad et al., 2013; Tavakkoli et al., 2017). It is also reported to have a role in cell survival, cell cycle and anti-apoptotic activity (Hannan et al., 2021) as well as it potentially ameliorates brain function related to cognition, depression, epilepsy, and memory due to high proportions of polyunsaturated fatty acids in black seeds composition. These compounds are essential for protecting nervous system against any neuronal injury or disorders (Hala et al., 2006). The reported therapeutic potential, easy availability, cost effective processing and no side effects of using black seeds render it an ideal candidate for treatment of MS using an animal model. In this study we have evaluated the therapeutic potential of honey, black seed and their combinations in regeneration of oligodendrocytes in cuprizone induced quail brain. The results will be helpful in validating the therapeutic potential of mixture of honey and black seed, and designing a new drug against MS with no side effect.

**MATERIALS AND METHODS**

**Site of study and ethical approval**

This study was conducted in the Department of Biosciences of COMSATS University, Islamabad. All the ethical and biosafety approvals were sought and granted by departmental ethical review committee.

**Animal model**

Due to unavailability of the relevant animal model for MS, this study was done in a new animal model, quail (*Coturnix japonica*). Quails have been proving an important tool in disease research (Baer et al., 2015). Since 1940, Quails have been utilized in genetics research (Shimakura, 1940) and have become an important model in developmental biology, behavior and biomedical studies (Minvielle, 2009). They have also been used as an animal model for albinism (Homma, 1968). The quail are relatively smaller in size and are easier to rear than chicken or even mouse (Hüss et al., 2008).

Adult males (14 weeks old) were obtained from Hamza Chicks Islamabad, Pakistan. Animals were housed under standard laboratory conditions at the Institute and fed with standard chicken diet and continuous supply of fresh drinking water. Demyelination was induced with CPZ at a dose of 200 mg/kg/day for 6 months by mixing the (CPZ, Sigma) powder in feed pellets. Weekly body weights of each animal were recorded basis throughout the treatment period along with clinical observation.

**Treatment**

From the laboratory reared animals, seven groups for bioassay were designated with five animals in each group (Table I). Animals in group 3 to 7 were exposed to a daily dose of CPZ while animals in group 1 and 2 were kept untreated. At the end of 6 months, animals in group 1 and 3 were slaughtered to assess the extent of demyelination. Animals in group 5 and 6 were given a daily dose of honey and black seed through oral gavage at 100 mg/kg/day respectively for a period of 6 weeks after induction of demyelination. Both honey and black seeds were administered in mixture, in equal quantity, to the animals of group 7 at a dose of 200 mg/kg/day for 6 weeks. Animals of group 4 did not receive any treatment. Animals in all the groups were perfused in 4% paraformaldehyde at the end of 6 weeks to isolate brains.

**Tissue processing for histological analysis**

Histological assessment was done as described by Akram et al. (2018). Tissues were fixed in a mixture of ethanol (60 ml), formaldehyde (30 ml), and glacial acetic acid (10 ml) for 24–48 h, and then proceeded for
embedding in paraffin. Sections of 8-10 μm thickness were cut on a microtome (Shandon, Finesse 325, Thermo Fisher Scientific, Luton, England) and stained with Harris hematoxylin (Sigma Aldrich, 638A) and eosin (Sigma-Aldrich, E4009). Sections were examined at different magnifications with light microscope (Olympus CH40 Olympus Optical Co. Ltd 2-43-2 Tokyo Japan Model BX41TF SN 5L20405) and focusing on the white matter of cerebellum. Images were captured using a microscope mounted camera (Model DP12-2 SN 5094926 B4 Tokyo Japan) and the number of cells were counted by processing the images with Image J software.

### Table I. Animals groups and treatment.

<table>
<thead>
<tr>
<th>Animal groups</th>
<th>Treatment</th>
<th>Exposure</th>
</tr>
</thead>
<tbody>
<tr>
<td>G1</td>
<td>Positive control</td>
<td>No. for 6 months</td>
</tr>
<tr>
<td>G2</td>
<td>Positive control</td>
<td>No. for 7.5 months</td>
</tr>
<tr>
<td>G3</td>
<td>Cuprizone</td>
<td>6 months</td>
</tr>
<tr>
<td>G4</td>
<td>Cuprizone</td>
<td>6 months + 6 weeks no treatment</td>
</tr>
<tr>
<td>G5</td>
<td>Cuprizone+Honey</td>
<td>6 months + 6 weeks</td>
</tr>
<tr>
<td>G6</td>
<td>Cuprizone+Black seed</td>
<td>6 months + 6 weeks</td>
</tr>
<tr>
<td>G7</td>
<td>Cuprizone+Honey and black seed</td>
<td>6 months + 6 weeks</td>
</tr>
</tbody>
</table>

### Behavioral bioassays

Prior to slaughtering of the animals, following behavioral studies were performed.

**Beam walking assay**

The beam walking test was performed in accordance with several studies (Carter *et al.*, 2001; Liu *et al.*, 2009; Chen *et al.*, 2011; Luong *et al.*, 2011). Albeit with modification. We overcome the animal size, as rectangular beam, by adjusting the diameter or rounded shape to support the movement of these quails. The set up consisted of one rectangular wooden beam of 2 cm width and 100 cm length. The beam was graduated from score 1-20. The ends of the beam were placed on two wooden blocks, 15 cm long, 15 cm wide and 30 cm high. These dimensions of the blocks allowed beam to stay at an elevation of 30 cm from the ground. The quail were trained on the beam twice a day for three days before test.

Experiment was performed with control and treated quail at the end of treatment i.e., CPZ and honey, black seed and the mixture. These quails were compelled to walk from one end of the graduated beam to the other. We recorded the graduated score of the beam where quail feet slipped. Maximum score 20 was given to the quail which crossed beam without any slip. Three observations were made for each quail get average scores ± SE.

### Flight assessment

Quails were stimulated to fly from a fixed height (150 cm). The length of their flight was measured and the quality of rhythmic fanning of wings and take off/landing were recorded. The data was presented as the average length of flight ± SE.

### Statistical analysis

Weekly body weight, beams score, flight length and number of cells (counted) in central part of cerebellum of all the treated and untreated control groups were recorded. The obtained data were expressed as mean ± SEM and analyzed using ANOVA after conforming the assumption of data homogeneity. Where the significant differences were observed, the pair wise post-hoc comparisons were performed using Tukey’s test at α =0.05. The data analyses were conducted using Statistical Package for the Social Sciences for Windows (SPSS 22.0, IBM SPSS Inc.) and GraphPad Prism 8.

### RESULTS

#### Body weight

Although not significantly different, a slight increase in weekly body weight of the animals was recorded when they were treated with black seed alone or together with honey (Fig. 1).
Behavioral studies

Beam walk test

The beam walk score showed significant difference between the control and CPZ treated animal. Similarly, all the treated groups i.e., honey, black seeds, and mixture treated groups showed significantly lower score than positive control but significantly higher than CPZ treated animal. Interestingly, there was no significance difference between animal groups treated with honey, black seed, and the mixture (Fig. 2).

Fig. 2. Beam walking score for five animal groups. Control group had significantly higher score as compared to all other groups. Honey, black seed and mixed groups showed significantly higher scores than CPZ treated animals. Scores are presented as mean ± SEM. Boxes with the same alphabet are not significantly different (P ≤ 0.05).

Table II. Oligodendrocyte number in demyelinated and remyelinated animals.

<table>
<thead>
<tr>
<th>Animal groups</th>
<th>No of cell in central cerebellar white matter</th>
<th>Percent- age (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Positive control 6 months</td>
<td>115</td>
<td>100</td>
</tr>
<tr>
<td>Positive control 7.5 months</td>
<td>122</td>
<td>100</td>
</tr>
<tr>
<td>Cuprizone 6 months</td>
<td>53</td>
<td>46.08</td>
</tr>
<tr>
<td>Cuprizone 7.5 months</td>
<td>76</td>
<td>62.29</td>
</tr>
<tr>
<td>Honey</td>
<td>80</td>
<td>65.50</td>
</tr>
<tr>
<td>Black seed</td>
<td>84</td>
<td>89.3</td>
</tr>
<tr>
<td>Honey and black seed</td>
<td>109</td>
<td>89.30</td>
</tr>
</tbody>
</table>

Flight test

Fig. 3. Flight length test results. Control group showed highest score followed by natural products treated groups and CPZ intoxicated group. Control group shows significantly higher length of flights than any other group. Treated groups have flight lengths less than control but significantly higher than animal treated with CPZ only. Data was presented as mean ± SEM. Bar with same alphabet means no significant difference (P ≤ 0.05).

Histopathological assessment

After six months of CPZ treatment, thickness of cerebellar white matter decreases as compared to positive control group (Fig. 4C, D, respectively). The central white matter of cerebellar and cerebrum showed more apoptotic glial cells, rare alive oligodendrocytes and shrinkage of axons in the groups treated with CPZ for six months compared to control group (Fig. 4E, F) cerebellar white matter, and (Fig. 4G, H) cerebral white matter). Effective remyelination can be seen in CPZ treated animal when left untreated for 6 weeks (Fig. 4J). Honey, black seed and the mixture treated groups showed variable rate of oligodendrocyte regeneration (Fig. 4K, L, M).

Cells counting

The population of oligodendrocytes reduces to 45% after six months of CPZ treatment (n=5) (Fig. 5, Table II). A spontaneous regeneration can be observed after the removal of CPZ treatment and oligodendrocyte number rises to 62% at the end of experiment i.e., 7.5 months (n=5). Animals treated with either honey (n=5) (65%) or black seed (n=5) (68%) also displayed increase in oligodendrocyte number but difference was non significant as compared to cuprizone only treated (n=5) animals at the end of experiment.
Fig. 4. Cerebellar and cerebral white matter (H & E x 20 and x 40). (A) positive control entire brain (1: cerebellum; 2: optic lobe; 3: cerebral hemisphere; 4: olfactory lobe). (B) positive control sagittal brain section (5: cerebral cortex; 6: cerebellar cortex; 7: deep cerebellar white matter; 8: pons; 9: medullar oblongata. (C) positive control after six months shows normal thickness of cerebellar white matter folds. (D) six months CPZ treated animals show loss of cerebellar thickness in white matter folds. (E) control group kept for six months without CPZ shows deep cerebellar white matter and normal oligodendrocyte population. (F) animals with CPZ treatment for six months show rare oligodendrocytes. (G) cerebral white matter of control animal after six months shows normal glial cells. (H) cerebral white matter of six months CPZ treated animal shows apoptotic oligodendrocytes (arrow) and shrinkage of axons. (I) normal oligodendrocyte population can be seen in control animal after seven and half months without any treatment. (J) spontaneous oligodendrocyte regeneration can be seen in animal with no treatment for six weeks after six months of CPZ treatment. (K) honey treated group showing regeneration of oligodendrocytes. (L) black seed treated group showing increase in glial cells population. (M) Mixture (honey and black seed) treated group shows significantly increased oligodendrocyte population.
Interestingly, animal group treated with honey and black seed mixture (n=5) showed significantly higher number of oligodendrocytes (89%) as compared to all other groups except control group. This is clear indication that honey and black seed mixture efficiently promotes oligodendrocyte regeneration in quail white cellular matter.

![Graph showing number of oligodendrocytes in different groups](image)

Fig. 5. The number of oligodendrocytes in cerebellar white matter of quail is depicted in the graph. There is a severe depletion of oligodendrocytes after six months of CPZ treatment. Oligodendrocyte no. increases after the arrest of treatment. A non-significant difference in the number of oligodendrocytes can be observed in CPZ, honey and black seed group. Animals in mixture group have significantly higher number of oligodendrocytes than the honey, black seed and CPZ group. Oligodendrocyte number has been presented as mean ± SEM. Bar with same alphabet means no significant difference (P ≤ 0.05).

**DISCUSSION**

Multiple sclerosis is a complex inflammatory autoimmune and neurodegenerative disorder that leads to the loss of myelin sheath and parts of axons in the central nervous system (CNS). The history of CPZ model begin more than 50 years ago (Peterson-Bollier, 1955). The direct or indirect supplementation of normal animal chow with CPZ lead to oligodendroglia cell death and consequent demyelination (Lindner et al., 2009). Though many models have been used for MS studies, CPZ is considered as one of the most widely used model to study demyelination and remyelination (Hussain et al., 2013; Zhan et al., 2020). It also helps to study intercellular interaction between glial cells (Remington et al., 2007).

In humans, the ventral zone of the spinal cord is the site of origin of oligodendrocytes that express myelin genes and platelet derived growth factor-α (PDGFαR). In birds and rodents, oligodendrocytes of the dorsal spinal cord initially originate from the ventral half of the spinal cord and then migrate towards the dorsal side. This indicates the common origin of oligodendrocytes in both vertebrate classes (Remington et al., 2007). Other investigations have reported similar origin between birds and human oligodendrocytes (Miller and Bell, 1996). Till date no study has been conducted to use quail as an in vivo model of MS.

After one week of CPZ induced demyelination, oligodendrocyte mitochondria get affected which leads to oligodendrocytes apoptosis and consequently downregulation of many myelin associated protein (Komoly, 2005; Gudi et al., 2011). It is well established that 0.2% CPZ induces extensive and reproducible demyelination without detrimental systemic effects in adult 8-10 weeks old mice. However, the effect differs among mouse strains (Irvine and Blakemore, 2006). Either 400 mg/kg/day (Zhen et al., 2017) or 200 mg/kg/day (Gudi et al., 2014) is recommended for obtaining extensive and consistent demyelination after 5 weeks of administration in rodents. Six months were required to obtain similar results in quail model. This may be due to double number of neurons in birds than mammals (Olkowicz et al., 2016). Although this bring novel insight on the duration to induced demyelination in birds, further study is required to find out the best CPZ doze. Demyelination is more prominent in the oligodendrocyte rich areas of the central nervous system such corpus callosum and cerebellar white matter (Silvestroff et al., 2012). As quail brain lacks a clearly defined corpus callosum so we focused on quail’s cerebellar white matter to assess the potential of honey and black seed in the regeneration of oligodendrocytes post CPZ administration.

Recent approaches tend to improve or even stop disease progression in patients, with primary progressive MS (PPMS) and secondary progressive MS (SPMS), by stimulating the neuroprotective activity or inhibiting the inflammation or promoting the spread of innate immune reaction and inducing the remyelination by promoting OPCs proliferation and differentiation (Zhan et al., 2020). In relapsing remitting MS, commercially available drugs are useful to regulate the activity of immune cells and episodes of the disease. Due to continuous loss of neuronal function in progressive stages of disease, stimulation of the endogenous remyelination could be a great progress in MS therapies (Kremer et al., 2019).

Medicinal plants are thought to have beneficial effects in cancers, diabetes and neurodegenerative diseases (Dehghan et al., 2016). An increased number of studies have emphasized on the herbal compounds to improve myelin repair (remyelination) and suppress the inflammation and symptoms (Piao and Liang, 2012).

Body weight is an important parameter to assess the health or disease condition of an animal. However, in the present study we did not find significant body weight loss in animals. The motor impairment was also less prominent...
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among the CPZ treated animals though we notice some dizziness and seizures. Approximately 5% of body weight loss was observed after six months of CPZ treatment in quails as compared to a loss of 10% in mice model fed with 0.2% CPZ for five weeks (Steelman et al., 2012). Either dose of CPZ was not sufficient to induce severe demyelination or the number of oligodendrocytes is higher in the quail brain because songbirds contain roughly the double number of neurons as compared to same brain size of mammals (Olkowicz et al., 2016). Although incomplete, the spontaneous remyelination was remarkable in the animals fed with CPZ for six months and exempted of any treatment for six weeks. The body weight was slightly increased at seven and half month compared to six months weight. The difference observed in the beam walk test and flight length test between the CPZ treated group and honey and black seed treated group further strengthened the existing and incomplete endogenous remyelination. Moreover, the no. of counted oligodendrocytes in positives control and negatives control groups clearly demonstrate spontaneous and incomplete remyelination. The incomplete endogenous remyelination always occurs in progressive form of MS (Kremert et al., 2019) which underlines the significance of medication to continuously stimulate the innate remyelination.

Honey is a natural compound that can be obtained from the plant nectar (Rodríguez et al., 2019). Its main properties differ based on the botanical and geographical origin, climate, and processing. It is mainly composed of carbohydrates, water with minor quantities of other ingredients like nitrogen, organic acids, minerals and vitamins which influence its biological potency (Machado et al., 2018).

Behaviorally, the observed difference between honey treated group and CPZ treated group at the end of experiment prove that honey enhances the endogenous remyelination. Comparing the numbers of oligodendrocytes in the honey treated group and CPZ treated, we observed that honey induces remyelination after arrest of CPZ treatment at the end of six months. Albeit the first study assessing the effect of honey in MS animal model, our data confirmed the anti-inflammatory and antioxidant potency of honey linked to one of his mains compounds, named polyphenols (Vazhcharickal, 2021; Hossen et al., 2017). In MS treatment, the strategies are, stopped the inflammation, increased the antioxidant enzymes and promoted the remyelination (Zhan et al., 2020).

*N. sativa* L. usually known as black seed or black cumin is being used for more than thousand years, due to its pharmacological properties (Mojaverostami et al., 2018). Thymoquinone is the most abundant constituent of the volatile oil of black seed, and is thought to be responsible for the most properties of the herb (Tavakkoli et al., 2017). Thymoquinone, is also well known for the removal of free radical and superoxide ions (Akhtar et al., 2014). This role conserve the activity of various antioxidant enzymes, such as catalase, glutathione peroxidase and glutathione-S-transferase (Woo et al., 2012).

Black seed has the potential to attenuate oxidative stress by activating the antioxidant defense system (Nrf2 signaling), inhibit inflammation by activating anti-inflammatory signaling (NF-κB and TLR signaling), induce immunity by modulating innate and adaptive immune components, prevent apoptosis by upregulating pro-survival signals and downregulating pro-apoptotic signals (PI3K/Akt, JNK, and mTOR signaling) (Hannan et al., 2021).

These properties are further confirmed by our data where a significant difference of beam walk and flight test scores was observed between animal treated with black seed and the CPZ treated animals. The counted number of oligodendrocytes in black seed treated group compared to CPZ treated group also demonstrated the capability of black seed to accelerate remyelination after the damage caused by CPZ. Our results are in agreement with (Noor et al., 2015; Fahmy et al., 2014) who reported enhanced remyelination in the CNS after *N. sativa* treatment in experimental autoimmune encephalomyelitis (EAE) model of MS. It was further described to reduce inflammation processes. The neuroprotective and antioxidant potentials of black seed have also been elucidated in lead induced neurotoxicity during development and early life in mouse models (Butt et al., 2018) and in rotenone induced Parkinson’s animal model (Ebrahimii et al., 2017).

After behavioral and histological analysis, the mixture of honey and black seed showed significantly higher beam walk and flight tests scores and oligodendrocyte population. Although the mixture is more efficient than the other treatment groups, it does not completely rectify the damage induced in the animal by CPZ. That may be due to short duration of the honey and black seed treatment. After discontinuation of CPZ treatment, the mixture of honey and black seed reduces inflammation and promote the recruitment and regeneration of OPCs. The loss of myelin can be restored during remyelination, which can prevent axonal degeneration and restore motor function.

The complementary effect of mixture of honey and black seed, revealed a new therapeutic dimension of the disease. A comparison of the efficacy of honey, black seed oil and the mixture has demonstrated that; honey, black seed and the mixture have accelerated the spontaneous regeneration of oligodendrocytes at 3%, 6%, and 27%, respectively as compared to CPZ treated animals at the end of experiment.
Overall, honey, black seed and the mixture have induced 20%, 23% and 43% regeneration of oligodendrocytes respectively in comparison with CPZ treated group at the end of six-month group. Black seed displayed 4% higher regeneration than honey while and the mixture effect was 20% and 24.6% superior than black seed and honey, respectively.

Honey is known world-wide as a medicine, energy source and well recognized for biological, physiological and pharmacological activities (Amin et al., 2018). In the universal honey phenolic acids, flavonoids, and antioxidants are found with synergic action (Bogdanov et al., 2008). Glucose, fructose, flavonoid, polyphenols and organic acids are thought to play a major role in quality and health benefits of honey (Cianciosi et al., 2018). Phenolic compounds have well established antioxidiant, antimicrobial, antiviral, anti-inflammatory, antifungal, wound healing, and cardio protective properties (Viuda et al., 2008). Medically honey has been found to have antioxidiant, anti-inflammatory, anti-bacterial, antiviral, anti-ulcer, anti-hyperlipidemic, antidiabetic and anticancer properties (Juszzak et al., 2016). The presence of phenols and flavonoids in honey is believed to be responsible for the exhibition of anti-oxidative and anti-inflammatory activities (Ahmed et al., 2018). Black seed also contains many anti-inflammatory and antioxidiant components like polyphenols, quercetin, kaempferol and thymoquinone which are efficient in relieving the oxidative stress by activating the antioxidiant defense system and inhibiting the inflammation by activating anti-inflammatory signaling (Hannan et al., 2021).

Similar anti-inflammatory, and antioxidiant potential of honey and black seed is essential for the therapeutic application in MS animal model. Collectively they promote efficient remyelination and recovery of saltatory action potential conduction to restore motor impairment.

Our data confirmed the therapeutic potential of honey and black seed in the MS animal model. However, more studies are required to qualitatively assess the effects of each of these medicinal products and their active ingredients.

CONCLUSION

This study is the first attempt to assess the therapeutic effects of honey, black seed, and the mixture in CPZ induced demyelinated quail model. After six months of CPZ induced chronic demyelination and six weeks of honey and black seed oil treatment, we found that honey and black seed oil are capable to induce and accelerate the remyelination and the mixture of two is more efficient than both separately. Our findings stress the need of future experiments to identify and characterize active ingredients to design effective drugs with low cost and no adverse side effects.

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IRB approval

The study project was approved by the Ethics Review Board and Institutional Biosafety Committee of Department of Biosciences, COMSATS University, Islamabad, 45550, Pakistan after due deliberations.

Ethics approval and consent to participate

Not applicable.

Consent for publication

Not applicable.

Availability of data and materials

All the generated and analyzed data of this study are included in this article and its supplementary material files.

Statement of conflict of interest

The authors have declared no conflict of interest.

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