INTRODUCTION

Inflammation is an evolutionary process in the body that gain a great benefit for maintaining homeostasis. The inflammation may cause due to a microbial infection, cellular damage, stress and other chemicals irritants, etc. [1]. Among vertebrates, the inflammatory cascade is a complex process associated with immunological, physiological, and behavioral responses associated with cytokines and other immunological signaling molecules [2]. The cascade mainly involves an increase in microvesSEL permeability, migration of cell types, the growth of new tissues, and blood vessels [3]. During inflammation, a diverse group of molecules or substances called as pro-inflammatory mediators such as histamines, bradykinins, nitric oxide (NO), serotonin, and prostaglandins was released. These substances may significantly contribute to the classical picture of heat, redness, pain, selling, and diminished function associated with inflammation and may cause hyperalgasia or allodynia [4]. Acute inflammation refers to the initial response of the immune system against foreign bodies and tissue injury. It is a rapid time-limiting process, mediated by various substances such as eicosanoids and vasoactive amines which significantly stimulate the plasma movement and leukocytes into the inflammation site [5]. During tissue inflammation, increases in vasodilatation and recruitment of capillaries and increases in vascular permeability cause extravasations of plasma to lead to tissue edema. This causes migration of leukocytes, especially neutrophils [6]. Free radical species such as hydrogen peroxide (H$_2$O$_2$), NO, molecular oxygen (O$_2$), and superoxide radicals may significantly contribute to tissue inflammation. Especially, NO plays a key role in cell survival and cell death, cause pro-inflammatory effects on various immune and associated cells [7]. Increases in NO show the significant increases in macrophages in inflammatory sites.

It has been reported that currently available anti-inflammatory drugs which include NSAIDs (nonsteroid anti-inflammatory drugs) and opioids were not functioning in all cases, due to their low potency. These drugs cause side effects on gastrointestinal complications such as ulcers, perforating, vomiting or nausea, bleeding, constipation, and cognitive impairment [8]. As a result, the identification or development of novel alternatives has necessary and beneficial [9]. Previous studies taken up by various research groups showed that the effect of plant-derived lectins can alter the immune system and thereby it blocks the inflammation cascade. Lectin from Caulerpa cupressoides possesses anti-inflammatory activity by inhibiting the leukocyte migration in a murine model of inflammation and lectin from Synadenium carinatum inhibits recruitment of immune cells to the lung in an asthma murine model by altering the expression of transcription factor necrosis factor (NF-κB) [10,11]. In the present report, we investigate the partially purified lectin, Praecitrullus fistulosus lectin-like protein (PfLP) which possesses anti-inflammatory activity against carrageenan-induced paw edema in mice model. We also evaluate the NO content in paw homogenate tissues. Decreases in NO content in treated group support the anti-inflammatory activity of PfLP. In future dietary, lectin may play a key role to develop a novel therapeutics against various diseases associated with inflammation.

METHODS

Materials
All the chemicals used in the present study were procured from Sigma-Aldrich and Hi-Media.

Collection and authentication of plant material
Samples were collected from local area farmers, Mysore, and authenticated by Dr. Sharvani KA, Assistant professor, Department of Botany, Yuvaraja’s College, University of Mysore, Mysuru. The herbarium (Accession number: YCM(UOM)0266) was prepared and deposited at the Department of Botany, Yuvaraja’s College, University of Mysore.
Preparation of phloem exudates

Phloem exudates were processed and prepared as described earlier and the partially purified lectin sample was prepared and named as PfLP [12]. Protein concentration was determined using the Bradford assay as described earlier [13].

In vivo anti-inflammatory activity

Animals

Male Swiss albino mice (27±2 g) were obtained from the Central animal facility, DOffs in Zoology, University of Mysore, Mysuru. The animals were maintained as per the Institutional Animal Ethical Committee (IAEC-Ref No: UOM/IAEC/10/2017) guidelines. They are fed with standard diet and water ad libitum procured from Krishna enterprises, Bengaluru. All animals were acclimatized for 1 week with a constant temperature of 22±2°C, with 12 h dark-light cycle, before the experimental sessions.

Preparation of 1% of carrageenan

Carrageenan was procured from Sigma-Aldrich. 1% of carrageenan was prepared by dissolving 0.1 g of carrageenan in 10 ml of 0.9% NaCl.

Carrageenan-induced paw edema assay

Anti-inflammatory activity of PfLP was evaluated by the acute inflammatory study as described earlier with slight modifications [14]. Acute inflammation was induced using 1% of carrageenan (100 µl/paw) into the right hind paw of each mouse. The edema was induced by injecting into subplantar tissue of the right hind paw of each mouse. Group I receives saline acts as control, Groups II and III receive 25 mg/kg and 50 mg/kg of PfLP respectively, and Group IV receives diclofenac (10 mg/kg) were administered intraperitoneally. 1 h before the injection of carrageenan and paw volume was measured using Vernier caliper and leg weight was recorded at post scarification of the animals.

Histopathology staining

Histological changes of treated and untreated samples were examined as described earlier in section. Briefly, collected paw edema tissues from various groups were fixed using 4% paraformaldehyde and embedded in paraffin wax and 10 µm sections were made using microtome (SLEE Cryostat). The sections were observed under low power (910X) light microscope to identify the histological changes.

MPO assay

Myeloperoxidase (MPO) activity was determined to evaluate the neutrophils concentration in the inflammation tissue, as described earlier with slight modifications [15]. Briefly, tissue from paw was homogenized using 1 ml of buffer (5 g HTAB/L in 50 mM potassium phosphate buffer, pH 6) using homogenizer, and homogenized samples were subjected to centrifugation at 10,000 rpm at 4°C for 10 min and supernatant was subjected to enzyme activity, as an index of migration of cells, using the method as described earlier [16]. Obtained 100 µl of the supernatant was incubated with 2.9 ml of phosphate buffer containing substrate (0.167 g mg/ml of O-dianisidine with 0.0005% H₂O₂). The absorbance was read spectrophotometrically at 458 nm using visible light for 2 min. Enzyme activity per gram tissue was calculated as follows:

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\text{MPO activity} \left(\frac{\text{Units}}{\text{gm}}\right) = \frac{(\Delta A460) \times (13.5)}{\text{paw tissue weight (g)}}
\]

Where, ΔA460 was changing in absorbance at 460 nm from 30 to 90 s (2 min measurements), 13.5 is coefficient determined that 1 unit MPO activity was the amount of enzyme that would reduce 1 mM peroxide/min.

No production assay

The nitrite level in carrageenan-treated control and treated mice was determined as described earlier [17]. Homogenize clear supernatant was used as an indicator of NO production using Griess reaction. Briefly, the 150 µl of supernatant were incubated with an equal volume of Griess reagent (1% sulfanilamide prepared in 3 N HCl and 0.1% N-ethylene dihydrochloride) for 10 min at room temperature, and absorbance was read at 540 nm using a microplate reader and nitrite concentration was determined by comparing with standard graph construct using sodium nitrite.

Statistical data analysis

Values are expressed as mean ± standard error of the mean for control and experimental samples. Statistical analysis was performed by analysis of variance, followed by the Bonferroni’s test and Student’s t-test using GraphPad software prism 5.1.

RESULTS

Effect of PfLP on carrageenan-induced paw edema in mice

The results of i.p administration of diclofenac (10 mg/kg) and PfLP (25 mg/kg and 50 mg/kg body weight) after 4-h induction on carrageenan-induced paw edema (mm³) against control group were summarized in Fig 1c. Change in leg weight of each mouse in a group (standard deviation ± mean value) also summarized in Fig 1d. On administration of PfLP shows a significant reduction in edema volume at all doses, i.e. 25 mg/kg shows 28.45±2.37% (**p<0.05) and 50 mg/kg shows 64.01±1.97% (**p<0.01) of inhibition against a control group. The standard drug diclofenac shows a significant reduction in edema of 68.87±3.21% (**p<0.01) at 10 mg/kg following its administration against to the untreated control group.

Examination of paw tissue supports the anti-inflammatory activity of PfLP effectively inhibits neutrophils accumulation in the inflamed area. A large number of neutrophils were accumulated in inflammatory tissue area when compared with the diclofenac-treated and PfLP-treated mice groups as shown in Fig 1a.

Effect of PfLP on MPO activity in carrageenan-injected mice

Results of i.p administration of PfLP on MPO activity (Abs at 450 nm) were evaluated that in carrageenan-injected mice hind paw tissue lysate is shown in Fig 2a. PfLP gradually decreases the MPO activity in PfLP-treated animals at the two different does, i.e. 25 mg/kg (0.721 OD; **p<0.01) and 50 mg/kg (0.411 OD; ***p<0.001) against a control group (0.925 OD), whereas diclofenac shows significant decreases in the MPO activity at 10 mg/kg (0.324-OD; ***p<0.001).

Effect of PfLP on NOx levels in carrageenan-injected mice

Effect of PfLP on NO production in treated and control group mice were determined by estimate the nitrite concentration (µM) using Griess reagent was summarized in Fig 2b. On treatment in a dose-dependent study with PfLP-treated group mice shows significant decreases in nitrite level up to 27.1±0.53 (µM) (**p<0.05) and 17.5±0.978 (µM) (**p<0.001) at 25 mg/kg and 50 mg/kg, respectively, compared to the untreated control mice of 36±2.01 (µM). Significant reduction of nitrite concentration is directly proportional to the NOx concentration during inflammation site. Whereas, mice received standard drug diclofenac (10 mg/kg) show a drastic reduction in nitrite level up to 12±1.45 (µM) (**p<0.001) level.

DISCUSSION

Inflammatory reactions, characterized by various cellular and vascular process, have an important role in the activation of lipoxigenase (LOX) and cyclooxygenase enzyme (COX) resulting in the release of inflammatory molecules such as prostaglandins and leukotrienes derivatives. During tissue damage/vascular inflammation, the number of events that taking place to cause inflammation such as migration of leukocytes, production of pro-inflammatory cytokines such as interleukins, NO, tumor necrosis factor-α, NF-kB transcription, tissue-specific expression of LOX, and COX cascades leads to the production of prostaglandins and leukotrienes [18]. Developing novel non-toxic therapeutics are one of the challenges ahead in the field of inflammation.
Our previous study exhibits promising lectin activity in phleum exudates of *Praecitrullus fistulosus* [21]. With the continuation of our work, we elucidate the role of lectin during inflammatory condition. On pre-treatment of PfLP before the subplantar injection of carrageenan exhibits promising anti-inflammatory activity in a dose-dependent manner. The decrease in the paw volume and leg weight supports our findings. Previous studies showed that lectin from plants competitively binds the glycosylated selectin on the extracellular membrane of leukocytes and endothelial cells and thereby it reduces the rolling and adhesion of neutrophils on the endothelium region [22]. From the study suggested that the lectins inhibit the vascular inflammation mediated by adhesion of immune cells [23]. Extravasations of neutrophils or migration of neutrophils into the inflamed area lead to cause activation of tissue-specific hydrolytic enzymes [24]. Histopathology studies of both treated and control mice suggested that a decrease in the accumulation of neutrophils in the inflamed region supports the anti-inflammatory activity of PfLP. Increase in MPO activity is considered as a hallmark of cell infiltration (mainly neutrophils) in the inflamed tissue region. MPO enzyme abundantly found in neutrophils and it low amount in macrophages and monocytes [25]. Neutrophils concentration in the inflamed tissue area was measured by estimating the MPO activity. Histopathology studies and MPO estimation in tissue area support the anti-inflammatory activity of PfLP.

NO is a free radical produced in mammalian cells, plays an important role in cell survival and death, and exhibits a pro-inflammatory effect in various types of cells [7]. NO causes a detrimental effect on normal cells which may act as toxic. Harmful effects of NO depend on the NO concentration, combination, and formation of toxic derivatives, pathophysiological condition, and responses of target tissue/cell [26,27]. Therefore, inhibition of NO production during the inflammatory process may be the potential therapeutic implications. In this study, we evaluate the NO concentration in the hind paw tissue in both control and treated mice, as shown in Fig. 2b. PfLP significantly inhibits the NO production in the carrageenan-injected mice at the dose of 50 mg/kg body weight compared to the control mice, in the late phase of the inflammation. This shows the anti-inflammatory effect of PfLP. From the above studies, we conclude that lectin may have the potential ability to inhibit or alter the inflammation responses by involved in the various pathways.

CONCLUSION

This work summarizes the anti-inflammatory activity of a PfLP from *Praecitrullus fistulosus* that is predominantly attributable inhibition of acute inflammation in a paw edema model. The study also suggested that PfLP may involve the inhibition of NO production by altering the NO enzyme activity, thereby it reduces the inflammatory associated pain. However, it has been suggested that further molecular mechanism needs to explore the involvement of PfLP in reducing inflammation, and it may consider as a promising candidate in the development of new therapeutics against inflammation-associated diseases.

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AUTHORS’ CONTRIBUTION

Sharada AC designed the study and Madhu CS performed the experimental work.

CONFLICTS OF INTEREST

The authors declare that there are no conflicts of interest.

REFERENCES


