AMELIORATING EFFECT OF LAWSONE AND ALPHA-PINENE ON L-ARGININE INDUCED ACUTE PANCREATITIS IN RATS

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ABSTRACT

Objective: This study is intended to study the protective effect of Lawsone and Alpha-pinene on L-Arginine induced acute pancreatitis in rats.

Method: In this study lawsone and alpha-pinene against L-arginine induced acute pancreatitis was determined at 24 h by determination of serum levels of amylase, lipase and proinflammatory cytokines [tumor necrosis factor (TNF)-α, C-reactive proteins and interleukin (IL)], pancreatic myeloperoxidase (MPO) activity, lipid peroxidation (thiobarbituric acid reactive substances (TBARS)), nitrate/nitrite levels, and the wet weight/body weight ratio.

Result: Lawsone, alpha-pinene and methylprednisolone treatments significantly attenuated the L-arginine-induced increases in pancreatic wet weight/body weight ratio, and decreased the serum levels of amylase and lipase, and TNF-α and IL-6 and significantly lowered pancreatic levels of MPO, TBARS, and nitrate/nitrite. The histopathological findings further proved the amelioration of pancreatic injury by lawsone and alpha-pinene. Further it also proved anti-inflammatory and antioxidant agent property of lawsone and Alpha-pinene.

Conclusion: This study had proved ameliorating effect of Lawsone and alpha-pinene on L-Arginine induced acute pancreatitis in rats.

Keywords: Acute pancreatitis, Anti-inflammatory, Cytokines, L-arginine, Lawsone, Pinene

INTRODUCTION

Inflammation of pancreatic gland called pancreatitis (AP) may leads to sever complication and high mortality without treatment. The pathogenesis is not fully understood, however the leukocyte activation, microcirculatory disturbances and oxidative stress are the major constituents of AP. This is characterized by activation of widespread inflammatory cell infiltration, leukocyte and digestive proteases. Reactive oxygen, nitrogen species and various kinds of inflammatory mediators are released in inflammatory process. Previously it was reported that several factors are responsible for the AP, like alcohol, gallstones, hereditary pancreatitis, hypercalcemia, hyperlipidemia, malnutrition, abdominal trauma, penetrating ulcers, malignancy, drugs like steroids, sulfonamides, furosemide, thiazides, infections like mumps, Coxsackie Virus, Mycoplasma Pneumoniae, Aasariv, Clostrhitis, and structural abnormalities like choledochocele and pancreas divisum[1,2]. Repeated attacks of acute pancreatitis have the potential to develop into chronic pancreatitis or pancreatic cancer characterized by fibrosis and loss of acinar cell function [1.2]. No specific treatment is available to treat AP. Many therapies and medical management is aimed to control the sign and symptoms of AP, using steroids, analgesics and anti-inflammatory agents. The use of the synthetic and semi-synthetic treatment has various kinds of drawbacks like photosensitivity skin reactions, intoxication and addiction. Apart from this, these compounds are very expensive and not reliable. Hence, there is need to explore potential antioxidant and anti-inflammatory agents available from natural sources, which are cost effective and have several advantages than the synthetic and semi-synthetic compounds. Anti-oxidant, anti-inflammatory and anti-carcinogenic activities[3,4] of alpha-pinene and anti-oxidant, anti-inflammatory and anti-carcinogenic activities[5,6] of lawsone have been reported. The alpha-pinene has shown to inhibit nuclear translocation of Nuclear Factor Kappa B [4] and lawsone has haematotoxic properties, leading to stimulation of cell proliferation; representing a sufficient explanation for a weak induction of late micronuclei[6]. Hence in this study the non-toxic dose of alpha-pinene and lawsone (100 and 200 mg/kg) were used to evaluate the potential effect to ameliorate pancreatic injury induced by L-arginine.

MATERIALS AND METHODS

Animals: Male Wistar rats (42) weighing 180-200 g obtained from Mahaveer Enterprises, Hyderabad were maintained at a constant room temperature (23±2 °C) with 12:12 h light-dark cycles and free access to water and standard laboratory chow. The rats were randomly divided into 7 groups of 6 in each and experiments were performed after 12 h of fasting. All the experimental procedures were carried out in accordance with Committee for the Purpose of Control and Supervision of Experiments on Animals (CPCSEA) guidelines. The study was reviewed and approved by the Institutional Animal Ethics Committee (320/CPCSEA dated 03-01-2001), G.Pulla Reddy College of Pharmacy, Hyderabad, India.

Chemicals: L-arginine (Sigma Aldrich Co Pvt Ltd, USA), lawsone (Sigma Aldrich Co Pvt Ltd, Japan), alpha-Pinene (Destilaciones Bordas-Chinchurreta SA), hexa-decyl-tri-methyl-ammonium bromide (HETAB) (Sigma Aldrich Co Pvt Ltd, Switzerland), o-dianisidine dihydrochloride, thiobarbituric acid (TBA) (Sigma Aldrich Co Pvt Ltd, Germany), Griess reagent (Sigma Aldrich Co Pvt Ltd, Germany) and vanadum trichloride (Sigma Aldrich Co Pvt Ltd, USA) were procured from Sigma Aldrich Chemical Co. All other chemicals and reagents were of highest commercial grade available locally.

L-arginine, powder: prepared as a solution by dissolving in 0.9% saline to a final concentration of 500 mg/mL and the pH was adjusted to 7 with 5 N HCl.

Lawsone was prepared as a solution by dissolving in 3% tween 80 and 0.9% saline to a final concentration of 100 and 200 mg/mL and the pH was adjusted to 7 with 0.1 N NaOH.

Pinene was prepared as a solution by dissolving in 3% tween 80 and 0.9% saline to a final concentration of 100 and 200 mg/mL and the pH was adjusted to 7 with 0.1 N NaOH.

L-arginine-induced pancreatitis model—Acute pancreatitis was induced in five groups of rats by two intraperitoneal (ip) injections of L-arginine (2.5 g/kg, 1 h apart). One hour following the last injection of L-arginine, the rats were treated orally as follows: Gr. 1 received the vehicle (3% Tween 80) of lawsone (vehicle control); Grs 2 and 3 were treated with lawsone (100 and 200 mg/kg, respectively), Gr 4 and 5 were treated with pinene (100 and 200 mg/kg, respectively), Gr 6 acted as positive control and received methylprednisolone (30 mg/kg), all in a volume of 10 mL/kg and Gr. 7, received saline (0.9%, NaCl, ip) in place of L-arginine and served as a normal control. After 24 h of the last
injection of L-arginine or saline, a midline laparotomy was performed. Rats under ether anaesthesia and blood samples were collected from the inferior vena cava, the rats were then exsanguinated, the whole pancreas was quickly removed and stored at -70°C until use. The pancreatic weight/body weight ratio was evaluated as an estimate of the degree of pancreatic edema (mg/g).

Macroscopic evaluation

Pancreas weight/body weight ratio—The pancreas was removed immediately after the blood collection, trimmed free of fat and weighed. The pancreatic weight/body weight ratio (mg/g) was calculated for each animal, to estimate the level of pancreatic edema.

Serum analysis—For serum analysis, blood samples were centrifuged at 3000 g at 4°C for 10 min. The serum amylase and lipase were determined by routine colorimetric methods using the commercial kits for amylase (Randox diagnostics), lipase (Accurex diagnostics), and C-reactive protein and interleukin-6 and expressed as U/l[7].

Statistical analysis—Statistical analysis was performed by one way ANOVA followed by Newman Keuls as post hoc test using GraphPad Prism 5. Values were presented as mean ± SE. The difference was considered to be statistically significant when P < 0.05.

RESULTS

Serum biochemical parameters and pancreatic edema—Induction of pancreatitis resulted in significant raise in the serum amylase, lipase and pancreatic edema. Treatment with lawson and pinene (100 and 200 mg/kg) dose dependently decreased the serum amylase, lipase and pancreatic edema (Table 1).

Pancreatic, lung, liver and kidney MPO and total protein—Induction of pancreatitis resulted in significantly increased the pancreatic MPO and decreased the pancreatic total protein levels. Treatment with lawson and pinene (100 and 200 mg/kg) dose dependently reversed the change in pancreatic MPO and total protein levels (Table 1).

Pancreatic, lung, liver and kidney MDA, nitrate/nitrite, GSH and antioxidant enzymes catalase and SOD—Induction of pancreatitis resulted in a significant raise in MDA, nitrate/nitrite, catalase and SOD and decline in GSH levels. Treatment with lawson and pinene (100 and 200 mg/kg) dose dependently reversed the change in MDA, nitrate/nitrite, catalase, SOD and GSH levels (Table 1).

Assessment of interleukins, TNF-α and C-reactive protein—Induction of pancreatitis resulted in a significant raise in interleukins, TNF-α and C-reactive protein. Treatment with lawson and pinene (100 and 200 mg/kg) dose dependently decreased the interleukins and C-reactive protein (Table 2).

Pancreatic histology—Histological examination of normal control group (saline treated) showed normal architecture and absence of edema, neutrophil infiltration, hemorrhage and necrosis (Fig 1). Whereas, pancreatic sections of disease control group showed extensive tissue damage characterized by acinar cell degeneration, necrosis, edema, mononuclear cell infiltration, hemorrhage and thus received significantly higher scores. Treatment with lawson (100 and 200 mg/kg), pine(e 100 and 200 mg/kg) and methyl prednisolone (30 mg/kg) ameliorated the inflammation, edema and more significantly acinar cell degeneration and necrosis and protected the pancreas from L. w. induced damage. Treatment with lawson dose dependently decreased the total pathological scores compared to disease control group.

Table 1: Effect of alpha-pinene and Lawsonia on pancreatic weight, total body weight, serum amylase, serum lipase, total nitrate, total protein, MDA, MPO and SOD after L-arginine induced acute pancreatitis.

<table>
<thead>
<tr>
<th>Parameter/Groups</th>
<th>N.C</th>
<th>D.C</th>
<th>STD</th>
<th>LW 100</th>
<th>LW 200</th>
<th>PIN 100</th>
<th>PIN 200</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pancreas weight</td>
<td>870.3±15.36</td>
<td>1015±19.0*</td>
<td>911.3±18.93a</td>
<td>843.3±46.33a</td>
<td>783.8±21.37a</td>
<td>862.8±58.67a</td>
<td>896.8±10.17a</td>
</tr>
<tr>
<td>Total body wt</td>
<td>187.7±4.889</td>
<td>202±2.36α</td>
<td>191.2±3.62α</td>
<td>180.7±5.57α</td>
<td>185.9±3.68β</td>
<td>188.9±8.75α</td>
<td>174±1.87</td>
</tr>
<tr>
<td>Pancreatic Ind (x10-3)</td>
<td>4.55±2.17</td>
<td>5.28±1.84α</td>
<td>4.67±1.57α</td>
<td>4.86±1.29β</td>
<td>4.16±1.22γ</td>
<td>4.58±1.28γ</td>
<td>4.99±1.24</td>
</tr>
<tr>
<td>Serum Amylase</td>
<td>2000±85.63</td>
<td>767±439α</td>
<td>331±10.8α</td>
<td>273±140.6α</td>
<td>317±67.8α</td>
<td>293±527.9a</td>
<td>320±46 a</td>
</tr>
<tr>
<td>Serum Lipase</td>
<td>191.7±4.014</td>
<td>566.7±30.84α</td>
<td>23.6±14.06α</td>
<td>260±73.03α</td>
<td>233±27.33a</td>
<td>250±24.49a</td>
<td></td>
</tr>
<tr>
<td>Total Nitrate</td>
<td>11.87±2.37</td>
<td>16.0±14.62α</td>
<td>7.06±0.93α</td>
<td>10.13±1.31α</td>
<td>8.5±0.77α</td>
<td>8.5±0.76α</td>
<td>7.05±0.83α</td>
</tr>
<tr>
<td>Total Protein</td>
<td>0.73±0.032</td>
<td>0.35±0.037α</td>
<td>0.91±0.067α</td>
<td>0.73±0.02α</td>
<td>0.82±0.05α</td>
<td>0.73±0.02α</td>
<td>0.82±0.05α</td>
</tr>
<tr>
<td>Kidney MDA</td>
<td>0.47±0.011</td>
<td>0.35±0.04α</td>
<td>0.71±0.14α</td>
<td>0.51±0.17α</td>
<td>0.64±0.09γ</td>
<td>0.10±0.04β</td>
<td>0.10±0.06α</td>
</tr>
<tr>
<td>Liver MDA</td>
<td>0.48±0.010</td>
<td>0.28±0.04α</td>
<td>0.67±0.03α</td>
<td>0.48±0.03β</td>
<td>0.59±0.04γ</td>
<td>0.08±0.03γ</td>
<td>0.08±0.04β</td>
</tr>
<tr>
<td>Lung MDA</td>
<td>0.50±0.020</td>
<td>0.28±0.04α</td>
<td>0.67±0.03α</td>
<td>0.51±0.02γ</td>
<td>0.60±0.04β</td>
<td>0.08±0.03γ</td>
<td>0.10±0.04β</td>
</tr>
<tr>
<td>Pancreatic MDA</td>
<td>1.62±0.661</td>
<td>1.04±0.4267α</td>
<td>0.39±0.79α</td>
<td>0.17±0.69γ</td>
<td>0.12±0.49β</td>
<td>0.75±0.71α</td>
<td>0.35±0.58β</td>
</tr>
<tr>
<td>Pancreas MPO</td>
<td>4.75±2.1</td>
<td>3.28±4.7γ</td>
<td>6.2±2.2α</td>
<td>1.72±3.7α</td>
<td>11.2±3.7α</td>
<td>19.2 ± 3.7α</td>
<td>14.2±3.7α</td>
</tr>
<tr>
<td>Liver MPO</td>
<td>7.7±9.042</td>
<td>38.9±2.259α</td>
<td>10.6±2.51α</td>
<td>24.9±6.03α</td>
<td>15.0±2.84α</td>
<td>26.5±4.83α</td>
<td>16.9±2.83α</td>
</tr>
<tr>
<td>Kidney MPO</td>
<td>7.88±5.419</td>
<td>31.7±2.22α</td>
<td>11.1±1.93α</td>
<td>26.1±7.26α</td>
<td>16.9±2.15α</td>
<td>28.5±4.32α</td>
<td>17.3±3.38α</td>
</tr>
<tr>
<td>Kidney MDA</td>
<td>3.55±0.123</td>
<td>32.5±9.03α</td>
<td>6.93±1.04α</td>
<td>13.3±1.67α</td>
<td>11.6±1.48α</td>
<td>21.9±2.94α</td>
<td>15.2±1.48α</td>
</tr>
<tr>
<td>Kidney Catalse</td>
<td>2.33±0.95</td>
<td>1.50±0.64α</td>
<td>2.61±1.08α</td>
<td>1.89±0.77α</td>
<td>1.89±0.77α</td>
<td>1.63±0.46</td>
<td>1.17±0.47</td>
</tr>
<tr>
<td>Liver Catalse</td>
<td>0.83±0.34</td>
<td>1.01±0.415α</td>
<td>1.97±0.80β</td>
<td>1.52±0.62γ</td>
<td>1.78±0.72β</td>
<td>1.32±0.54β</td>
<td>1.63±0.66</td>
</tr>
<tr>
<td>Liver SOD</td>
<td>1.04±0.42</td>
<td>1.03±0.42α</td>
<td>3.44±1.4α</td>
<td>2.06±0.84α</td>
<td>3.83±1.35α</td>
<td>1.04±0.42</td>
<td>2.25±0.91α</td>
</tr>
<tr>
<td>Pancreases SOD</td>
<td>2.09±0.85</td>
<td>1.51±0.61α</td>
<td>1.42±0.5α</td>
<td>1.32±0.54</td>
<td>1.81±0.74</td>
<td>1.5±0.61α</td>
<td>1.04±0.42</td>
</tr>
</tbody>
</table>

*p < 0.0001 when compared with normal control, +p 0.001, +p 0.001, +p 0.01 when compared with disease control group. LW1: Lawson 100, LW2: Lawson 200, PI 100: pinene 100 and PI200: pinene 200 mg/kg. Values are mean ± SEM from 6 animals in each group.
Table 2: Effect of alpha-pinene and Lawsone on IL-6, TNF-α and CRP after L-arginine induced acute pancreatitis.

<table>
<thead>
<tr>
<th>Parameter/Groups</th>
<th>NC</th>
<th>DC</th>
<th>LW1</th>
<th>LW2</th>
<th>PIN1</th>
<th>PIN2</th>
</tr>
</thead>
<tbody>
<tr>
<td>IL-6</td>
<td>29.3±1.569</td>
<td>90.48±1.689*</td>
<td>62.42±1.358γ</td>
<td>33.57±1.182γ</td>
<td>66.9±1.131</td>
<td>35.35±2.464γ</td>
</tr>
<tr>
<td>TNF-α</td>
<td>19.3±1.541</td>
<td>26.32±3.036</td>
<td>21.58±0.9711</td>
<td>18.57±1.316</td>
<td>21.9±1.987</td>
<td>17.35±1.383</td>
</tr>
<tr>
<td>CRP</td>
<td>415.3±7.762</td>
<td>1640.3±119*</td>
<td>861±101.7γ</td>
<td>546.8±9.874γ</td>
<td>743±63</td>
<td>562.8±17.44γ</td>
</tr>
</tbody>
</table>

N.C- Normal Control, D.C- Disease Control, LW 100- Lawsone 100 mg/kg, LW 200- Lawson 200 mg/kg, PI 100: pinene 100 and PI200: pinene 200 mg/kg. *P < 0.0001 when compared with normal control, γP < 0.01 when compared with disease control group.

Histopathology

Fig. 1—Effect of lawsone on pancreatic histopathological changes after L-arginine induced acute pancreatitis [(a) normal control, (b) standard control, (c) disease control 24 h hemorrhage, (d) disease control 24 h edema and necrosis, (e) lawsone 100 mg/kg, ip, 24 h, (f) pinene 200 mg/kg, ip, 24 h, (g) pinene 100 mg/kg, ip, 24 h, (h) pinene 200 mg/kg, ip, 24 h] (H&E ×200)
DISCUSSION

The present study demonstrated that treatment with lawsone (200 mg/kg) and pineine (200 mg/kg) efficiently reduced the severity of L-arginine induced acute pancreatitis in rats. In consistent with previous reports[2,19,20], in the present study administration of L-arginine significantly developed the acute pancreatitis characterized by raised levels of serum amylase, lipase and acinar cell necrosis. Serum amylase and lipase levels are the important diagnostic markers for acute pancreatitis. They usually rise within 4.8 hours of the initial attack, peaks at 24 hours[4,19,21]. Similarly, in accordance with previous reports, in the present study induction of pancreatitis significantly increased the serum amylase and lipase levels at 24 hours. Treatment with lawsone decreased the serum amylase and lipase levels, indicates protective effect of lawsone at early stage of the disease progression. In consistent with previous reports[34,19], in the present study induction of pancreatitis significantly increased the pancreatic MPO, MDA, nitrite, catalase and SOD and decreased the GSH levels. MPO, a marker of neutrophil infiltration is an enzyme found in neutrophils and its activity is linearly related to infiltration of neutrophils[22,23]. In agreement with previous reports[3,24], in the presence study induction of pancreatitis with L-arginine increased the pancreatic MPO levels. Inhibition of the neutrophils infiltration can attenuate the paracutaneous injury[25]. Treatment with lawsone significantly decreased the pancreatic MPO levels probably due to its anti-inflammatory action. MDA, a marker of lipid peroxidation was elevated in Larginine treated rats. Lipid peroxidation is a process mediated by free radicals, which results in impairment of the membrane functional and structural integrity[4,20,26]as a consequence of oxidative deterioration of polyunsaturated fatty acids of cell membrane. It could be attributed to the accumulation of free radicals proposed to be generated by L-arginine. The change in levels of catalase and SOD remains controversial. A few studies[26] reported the fall in these enzyme levels at 24 hours and few studies[20] reported asraised levels of these enzymes. In consistent with previous reports[2], in the present study significant increase in SOD and catalase level was observed. It indicates that oxidative stress caused by L-arginine may up-regulate the activity of antioxidant enzymes to facilitate rapid removal of accumulated reactive oxygen and nitrogen species[20]. It is well known that GSH is found to be decreased in L-arginine treated rats indicating enhanced oxidative stress as the disease progresses[20]. The role of NO in the initiation and progression of acute pancreatitis remains controversial[27]. Some studies[28,29,30][31] reported that NO increase the pancreatic blood flow and/or secretion in response to endothelium derived NO and ameliorates the pancreatic dysfunction, whereas others[32,33,34] suggested that NO aggravates pancreatic oxidative stress and damage. In agreement with previous reports[34], in the present study significant increase in NO and pancreatic edema was observed in L-arginine received rats. Previous investigations[34] demonstrated that, administration of excess L-arginine could induce iNOS activity and increase the NO levels in pancreas. The raised levels of NO can increase vascular/micro capillary permeability and may contribute to the pancreatic edema and acinar cell damage[34]. Treatment with lawsone significantly restored the pancreatic MDA, nitrite, edema, catalase, SOD and GSH in L-arginine received rats. Passaglia[35] stated that acinar cells are the protein factory of the body. In acute pancreatitis, catabolism of proteins could increase up to 80%. Consequently, a sharp decline in protein content was observed in pancreas. In consistent with previous reports[4,36,37,38] pancreatic total protein content, a marker of the tissue damage was found to decrease in L-arginine received rats, this was also corrected from previous results. Treatment with lawsone significantly increased the total protein content. It is well known that the extent of pancreatic tissue damage in acute pancreatitis correlates with the levels of inflammatory mediators and free radicals. In agreement with previous reports[4,24,21,39], in the present study, histopathological assessments revealed that, induction of pancreatitis resulted in pancreatic damage characterized by acinar cell necrosis, mono nuclear cell infiltration, edema and hemorrhage. Treatment with lawsone protected the pancreas from L-arginine induced injury. In conclusion, the present study suggests that treatment with lawsone significantly ameliorated the severity of L-arginine induced pancreatitis by reducing the neutrophil infiltration and oxidative stress markers and this effect may be due to antioxidant and anti-inflammatory properties of the lawsone.

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REFERENCE


