Curcumin has Curative Effect on Isoniazid-Rifampicin Induced Hepatotoxicity

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ABSTRACT

Background: Anti-tuberculosis drugs-induced hepatotoxicity is associated with oxidative stress. Curcumin is a powerful antioxidant and has been found to protect the liver from the damaging effects of oxidative stress. The study aimed to assess the curative and protective effects of Curcumin against hepatotoxicity induced by anti TB- drugs (Isoniazid-Rifampicin) by using an experimental model of Albino rabbits.

Methods: Albino rabbits (n=24) were divided into four groups. Group A was the control group. Group B administered Isoniazid (INH) (50 mg/kg/d) and Rifampicin (RMP) (100 mg/kg) alone, Group C received both INH+RMP and Curcumin (60 mg/kg) before and during induction. Group D received INH (50 mg/kg/d) and RMP (100 mg/kg/d) for 7 days, followed by 7 days of Curcumin (60 mg/kg/d). Biochemical testing and liver morphological histopathology was done for all groups. All values were recorded in mean ± standard deviation.

Results: Anti-Tuberculosis drugs increased Alanine Transaminase (ALT), Aspartate Transaminase level (AST), Alkaline Phosphatase level (ALP), Total Bilirubin, and Albumin 62.0±2.5, 172.5±1.0, 128±1.5, 0.80±0.05, 5.0±0.5 respectively and decreased Total protein levels (2.05±1.0). Whereas, Curcumin lowered liver enzymes 37.0±2.8, 126.12±1.5, 90.5±1.0, 0.40±0.01, 3.50±0.5 respectively, and increased levels of total protein (5.00±0.5). Group A exhibited normal liver morphology, whereas, Group B had ballooning degeneration, focal cell necrosis, and liver inflammation. Group C had moderate fatty liver but no centrilobular degeneration or focal cell necrosis and Group D exhibited minor liver inflammation and normal liver morphology.

Conclusion: Curcumin was found preventative and therapeutic remedy which can be used for the treatment of hepatotoxicity.

Keywords: Curcumin; Hepatotoxicity; Hepatoprotective; Isoniazid; Rifampicin.

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INTRODUCTION
Tuberculosis (TB) is an infectious and communicable illness caused by Mycobacterium that continues to endanger worldwide public health. In 2019, around 10 million individuals were diagnosed with TB, and 1.4 million died as a result of the disease. On the other hand, Isoniazid’s introduction in the 1950s offered an excellent method of TB control. It is still the first-line treatment for both active TB and tuberculosis prevention. Despite its great antibacterial activity, its usage is limited due to significant hepatotoxicity and fatal liver damage. As a result, it is critical to identify the hepatotoxicity mechanism as well as novel treatment targets and more effective techniques to prevent isoniazid-induced hepatotoxicity.

One of the most common drugs used in TB treatment includes Isoniazid (INH) and Rifampicin (RMP) which are used in combination to have an effective TB treatment but increase the risk of hepatic injury. Hepatotoxicity which is caused by TB medications can influence the hepatocytes or vasculature and affect the biliary epithelium, but the fundamental component and contributing elements which causes hepatic harm are not known. Brief and asymptomatic expanded transaminase levels can be minor and have non-dynamic harm to mitochondria, cell layers, or different structures of hepatocytes. Furthermore, different studies have been carried out to develop effective ways to reduce the hepatotoxicity of anti-TB drugs, which can be done by either using synthetic drugs or natural botanical products without interfering with their therapeutic actions.

Curcumin has been used which is a natural product. Contrarily, curcumin, a yellow substance generated and derived from the Curcuma longa plant, is a very potent antioxidant and has potent anti-inflammatory properties. According to a previous study, it was found that Curcumin had a protective effect against chronic liver failure as it helps to reduce harmful oxidative stress. Due to Curcumin’s chemical structure, it is a strong antioxidant agent that can prevent free radicals. The expected preventive or putative effects of Curcumin have also been associated with its antioxidant property. Therefore, this study aimed to assess the curative and protective effects of Curcumin against hepatotoxicity induced by anti-TB drugs in Albino rabbits.

METHODS
This experimental study was conducted at Shaheed Zulfiquar Ali Bhutto Institute of Science and Technology (SZABIST) in the faculty of Life Sciences. Initially, rabbits (n=24) were kept in polypropylene cages with stainless steel tops, in the SZABIST animal room to protect them from intense sunlight, and the cage bases were lined with paddy husk. It was ensured that the environmental conditions must be controlled. The temperature was maintained between 25-30°C, and the relative humidity at 55%-75%. While, the identification of the rabbits was done by marking their cages with their specific groups, and the rabbits corresponding to each group had been marked and the groups have been assigned randomly.

Albino rabbits of both genders were taken, which weighed an average of 1.2 kg. Rabbits were kept for acclimatization for 10 days under controlled temperature and light cycle. Normal feed and water were given for the entire period. After the acclimatization period, the rabbits were separated into 4 groups, with 6 rabbits in each group; Group A in which rabbits were given normal feed, and in Group B the rabbits were induced to hepatotoxicity by a combination of two drugs; isoniazid and rifampicin orally as per their body weight for 7 days. Group C rabbits received Curcumin for 3 days as a pretreatment, and after 3 days hepatotoxicity was induced by drug combination and Curcumin was also given throughout the induction period simultaneously to check the protective effects. Group D received rifampicin and isoniazid for initial 7 days and was then given Curcumin for 7 days to check the curative effects of Curcumin against hepatotoxicity.

Isoniazid and Rifampicin, the two medications, are administered in combination to have significantly stronger effects in the effort to treat tuberculosis. Isoniazid was set to be administered in doses of 50 mg per kilogram of rabbit weight, whereas rifampicin was set to be given in doses of 100 mg per kilogram of rabbit weight. However, 60 mg of curcumin per kilogram of the rabbits’ weight was given. A combination of INH and RMP was used to cause liver damage, these drugs were given orally for 7 days to induce hepatotoxicity. After 7 days, blood samples were collected to check whether hepatotoxicity was induced or not. A Liver Functioning Test was done to see whether the level of liver enzymes and proteins was elevated or decreased. The levels were compared with the control group.

After completion of the model, blood (2 ml) was drawn from rabbits’ lateral ear veins and the samples were collected in a yellow top vacutainer and labeled according to the groups. Xylene was applied on the rabbits’ ear for visibility of the vein. To obtain serum, blood was centrifuged for 15 minutes at 2200- 2500 RPM. The liver function test was assessed by levels of Albumin, Alanine Transaminase level (ALT), Aspartate Transaminase level (AST), Alkaline Phosphatase level (ALP), total bilirubin and total protein. The levels were compared with the control group to check the intensity of liver damage. The liver of rabbits was isolated and preserved in 10% formalin. Sections of 5 μM were prepared; Hematoxylin and eosin were used to stain slides and then they were observed under a light microscope. Statistical analysis was done by marking their cages with their specific groups and the rabbits corresponding to each group had been marked and the groups have been assigned randomly.
formalin. Sections of 5 µM were prepared; Hematoxylin and eosin were used to stain slides and then they were observed under a light microscope. Statistical Package for Social Science program (SPSS) by IBM was used to perform statistical analysis. All tests were compared with Group A and all data was expressed as mean and standard deviation (±).

RESULTS
Out of the all groups of rabbits (n=24), experimental group C had decreased average liver weights compared to group D. Figure 1 represents the average liver weight of each group. However, the protective group had a mean liver weight that was more comparable to the control group. The raised hepatic weight of group D is a clear physical sign of hepatotoxicity.

![Figure 1: Mean hepatic weight of each group.](image)

Table 1: Observational analysis of faces and urine.

<table>
<thead>
<tr>
<th>Group A (Control)</th>
<th>Stool Consistency</th>
<th>Urine Coloration</th>
</tr>
</thead>
<tbody>
<tr>
<td>Day 1</td>
<td>Round consistent shape and amount</td>
<td>Pale white, clear</td>
</tr>
<tr>
<td>Day 3</td>
<td>Round consistent shape and amount</td>
<td>Pale white, clear</td>
</tr>
<tr>
<td>Day 5</td>
<td>Round consistent shape and amount</td>
<td>Pale white, clear</td>
</tr>
<tr>
<td>Day 7</td>
<td>round consistent shape and amount</td>
<td>Pale white, clear</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Group B (Anti-Tb Drug)</th>
<th>Stool Consistency</th>
<th>Urine Coloration</th>
</tr>
</thead>
<tbody>
<tr>
<td>Day 1</td>
<td>Round consistent shape and amount</td>
<td>Pale white, clear</td>
</tr>
<tr>
<td>Day 3</td>
<td>Cecotropes, sticky</td>
<td>Pale white, turbid</td>
</tr>
<tr>
<td>Day 5</td>
<td>Sticky/slimy</td>
<td>Amber, turbid</td>
</tr>
<tr>
<td>Day 7</td>
<td>watery diarrhea</td>
<td>Red, turbid</td>
</tr>
</tbody>
</table>

During the pharmacological induction of hepatotoxicity, an observational characterization of the feces and urine of all groups was carried out. Following the induction of hepatotoxicity in Group D, another observation analysis was carried out within the allotted time frame of fourteen days. At the end of the 14th day of the curcumin treatment, the feces were more Cecotropes and sticky when compared to the feces that were produced by Group A. After seven days Group B (Anti-Tb Drug) experienced watery diarrhea and red, cloudy urine which can be an indication of hematuria. Group C, on the other hand, was characterized by its small size, mixed shape, and dry, and dark-colored feces (Table 1). While it was discovered that the color of urine was amber and possessed turbidity, whereas group A, includes urine that is transparent with a pale white color, and normal stool consistency. This demonstrates that there had been internal modifications in the gastrointestinal tract and urinary tract of rabbits in which hepatotoxicity was induced.
Following a study in Poland, fosfomycin resistance is common compared to hospitalized patients. This is in close association with a study conducted in Singapore. Their study revealed that fosfomycin resistance (7.8%) is also reported in a study by Micahopoulos et al. in 2022.

Lower levels of estrogen lead to a reduction of a short urethra and decreased level of estrogen. This can be endorsed to the truth that females have shorter urethra. This is also detected by a study on women's health in Asia. Fosfomycin resistance is around 23%. However, delineated that it is difficult to use Fosfomycin as empirical therapy. A study from the Czech Republic reported that it is difficult to use Fosfomycin as empirical therapy. A study from the Czech Republic reported that it is difficult to use Fosfomycin as empirical therapy.

Table 2: Effects of curcumin on enzyme and protein levels of the liver in isoniazid-rifampicin induced hepatic toxicity in rabbits.

<table>
<thead>
<tr>
<th>Groups</th>
<th>Stool Consistency</th>
<th>Urine Coloration</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Group C (Pre-Anti TB Drugs)</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Day 1</td>
<td>Round consistent shape and amount</td>
<td>Pale white, clear</td>
</tr>
<tr>
<td>Day 3</td>
<td>Cecotropes, sticky</td>
<td>Pale white, clear</td>
</tr>
<tr>
<td>Day 5</td>
<td>Cecotropes, sticky</td>
<td>Amber, turbid</td>
</tr>
<tr>
<td>Day 7</td>
<td>Small, mixed shape, dry and dark</td>
<td>Amber, turbid</td>
</tr>
<tr>
<td><strong>Group D (Post-Anti-TB Drugs)</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Day 1</td>
<td>Round consistent shape and amount</td>
<td>Pale white, clear</td>
</tr>
<tr>
<td>Day 3</td>
<td>Cecotropes, sticky</td>
<td>Pale white, turbid</td>
</tr>
<tr>
<td>Day 5</td>
<td>Sticky/slimy</td>
<td>Amber, turbid</td>
</tr>
<tr>
<td>Day 7</td>
<td>Watery diarrhea</td>
<td>Red, turbid</td>
</tr>
<tr>
<td>Day 9</td>
<td>Cecotropes, sticky</td>
<td>Pink, turbid</td>
</tr>
<tr>
<td>Day 11</td>
<td>Cecotropes, sticky</td>
<td>Amber, turbid</td>
</tr>
<tr>
<td>Day 14</td>
<td>Cecotropes, sticky</td>
<td>Amber, clear</td>
</tr>
</tbody>
</table>

Group B expressed that INH and RMP elevated the levels of ALP, AST, ALT, total bilirubin, Albumin (62.0±2.5, 172.5±1.0, 128±1.5, 5.0±0.5, 5.0±0.5 respectively) and decreased total protein levels (2.0±1.0) as compared to Group A. Whereas Group C receiving co-administration of Curcumin and INH-RMP and Group D receiving Curcumin after hepatotoxicity induction showed decreased levels of liver enzyme (37.0±2.8, 126.12±1.5, 90.5±1.0, 0.40±0.01, 3.50±0.5 respectively) increased levels of total protein (5.00±0.5) when compared to Group B as shown in Table 2.

In addition, Figure 2 represented histological images in which Group A showed normal liver morphology whereas Group B in which hepatotoxicity was induced showed ballooning degeneration, focal cell necrosis and liver inflammation. Group C was pre and co-treated with Curcumin indicated no centrilobular degeneration or focal cell necrosis however it only had mild fatty liver. Group D was post-treated with Curcumin which indicated mild liver inflammation and normal liver morphology.
Among different common infections, urinary tract infections are frequently detected in the community and urinary tract infection is the most common infection in patients, this fact collaborates with another study by 23.

Isolates from urinary samples, this 22 - 25 - 2020 was found in association with our study Fosfomycin resistance (14.1%) in isolates of therapy study was conducted at Rawalpindi and Islam however, delineated E. coli -

resulted in 900 fatalities for every 100,000 people. This where an anti-tuberculosis drug increases liver documented tubercles, abscesses development, beginning, its pathological and anatomical dating the isolation of consumptives before their
discrimination showed decreased levels of liver enzyme, inflammation and cell death at focal and centrilob microscopic examination of these cells indicates to produce hepatotoxicity in rabbit liver cells, and found isoniazid and rifampicin had been observed isoniazid-rifampicin drug on liver histological studies of hepatotoxicity in rabbits. J Pharmacol Toxicol Methods. 1995;34(2):109-16. Throm

- 30. doi 10.1016/j.pulmoe.2019.09.005

Hepatitis caused by isoniazid and rifampicin is linked
to the death of cells in certain regions of the liver as well as centrilobular degeneration\textsuperscript{21}. The study found isoniazid and rifampicin had been observed to produce hepatotoxicity in rabbit liver cells, and microscopic examination of these cells indicates inflammation and cell death at focal and centrilobular regions\textsuperscript{22,23}. This was following the characteristics observed in this study which validate the current study animal model. The buildup of a toxic intermediate of isoniazid and rifampicin metabolism is thought to be the cause of liver damage\textsuperscript{24}. Thrombocytopenia can be caused by any of the basic anti-tubercular medicines\textsuperscript{25}. It happens as a hematological response in the case of isoniazid. One of the indications of thrombocytopenia is blood in the rabbit’s urine or stool\textsuperscript{26}. Although thrombocytopenia is uncommon, it can occur with anti-tubercular medications such as Rifampicin and Isoniazid, which can have deadly side effects. Bilirubin is not effectively processed due to hepatotoxicity or a damaged liver\textsuperscript{27}. Thus, drastically elevated bilirubin levels in the blood can induce jaundice. A high level of bilirubin in the circulation can induce leakage and spread into adjacent tissues\textsuperscript{28-30}. In line with this, another research demonstrated comparable biochemical changes, such as elevated serum ALT levels, in rabbits who were subjected to INH-induced hepatotoxicity, indicating aberrant functioning integrity of the cells that make up the liver\textsuperscript{29}.

In this study, Curcumin after hepatotoxicity induction showed decreased levels of liver enzyme, however, curcumin and its derivatives have been shown in studies to have antimycobacterial and antitubercular activity via influencing the human immune response throughout infection. While recent trials on curcuminoids have shown that they are safe for human intake, an examination of the derivatives is required. The goal of this study was to assess the role in the treatment of curcumin and its compounds in tuberculosis. As a result, the current study provides evidence that curcumin might be regarded as a potent antitubercular agent and may be exploited to develop innovative medications for TB protection and therapy. While rising evidence supports the use of curcumin and its derivatives for tuberculosis therapy, further clinical and preclinical research is needed before recommending curcumin formulations in public health\textsuperscript{30}.

**CONCLUSION**

This study found curcumin as a protective and curative medication for preventing liver failure by tuberculosis medications-induced hepatotoxicity. Therefore, curcumin is a useful anti-oxidant in the treatment of hepatotoxicity.

**ACKNOWLEDGEMENTS**

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**CONFLICT OF INTEREST**

There was no conflict of interest.

**ETHICS APPROVAL**

The Ethics Reviewing Board at Shaheed Zulfikar Ali Bhutto Institute of Science and Technology approved this study, according to the declaration of Helsinki principles.

**FUNDING**

The research study was funded by the Shaheed Zulfikar Ali Bhutto Institute of Science and Technology (SZABIST).

**AUTHORS’ CONTRIBUTION**

MAK contributed to the conception and design of the study, the input of intellectual data and the drafting of the manuscript. SAH assisted in the acquisition, analysis and interpretation of data and paper drafting. SM was the supervisor of the study while AMK helped in the sample analyses for biochemistry and histopathology results. All authors approved the final version of the manuscript to be published.

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