PROTECTION OF *TERMINALIA BEleriCA ROXB*. AGAINST IRON OVERLOAD INDUCED LIVER TOXICITY: AN ACCOUNT OF ITS REDUCING AND IRON CHELATING CAPACITY

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ABSTRACT
Excess iron deposition in the liver catalyses the production of Reactive Oxygen Species (ROS) which in turn initiate oxidative damage of protein and nucleic acids leading to several human diseases. The fruits of *Terminalia BeleriCa* Roxb. (TB) has been most commonly used not only in stimulating gastrointestinal health, but also in treatment of various hepatic disorders. The present study was aimed to evaluate the ameliorating effect of 70% methanol extract of TB (TBME) on iron overload induced liver injury, along with its *in vitro* iron chelating and DNA protection studies. Iron overload was induced by intraperitoneal administration of iron-dextran into mice. The biochemical markers of hepatic damage, liver iron, protein carbonyl and hydroxyproline content were measured in response to the oral administration of TBME. The reductive release of ferritin iron by TBME was further studied. The extract exhibited significant iron chelation with IC$_{50}$ of 27.70 ± 2.27 µg mL$^{-1}$ and considerable DNA protection with [P]$_{50}$ of 1.30 ± 0.01 µg mL$^{-1}$. Treatment with different doses (50, 100 and 200 kg body weight) of TBME showed dose dependent reduction in liver iron, lipid peroxidation, protein oxidation, liver fibrosis, serum enzymes and ferritin. The antioxidant enzymes levels were enhanced and the reductive release of ferritin iron increased significantly with gradually increasing concentrations of TBME. From the present results, it appears possible to support that TBME represents beneficial effects on iron overload mediated liver toxicity and hence possibly useful as iron chelating drug for iron overload diseases.

Keywords: Ferritin, Iron Overload, Lipid Peroxidation, Liver, Protein Carbonyl, *Terminalia BeleriCa*

1. INTRODUCTION
The liver is considered to be one of the most vital organs functioning as a centre of nutrient metabolism and waste metabolite excretion. Hepatic damage is associated with distortion of these metabolic functions (Wolf, 1999) and sometimes, resulting in the serious health problems. Hepatotoxicity is the most common finding in patients with iron overloading as liver is mainly responsible for taking up and storing excessive amount of iron (Papanastasiou *et al.*, 2000). Iron, the most abundant transition metal in the body, is required by all mammalian cells for growth and survival. But, sometimes, extreme iron deposition increases oxidative stress in the liver and will lead to further injuries such as hepatocellular necrosis (Olynyk *et al.*, 1995), inflammation (Deugnier *et al.*, 1992), fibrosis (Gardi *et al.*, 2002; Arezzini *et al.*, 2003) and even to cancer (Niederau *et al.*, 1985). In all iron overload induced diseases, iron removal by iron chelation therapy is an effective lifesaving strategy. The iron chelating agents currently available for clinical use are deferoxamine, 1,2-dimethyl-3-hydroxypyrid-4-one (deferiprone, L1) and deferasirox. However, such compounds are inadequate and exert several side effects and disadvantages (Al-Refaie *et al.*, 1992; Kontogiorghe, 2003). In view of severe undesirable side effects of synthetic agents, there is growing focus to develop more effective and safe
2. MATERIALS AND METHODS

2.1. Chemicals

Iron-dextran and guanidine hydrochloride were purchased from Sigma-Aldrich, USA. Trichloroacetic Acid (TCA), Nitro Blue Tetrachloride (NBT), reduced Nicotinamide Adenine Dinucleotide (NADH), Phenazine Methosulfate (PMS), ferrozine, glutathione reduced, bathophenanthroline sulfonate disodium salt, Thiobarbituric Acid (TBA), L-Ascorbic acid and 5,5'-Dithiobis-2-Nitrobenzoic Acid (DTNB) were obtained from Sisco Research Laboratories Pvt. Ltd, Mumbai, India. Hydrogen peroxide, KCl, ammonium iron (II) sulfate hexahydrate [(NH₄)₂Fe(SO₄)₂·6H₂O], 1-chloro-2,4-dinitrobenzene (CDNB), chloramine-T, hydroxylamine hydrochloride, Dimethyl-4-aminobenzaldehyde and 2,4-Dinitrophenylhydrazine (DNPH) were obtained from Merck, Mumbai, India. Ferritin was purchased from MP Biomedicals, USA. Streptomycin sulphate and N-(2-Hydroxyethyl)piperazine-N’-2-ethanesulphonic acid (HEPES buffer) were obtained from HiMedia Laboratories Pvt. Ltd, Mumbai, India. The standard oral iron chelating drug, desirox (parent group deferasirox), was obtained from Cipla Ltd., Kolkata, India. Desferal (deferoxamine mesylate USP), was obtained from Novartis, USA.

2.2. Plant Material

The fruits of Terminalia Belerica (TB) were collected from the Bankura district of West Bengal, India. It was identified and authenticated through the Central Research Institute (Ayurveda), Kolkata, India and a voucher specimen (CRHS 114/08) was submitted there.

2.3. Animals

Male Swiss albino mice (20±2 g) were purchased from Chittaranjan National Cancer Institute (CNCI), Kolkata, India and were maintained under a constant 12 h dark/light cycle at an environmental temperature of 22±2°C. The animals were provided with normal laboratory pellet diet and water ad libitum. All experiments were performed after obtaining approval from the institutional animal ethics committee and care of the animals was taken as per the guidelines of the Committee for the Purpose of Control and Supervision of Experiments on Animals (CPCSEA), Ministry of Environment and Forest, Government of India.

2.4. Extract Preparation

The fruits of TB were dried at room temperature for 7 days, finely powdered and used for extraction. The powder (100 g) was mixed with 500 mL methanol: Water (7:3) using a magnetic stirrer for 15 h, then the mixture was centrifuged at 2850 g and the supernatant was decanted. The process was repeated again with the precipitated pellet. The supernatants were collected, concentrated in a rotary evaporator and lyophilized. The dried extract was stored at -20°C until use. A 46.8% yield of the extract was obtained.

drugs (Zhang et al., 2006; Pardo-Andreau et al., 2008) which may raise the therapeutic benefits for patients.

Phenolics and flavonoids are plant metabolites widely spread throughout the plant kingdom. It has been found that most of them are effective antioxidants (Rice-Evans et al., 1996; Lopez et al., 2003) and iron chelation is very important part of their antioxidant activity (Cook and Samman, 1996). Phenolics and flavonoids therefore, could have a double, synergistic action, making them effective antioxidants, particularly active in pathological situations involving iron overload, e.g. β-thalassemia, Friedreich’s ataxia and hemochromatosis.

Terminalia Belerica (TB) (Gaertn.) Roxb. (Family-Combertaceae), popularly known as ‘Belleric myrobalan’, is a deciduous tree growing wild throughout the Indian subcontinent, Sri Lanka and south-east Asia. It is an integral part of traditional laxative formulation, ‘Triphala’ which is used for a huge variety of ailments for generations in Ayurvedic medicine (Chandra, 1996; Lopez et al., 1996; Lopez et al., 1999; Lopez et al., 2003) and iron chelation activity. TB is also reported to have hepatoprotective property of 70% methanol extract of TB (TBME) has been reported with the finding of significant amount of iron chelating activity of TBME and whether this activity along with reducing power can normalise the damage caused to liver by iron overload.
2.5. In vitro Study

2.5.1. Iron Chelation

The Fe$^{2+}$ chelating activity of TBME was evaluated according to a standard method (Haro-Vicente et al., 2006). In a HEPES buffer (20 mM, pH 7.2) medium, TBME (0-120 µg mL$^{-1}$) and positive control EDTA (0-20 µg mL$^{-1}$) were separately added to a 12.5 µM ferrous sulfate solution and 75 µM ferrozine was added to start reaction. The mixture was shaken vigorously and left standing for 20 min at room temperature and then the absorbance was taken at 562 nm. All tests were performed six times.

2.6. DNA Protection

pUC18 plasmid DNA was used for DNA protection study by TBME, according to a previously described method with minor modifications (Hermes-Lima et al., 1998). In HEPES buffer (pH 7.2, 100 mM), FeSO$_4$ solution (750 µM), TBME of varying doses (0-5 µg mL$^{-1}$), DNA (0.5 mg mL$^{-1}$) and water were added to make an initial reaction mixture. Finally, H$_2$O$_2$ solution (7.5 mM) was added to start the reaction. Desferal (0.2 mM) was used to stop the reaction after 10 min. 25 µL of each reaction mixture was loaded in 1% agarose gel. After migration, the gel was stained with ethidium bromide and visualized in a UV transilluminator. The DNA bands were quantified through densitometry, with the parameter Volume = Intensity X Area and the following formulae were used to calculate the percentage of protection:

$$\% \text{ SC} = \frac{[1.4 \times \text{SC}/(\text{OC}+(1.4 \times \text{SC}))]}{100}$$

where, SC = supercoiled; OC = open circular; 1.4 = correction factor; $\%$ protection = 100 X [(control SC-chelator SC)/(control SC-no chelator SC) - 1].

The ability of the plant extract to protect the DNA supercoil can be expressed by the concentration of sample required for 50% protection, designated as the $[P]_{50}$ value.

2.7. In Vivo Study

2.7.1. Experimental Design

Out of six groups each comprising of six mice, one group received normal saline only and served as Blank (B). The mice of other five groups were i.p injected with 100 mg kg$^{-1}$ b.w. each, of iron-dextran saline given five doses (one dose every two days). Normal saline was administered to one iron-dextran group (C) and other four groups were orally treated with 50 mg kg$^{-1}$ b.w. (S50), 100 mg kg$^{-1}$ b.w. (S100), 200 mg kg$^{-1}$ b.w. (S200) TBME and 20 mg kg$^{-1}$ b.w. Desirox (D), respectively, for three consecutive 7 day periods, started from the day after the first iron-dextran injection.

2.8. Sample Collection and Tissue Preparation

At the end of oral treatment, mice were fasted overnight before sacrifice. Blood was collected by cardiac puncture post terminal anesthetization with ethyl ether. Blood sera were separated by centrifugation and stored at -80°C until analysis. Blood cell from dissected liver was rinsed off with ice-cold saline. Half portion of the liver was cut, weighed and homogenized in 10 volume of 0.1 M phosphate buffer (pH 7.4) containing 5 mM EDTA and 0.15 M NaCl and centrifuged at 8000 g for 30 min at 4°C. Supernatant collected was used for the determination of lipid peroxidation, protein oxidation, hydroxyproline content and enzyme activities. Standard BSA graph was used to estimate the protein concentration in the homogenate. The other half of the liver samples were weighed and digested with 1:1 mixture of sulphuric acid and nitric acid to analyze the iron content.

2.9. Serum Enzymes

Alanine Amino Transferase (ALAT), Aspartate Amino Transferase (ASAT) and bilirubin in serum samples were measured using the commercial kits of Merck, Mumbai, India. Alkaline Phosphatase (ALP) was estimated using the kit supplied by Sentinel Diagnostics, Italy.

2.10. Antioxidant Enzymes

Superoxide Dismutase (SOD) was assayed by measuring the inhibition of the formation of blue colored formazan at 560 nm (Kakkar et al., 1984). Catalase (CAT) activity was measured through time course decomposition of H$_2$O$_2$ at 240 nm (Bonaventura et al., 1972). A formerly reported method was followed to determine Glutathione-S-Transferase (GST) based on the formation of GSH-CDNB conjugate (Habig et al., 1974). Reduced Glutathione (GSH) level was measured spectrophotometrically at 412 nm by a standard procedure (Eilman, 1959).

2.11. Lipid Peroxidation Products

Lipid peroxidation levels in liver homogenates were determined in terms of Thiobarbituric Acid Reactive Substances (TBARS), as an index of malondialdehyde accumulation, in accordance to a formerly reported method (Buege and Aust, 1978).
2.12. Protein Carbonyl Content

Protein carbonyl, a marker of protein oxidation, was done according to a spectrophotometric method (Reznick and Packer, 1994). Briefly, 50 µL streptomycin sulphate (10% w/v) was added to 450 µL sample homogenate and then centrifuged at 2800 g for 15 min. Then 200 µL of the supernatant was incubated with the same volume of 10 mM DNPH in 2 M HCl at room temperature for 20 min. 10% cold TCA was added to end the reaction and precipitate the proteins that were washed with ethyl acetate-ethanol mixture (1:1) to remove unreacted precipitate the proteins that were washed with ethyl acetate-ethanol mixture (1:1) to remove unreacted DNPH. The final protein pellet was dissolved in 1 mL 6 µmol/µL of guanidine hydrochloride solution and the absorbance was measured at 370 nm, using the molar extinction coefficient of DNPH, \( \varepsilon = 2.2 \times 10^4 \text{M}^{-1} \text{cm}^{-1} \).

2.13. Hydroxyproline Content

Liver fibrosis gives rise to collagen which is alternately represented by hydroxyproline content. Hydroxyproline of liver samples hydrolized in 6 M HCl was measured by Ehrlich’s solution using a standard procedure (Bergman and Loxley, 1963). A standard curve (R² = 0.9907) of 4-hydroxy-L-proline was measured and results were calculated after taking absorbances at 558 nm. The collagen content was determined by multiplying amount of total hydroxyproline content in each sample by a factor of 7.69 (Kivirikko et al., 1967). Results are expressed as milligrams of collagen per liver (wet weight).

2.14. Histopathological Study

The liver samples were excised, washed with normal saline and processed separately for histological observations. Initially, the material was fixed in 10% buffered neutral formalin for 48 h. A paraffin embedding technique was carried out and sections were taken at 50 µm thickness, stained with hematoxylin and eosin and examined microscopically for histopathological changes.

2.15. Liver Iron and Serum Ferritin

Liver iron was measured according to a formerly reported colorimetric method (Barry and Sherlock, 1971). Samples were incubated with bathophenanthroline sulfonate for 30 min at 37°C and absorbances were measured at 535 nm. Serum ferritin levels were measured using enzyme-linked immunosorbent assay kit (from Monobind Inc., USA) according to the manufacturer’s instructions.

2.16. Iron Release from Ferritin

Iron release assay was performed using ferrozine, a classic Fe²⁺-chelator, as previously described (Hynes and Coinceanainn, 2002). The reaction mixture (3 mL final volume) contained 200 µg ferritin, 500 µM ferrozine, in 50 mM pH 7.0 phosphate buffer. Time dependent release of iron in the reaction mixture on addition of different doses of TBME (100-500 µg) was measured spectrophotometrically at 560 nm for 20 min. Reaction mixture of buffer, ferritin and ferrozine, devoid of plant extract was used as the reference solution.

2.17. Statistical Analysis

All data are reported as the mean ± SD of six measurements. Statistical analysis was performed using KyPlot version 2.0 beta 15 (32 bit) and Origin professional 6.0. Comparisons among groups were made according to pair t-test. In all analyses, a p value of < 0.05 was considered significant.

3. RESULTS

3.1. In vitro study

3.1.1. Iron Chelation

The formation of violet colored Fe²⁺-ferrozine is disrupted in presence of an iron chelator. The results (Fig. 1a and b) demonstrated that the formation of Fe²⁺-ferrozine complex is inhibited dose dependently in the presence of TBME and reference compound, EDTA. The IC₅₀ values of the TBME and EDTA were 27.70 ± 2.27 µg mL⁻¹ and 1.27 ± 0.05 µg mL⁻¹, respectively. TBME inhibited 68% of the complex formation at the highest dose, i.e., 120 µg mL⁻¹.

3.2. DNA Protection

The protective effect of TBME against Fe²⁺-H₂O₂ mediated DNA breakdown was demonstrated in Fig. 2a. pUC18 supercoiled DNA was used as control (lane 1). Lane 2 comprised only of the open circular form of DNA generated by Fenton reaction. Addition of gradually increasing concentrations of TBME resulted in the restoration of DNA to the supercoiled form (lane 3-12). The results in Fig. 2b showed the dose dependant protection by TBME with a [P]₅₀ value of 1.30 ± 0.01 µg mL⁻¹.

3.3. In vivo Study

3.3.1. Serum Marker Enzymes

As shown in Table 1, increase in the levels of serum enzymes, viz., ALAT (105.92%), ASAT (94.12%), ALP (141.62%) and billirubin (268.41%), clearly signifies iron induced liver damage. Eventually, significant dose dependent decrease in the individual levels is observed in the TBME treated mice.
Fig. 1. Effect of (a) TBME and (b) standard EDTA on ferrozine-Fe$^{2+}$ complex formation. The data expressed as % inhibition of chromogen formation. The results are mean ± S.D. of six parallel measurements. ***p < 0.001 vs 0 µg/ml

3.4. Antioxidant Enzymes

The overload of iron in liver results in oxidative stress and affects the decrease in the levels of antioxidant enzymes. The reduction in SOD (87%), CAT (59%) GST (80%) and GSH (35%) levels which was decreased in group C compared to group B, was found to be elevated dose dependently after treatment with TBME (Table 2).

3.5. Lipid Peroxidation

The present result showed, 63% increment of lipid peroxidation in liver homogenates of iron injected mice than blank was significantly reduced by 7, 40 and 42% in mice fed with S50, S100 and S200 respectively (Fig. 3).

3.6. Protein Carbonyl Content

Protein carbonyl formation is an early marker for protein oxidation. A significant elevation (137%) of protein carbonyl content in iron overloaded mice was found to be arrested by 25, 51 and 66% in S50, S100 and S200 group respectively (Fig. 4) as treated with TBME.

3.7. Hydroxyproline Content

The hydroxyproline content was determined as it signifies the enhanced level of collagen content in liver fibrosis. Hydroxyproline content was observed to rise by 83% in group C than group B, as shown in Fig. 5. This upsurge of collagen content was gradually reduced by 42, 43 and 71% in TBME treated mice (S50, S100 and S200 respectively).
Fig. 2. Protection against oxidative damage to pUC18 by TBME. Picture of agarose gel of pUC18 DNA showing bands of supercoiled (SC) and open circular (OC) forms. Lanes on the gel represent: (Lane 1) control DNA (no H$_2$O$_2$ or Fe$^{2+}$); (Lane 2) reaction mixture without extract; (Lane 3-12) reaction mixture with extract of increasing concentration (0.5-5 µg/ml). **p < 0.01 and ***p < 0.001 vs 0 µg/ml

Table 1. The effect of TBME on serum marker enzymes (ALAT, ASAT and ALP) and bilirubin in iron overloaded mice

| Treatment | ALAT Unit/L  | Change (%) | ASAT Unit/L  | Change (%) | ALP Unit/L  | Change (%) | Billirubin Unit/L  | Change (%) |
|-----------|--------------|------------|--------------|------------|-------------|------------|-------------------|------------|
| B         | 19.01±3.33   | ---        | 33.64±1.57   | ---        | 122.77±8.12 | ---        | 1.18±0.12         | ---        |
| C         | 39.15±1.51$^{X1}$ | 105.92 | 65.30±3.23$^{X2}$ | 94.12 | 296.64±11.16$^{X1}$ | 141.62 | 4.36±0.21$^{X2}$ | 268.41 |
| S50       | 36.16±0.62$^{X2}$ | 90.20 | 51.58±0.34$^{X1Y1}$ | 53.34 | 285.28±16.01$^{X2}$ | 132.37 | 3.51±0.34$^{X1Y1}$ | 195.94 |
| S100      | 33.91±0.69$^{X1Y2}$ | 78.37 | 42.56±0.98$^{X2Y1}$ | 26.53 | 249.22±1.94$^{X1Y1}$ | 103.00 | 3.27±0.17$^{X1Y1}$ | 176.23 |
| S200      | 25.84±2.78$^{X2}$ | 35.92 | 35.66±1.91$^{X1Y2}$ | 6.00 | 169.46±30.16$^{X1Y1}$ | 38.03 | 2.60±0.35$^{X1Y1}$ | 119.71 |
| D         | 23.55±1.39$^{Y1}$ | 23.88 | 44.68±5.53$^{X1Y1}$ | 32.81 | 151.58±9.92$^{X1Y1}$ | 23.47 | 1.60±0.12$^{X1Y2}$ | 34.78 |

Values are mean ± SD of six observations.
X: significant difference from normal mice (B) group (X1: p ≤ 0.05; X2: p ≤ 0.01 and X3: p ≤ 0.001)
Y: significant difference from iron overloaded (C) group (Y1: p ≤ 0.05; Y2: p ≤ 0.01; Y3: p ≤ 0.001)
### Table 2. The effect of TBME on antioxidant enzymes (SOD, CAT and GST) and non-enzymatic antioxidant, GSH in iron overloaded mice

| Treatment | Unit/mg protein | Change (%) | Unit/mg protein | Change (%) | Unit/mg protein | Change (%) | µg/mg protein | Change (%) |
|-----------|-----------------|------------|-----------------|------------|-----------------|------------|--------------|------------|
| B         | 0.68 ± 0.05     | ---        | 22.2 ± 1.35     | ---        | 5.26 ± 0.64     | ---        | 0.49 ± 0.04  | ---        |
| C         | 0.09 ± 0.02     | X3         | 9.07 ± 0.64     | X3         | 1.05 ± 0.19     | X3         | 30.01        | X3         |
| S50       | 0.16 ± 0.06     | X3Y2       | 11.29 ± 0.71    | X3Y2       | 2.19 ± 0.22     | X3Y2       | 58.44        | X3        |
| S100      | 0.45 ± 0.04     | X3Y2       | 15.44 ± 0.79    | X3Y2       | 3.14 ± 0.41     | X3Y2       | 40.36        | X3        |
| S200      | 0.47 ± 0.03     | X3Y3       | 16.37 ± 0.74    | X3Y3       | 4.13 ± 0.41     | X3Y3       | 21.44        | X3        |
| D         | 0.39 ± 0.06     | X3Y3       | 18.97 ± 0.81    | X3Y3       | 3.13 ± 0.53     | X3Y3       | 40.43        | X3        |

Values are mean ± SD of six observations.

X: significant difference from normal mice (B) group (X2: p ≤ 0.01 and X3: p ≤ 0.001)

Y: significant difference from iron overloaded (C) group (Y2: p ≤ 0.01; Y3: p ≤ 0.001)

### 3.8. Histopathological Analysis

Histology of the liver sections of normal mice showed normal cell morphology with well-preserved cytoplasm, prominent nucleus and well brought out central vein (Fig. 6A). Iron-dextran control mice showed various degrees of pathological changes including hepatocellular necrosis, ballooning degeneration and loss of cellular boundaries (Fig. 6B). In contrast, the liver sections taken from TBME treated mice showed attenuation of the pathological changes and revealed marked reduction in hepatic lesions (Fig. 6C-E). Figure 6F exhibited the improved histology of liver sections taken from desirox treated group.

### 3.9. Liver Iron and Serum Ferritin Levels

Liver iron content was elevated about 151% after intraperitoneal administration injection of iron dextran. The TBME treatment (S50, S100 and S200) evidently lowered the iron content at a level of 6, 22 and 35% respectively (Fig. 7). Significant increase (182%) of serum ferritin level in iron loaded mice was substantially reduced to 58, 86 and 122% as treated with TBME dose dependently to S50, S100 S200, respectively (Fig. 8).

### 3.10. Reductive Release of Ferritin Iron and its Correlation with Reducing Power

Figure 9 showed the reductive release of ferritin iron by TBME, that was measured with a ferrous complex of ferrozine [Fe(ferrozine)₂]⁺. Control experiments without TBME produced negligible amounts of [Fe(ferrozine)₂]⁺, whereas, after dose dependant addition of TBME the [Fe(ferrozine)₂]⁺ complex formation was increased significantly with time. The correlation graph of reducing power and released ferritin iron (%) by TBME was depicted in Fig. 10.
Fig. 4. Inhibitory effect of TBME on protein oxidation levels in iron overloaded mice. Mice were randomly divided into six groups (blank, B; control, C; 50 mg kg⁻¹ b.w. TBME, S50; 100 mg kg⁻¹ b.w. TBME, S100; 200 mg kg⁻¹ b.w. TBME, S200; desirox group, D) and treated as described in ‘experimental design’ section. Protein carbonyl content was assayed to measure the extent of protein oxidation. Values are expressed as mean ± SD (n = 6). ***p < 0.001 compared with blank and ##p < 0.01, ###p < 0.001 compared with control.

Fig. 5. Collagen content in different treated mouse liver. Mice were randomly divided into six groups (blank, B; control, C; 50 mg kg⁻¹ b.w. TBME, S50; 100 mg kg⁻¹ b.w. TBME, S100; 200 mg kg⁻¹ b.w. TBME, S200; desirox group, D) and treated as described in ‘experimental design’ section. Hydroxyproline content was assayed to determine the liver fibrosis. Values are expressed as mean ± SD (n = 6). ***p < 0.001 compared with blank and ###p < 0.001 compared with control.
Fig. 6. Photomicrograph of mice liver sections (staining with haematoxylin and eosin) x 40; (A) Normal mice liver. (B) Iron intoxicated (iron-dextran, 100 mg kg\(^{-1}\) b.w.) liver section showing necrosis, ballooning degeneration and loss of cellular boundaries. (C) Liver section treated with iron-dextran + 50 mg kg\(^{-1}\) b.w. TBME. (D) Liver section treated with iron-dextran + 100 mg kg\(^{-1}\) b.w. TBME. (E) Liver section treated with iron-dextran + 200 mg kg\(^{-1}\) b.w. TBME. (D) Liver section treated with iron-dextran + 20 mg kg\(^{-1}\) b.w. desirox also shows reduced necrotic area and increased number of hepatocytes.

Fig. 7. Effect of TBME on hepatic iron content in different treated mouse liver. Mice were randomly divided into six groups (blank, B; control, C; 50 mg kg\(^{-1}\) b.w. TBME, S50; 100 mg kg\(^{-1}\) b.w. TBME, S100; 200 mg kg\(^{-1}\) b.w. TBME, S200; desirox group, D) and treated as described in ‘experimental design’ section. Values are expressed as mean ± SD of six mice. ***p < 0.001 compared with blank and ###p < 0.001 compared with control.
Fig. 8. Serum ferritin levels in different treated mouse. Mice were randomly divided into six groups (blank, B; control, C; 50 mg/kg b.w. TBME, S50; 100 mg/kg b.w. TBME, S100; 200 mg/kg b.w. TBME, S200; desirox group, D) and treated as described in ‘experimental design’ section. Serum ferritin levels were assayed to demonstrate the degrees of iron overload. Values are expressed as mean ± SD of six mice. *p < 0.05, **p < 0.01, ***p < 0.001 compared with blank and #p < 0.05, ##p < 0.01 compared with control.

Fig. 9. Dose dependent formation of the [Fe(ferrozine)$_3$]$^{2+}$ complex following release of Fe$^{2+}$ from ferritin by TBME with time. The reductive release of ferritin iron was quantified by measuring the formation of the ferrous complex of ferrozine, [Fe(ferrozine)$_3$]$^{2+}$ at 562 nm using a Shimadzu UV-VIS spectrophotometer.
Results showed a positive correlation coefficient ($R = 0.9161$) between the reducing power and the amount of released ferritin iron (%), which is highly significant ($p<0.001$). The results signified the higher reducing power of TBME resulted in the release of higher amount of iron from ferritin.

4. DISCUSSION

Iron is the most common cofactor within the oxygen handling biological machinery and specifically, lipid peroxidation of biological membranes is the main pathogenic mechanism of iron overload induced tissue damage (Bonkovsky, 1991). In hepatic iron overload, iron begins to initiate and propagate ROS induced oxidative damage to all biomacromolecules (proteins, lipids, sugar and DNA) ultimately leading to critical failure of biological functions and cell death (Sayre et al., 2005). An effective therapeutic approach could be sequestering of catalytically active iron present in the cytosol or released by chelation from other cell iron stores (Rothman et al., 1992) and at the same time, protecting membranes from free radicals produced in the cells (i.e., antioxidant activity).

The results from Fig. 1 suggest that, TBME showed impressive results indicating that it has iron chelating activity. Fe$^{3+}$-mediated Fenton reaction induced nicking of pUC18 plasmid DNA was also studied and it was observed that the DNA strand scission was significantly subdued with the addition of TBME in a dose dependent manner. The significant reduction in the formation of nicked DNA and increase in supercoiled DNA in the presence of the TBME reveal its excellent iron chelating activity.

Serum enzymes and bilirubin are the important parameters measured for the diagnosis of liver diseases. Iron overloaded hepatic injury leads to leakage of cellular enzymes into the bloodstream, resulting in amplified levels of serum ALAT, ASAT, ALP and bilirubin (Reddy and Lokesh, 1996). TBME reduced the serum enzymes as well as the total bilirubin levels, indicating its protective effect over liver and improvement in its functional efficiency.

Intrinsic defence mechanism of cells against oxidative stresses involves the range of antioxidant enzymes such as SOD, CAT, GST or compounds such as GSH (Beckman and Ames, 1998). Iron overloaded situation gives rise to major oxidative stress and lowers the levels of these enzymes involved in body’s antioxidant defence system. Oral administration of TBME significantly mended the levels antioxidant enzymes and thus helping revival from iron induced damage.

The most important hypotheses for the mechanism of hepatocellular injury in chronic iron overload is the peroxidative damage to the lipid membranes of cellular organelles resulting in structural and functional alterations in cell integrity through loss of lysosomal membrane (Bonkovsky et al., 1981). The present study demonstrates the lipid peroxidation inhibiting capability of TBME and this activity supposed to be associated with its iron chelating activity.

Protein oxidation is another significant factor in the pathogenesis of iron overload induced hepatic damage. The iron overload induced ROS can lead to oxidation of protein backbone resulting in modification of catalytic and structural integrity of various important proteins (Dalle-Donne et al., 2003). The spectrophotometric
DNPH assay was used for quantification of protein CO
-groups, as marker of oxidative damage and the results
presented in this work clearly establish that TBME
efficiently reduced the protein carbonyl content produced
in protein oxidation.
Liver fibrosis is the excessive accumulation of
extracellular proteins including collagen that occurs in
most types of chronic liver diseases (Ramm and Ruddell,
2005). Iron overload cause a significant increase of
hydroxyproline, which is the marker of fibrosis.
Treatment with TBME significantly reduced
hydroxyproline content in iron intoxicated mice,
demonstrating the hepatic fibrosis inhibitory potency of
the plant extract.
Histological observations are done along with the
level of various biochemical parameters in circulation to
mark the extent of hepatic damage. Normal hepatic
parenchyma surrounding veins are observed in liver
sections from normal mice, whereas extremely damaged
cytoarchitecture is found to be prevalent in those from
iron overloaded group. As can be found from other
serum and liver parameter studies, the rejuvenation of
damaged liver was corroborated in the improvement in
histopathological alterations which signifies the in-situ
evidence of hepatoprotective effect of plant extract.
The decrease in liver iron deposition induced by
TBME treatment support its iron chelating potency.
Body’s iron level is positively correlated with a
ubiquitous intracellular protein, ferritin, that stores iron
in a non-toxic form and also helps prevent iron from
mediating oxidative damage to cell constituents (Ford
et al., 1984). In general, the amount of ferritin in blood reflects
the extent of iron stores in the liver, making it one of the
key parameters in the management of hemochromatosis
and other iron overload diseases. In this study, the
ferritin level was found enhanced in iron overloaded
mice, whereas, the level significantly reduced after the
treatment with TBME.
Within cell, ferritin served as the storage protein for
excess iron in ferric state. In case of iron overload,
various iron chelators are administered to attenuate the
situation, but, most of these chelators have limited
binding activity for ferric iron as well as iron in ferritin is
not properly accessed to them. So, iron chelation therapy
is dependent on the reductive release of ferritin iron,
which is achieved by supplemented addition of a
reducing agent such as ascorbate to increase the
availability of storage iron to chelators (O’Brien, 1974).
However, it is obvious that intrinsic reducing property of
an iron chelator should definitely increase the efficiency of
iron chelation therapy to treat iron overload. Previous
results had shown the reductive ability of TBME
(Hazra et al., 2010) as well as in the present study, a
significant positive correlation between reducing power
and iron released from ferritin has been well established.
The current result exhibited the reductive ability of TBME
to release ferritin iron dose dependently with time and can
be used as iron chelating drug to treat iron overload.

5. CONCLUSION

To summarize, it can be concluded that TBME,
which possess both reducing power and iron chelating
activity, can reduce the toxic level of iron in iron
overload mice and hence protect liver from oxidative
stress and fibrosis. Serum enzyme and serum ferritin
levels, both indicators of severe iron overload, are also
effectively lowered owing to its administration. Taken
together, the current findings may be of use in
elucidating the pharmacology and application of TBME
as a hepatoprotective or potential iron chelating drug in
the treatment of iron overload diseases.

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