Original research

Hypertension potentiates cataractogenesis in rat eye through modulation of oxidative stress and electrolyte homeostasis

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Abstract

Purpose: To evaluate modes of cataractogenesis in the hypertensive state by using different hypertensive animal models, including fructose, cadmium chloride (CdCl2), Nω-nitro-L-arginine methyl ester (L-NAME), and two-kidney, one clip (2K1C) method.

Methods: Male Sprague–Dawley albino rats (150–180 g) were divided into different groups, each group containing six animals. Hypertension was induced in animals via six weeks administration of fructose (10% solution in drinking water), CdCl2 (0.5 mg/kg/day, i.p.), and L-NAME (20 mg/kg/day, p.o.) in their respective groups and NaCl (0.9% solution in drinking water) in the 2K1C group. The Ramipril-treated group (2 mg/kg/day, orally) served as a standard group for the 2K1C animal model. Blood pressure was measured biweekly using non-invasive blood pressure system. The biochemical parameters in serum and eye lenses were evaluated after six weeks of the experimental protocol.

Results: Hypertensive animal models showed significant induction of systolic and diastolic blood pressure and modulation of oxidative stress through depletion of antioxidants, including glutathione peroxidase, catalase, superoxide dismutase, glutathione, and elevation of malondialdehyde in serum and eye lenses. A significant elevation of ionic contents (Na+ and Ca2+) and reduction of total protein and Ca2+ ATPase activity in eye lenses were observed in all hypertensive animal models except L-NAME when compared with the normal group. The significant restoration of the antioxidants, Malondialdehyde (MDA) total protein, and ionic contents in the eye lenses concomitant with reduction of blood pressure were observed in the ramipril-treated group as compared to the 2K1C animal model. The results indicate that the fructose, CdCl2, and 2K1C models showed pronounced cataractogenic effects in the rat eye lenses.

Conclusion: Based on our findings, it can be concluded that systemic hypertension significantly increases the risk of cataract formation in the rat eyes via modulation of the antioxidant defense mechanism and electrolyte homeostasis.

Keywords: Hypertension; Cataract; Oxidative stress; Fructose; CdCl2; Two-kidney, one clip

Introduction

Cataract is the leading cause of blindness in the world and the most prevalent ocular disease.1 The number of cataract-blind is expected to increase dramatically in coming decades as the number of elderly in the world's population increases. It is suggested that the number of cataract-blind could reach close to 40 million by the year 2025.2 There are several risk factors which are associated with induction of cataractogenesis, such as diabetes, oxidative stress, ultraviolet radiation, age, etc.3 Several epidemiological studies revealed that hypertension is also associated with cataract, but sometimes it is significant1,4,5 and sometimes not.6 Their studies, however, lacked the exact exploitation of biological mechanism involved in the entire process regarding the exacerbation of cataract in hypertensive state. Some preclinical studies
indicate that systemic hypertension may alter the electrolyte homeostasis through inhibition of $\text{Na}^{+} \text{K}^{+}$ ATPase pump activity in the lens which causes the cataract formation, but at present, another possible mechanism involved in cataractogenesis through hypertension is still unclear. In this perspective, we designed an experimental study to evaluate modes of cataractogenesis in the hypertensive state by using different hypertensive animal models, such as fructose, cadmium chloride (CdCl$_2$), $\text{N}_\omega$-nitro-L-arginine methyl ester (L-NAME), and two-kidney, one clip (2K1C), which have different modes of cellular pathogenesis. In fructose animal model, chronic fructose administration induces hypertension by systemic oxidative stress, sympathetic overactivity, and increased production of vasoconstrictor molecules, viz endothelin-I and angiotensin-II. CdCl$_2$ induced hypertension is related to Ca$^{2+}$ mimicking contractile activity of cadmium ion on vascular smooth muscles, oxidative damage, and vascular endothelial dysfunction. The 2K1C model elevates the plasma renin activity, which activates the angiotensin II-mediated hypertensive actions, and L-NAME modulates the nitric oxide (NO) level in smooth muscles, which lead to elevation of blood pressure.

**Methods**

**Drugs and chemicals**

L-NAME was purchased from Sigma–Aldrich (St. Louis, MO, USA). $\alpha$-fructose and CdCl$_2$ were purchased from HiMedia chemicals (Mumbai, India). Ramipril was obtained from Cipla Limited (Mumbai, India) as a gift sample. Other chemicals and reagents used were of analytical grade.

**Experimental animals**

Sprague–Dawley albino male rats (150–180 g) were used for the experimental study and were housed under standard environmental condition (23 ± 2 °C, with 55 ± 5% humidity and 12 h light/dark cycle) according to the guidelines of Committee for the Purpose of Control and Supervision on Experiments on Animals (CPCSEA), Govt. of India, and were fed a standard pellet diet with ad libitum water for six consecutive weeks. The systolic (SBP) and diastolic blood pressure (DBP) in each group were monitored biweekly via non-invasive blood pressure system (NIBP; CODA-08 Channel, Kent scientific, USA), and biochemical parameter in serum and eye lens were determined after six weeks in sacrificed animals.

**Surgical procedure for 2K1C model**

2K1C was performed on all the rats by anesthetizing with ketamine and xylazine (60:10 mg/kg, i.p.). The kidney was visualized by a left lateral abdominal incision, and the left renal artery and ureter were ligated by a silk thread. The muscle and skin layer (incision site) were sutured with highly sterile suture needles. After surgery, rats were allowed to drink water ad libitum, with no further treatment. All uninephrectomized animals were given 0.9% NaCl in the drinking water for six consecutive weeks.

**Blood collection**

Animals were sacrificed after six weeks, blood was collected from each group via cardiac puncture, and serum was separated and stored at 2–8 °C for further biochemical analysis.

**Preparation of lens homogenate**

The eyeball was isolated from the sacrificed animals. Lenses were dissected via posterior approach, washed with cold saline, and stored with saline at −20 °C until analysis. Lens homogenate was prepared from both lenses of each animal in 10 volumes of 0.1 M potassium phosphate buffer (pH 7). The homogenate was centrifuged at 10,000 rpm for 1 h, and the supernatant was separated and used for biochemical analysis.

**Determination of lenticular opacity**

The lenticular opacity of the experimental groups was determined by the photographic method based on the appearance of graph lines through the lens. The eye lenses were dissected via a posterior approach, put on graph paper immediately, and photographed by a digital camera (Sony Cybershot DSC-W810). The graph lines would appear clearly in the transparent lens and cloudy or not visible in the cataractous lens.
**Biochemical parameters**

**Enzymatic and non-enzymatic antioxidants in serum and lens**

**Glutathione peroxidase (GPx).** The activity of the GPx was assayed by using the method of Tappel (1978). Briefly, 0.2 ml of the test sample was reacted with 0.2 ml of 0.4 M tris buffer, 0.1 ml of 10 mM sodium azide, 0.1 ml of 0.2 mM hydrogen peroxide, and 0.2 ml of glutathione. The reaction mixture was incubated at 37 °C for 10 min, and then reaction was arrested by the addition of 0.4 ml of 10% trichloroacetic acid (TCA). The absorbance was read at 340 nm.23

**Catalase (CAT).** The CAT activity was monitored at 240 nm for 30 s at 25 °C by using the method of Aebi et al (1984). One unit of CAT is defined as the amount of enzyme required to decompose 1.0 M of hydrogen peroxide into water per minute at pH 7.0 and 25 °C.20

**Superoxide dismutase (SOD).** The reaction mixture contained 1 ml of homogenate (10% w/v in 0.25 M sucrose buffer), 1.2 ml of sodium pyrophosphate buffer (0.052 M, pH 8.3), 0.1 ml of 186 μM phenazonium methosulphate, 0.3 ml of 300 μM nitro blue tetrazolium, and 0.2 ml of 780 μM NADH. The reaction mixture was diluted up to 3 ml by distilled water and incubated at 30 °C for 60 s. The reaction was arrested by addition of 1.0 ml glacial acetic acid and stirred vigorously. 4.0 ml of n-butanol was added to the mixture, shaken well, allowed to stand for 10 min, and centrifuged at 2500 rpm for 5 min. Butanol layer was separated out, and absorbance was measured at 560 nm against a butanol blank. Xanthine oxidase enzyme served as the control.21

**Glutathione (GSH).** The lens and serum glutathione level were measured using Ellman’s reagent (Ellman, 1959). Briefly, the homogenized lens (10% w/v in cold 20 mM EDTA) and serum were deproteinized with 0.5 ml of 10% TCA and centrifuged. The protein free supernatant (0.2 ml) was treated with 1 ml of 2.5% ammonium molybdate and 0.4 ml of amino naphth sulfonic acid. The absorbance of the resultant color product was measured at 412 nm. The pure glutathione was used as a standard for establishing the calibration curve.22

**Malondialdehyde (MDA) in serum and lens**

MDA, an end product of lipid peroxidation, was estimated in the lens homogenate and serum by using the method of Ohkawa (1979). Briefly, 0.2 ml of test sample was reacted with 0.2 ml of 8.1% sodium dodecyl sulphate, 1.5 ml of 20% acetic acid (pH 3.5), and 1.5 ml of 0.81% thiobarbituric acid in succession. The mixture was heated in boiling water for 60 min. After cooling at room temperature, 5 ml of butanol: pyridine (15:1 v/v) solution was added, and the mixture was centrifuged at 5000 rpm for 15 min. The upper organic layer was separated out, and intensity of the resultant pink color was read at 532 nm. Tetra methoxy propane was used as a standard.23

**Total protein content in lens**

The total protein content was measured by using the method of Lowry et al (1951). Briefly, 0.1 ml of lens homogenate was treated with 4.0 ml of alkaline copper solution and allowed to stand for 10 min, and then 0.4 ml of phenol reagent was added very rapidly, mixed quickly, and incubated at room temperature for 30 min for color development. The absorbance of the resultant color product was measured at 610 nm against blank. Bovine serum albumin was used as a standard for establishing the calibration curve.24

**Ca^{2+} ATPase activity in lens**

The Ca^{2+} ATPase activity was determined by using the method of Rorive and Kleinzeller (1974). Briefly, 0.1 ml of the lens homogenate (10% w/v in 0.25 M sucrose) was treated with 0.2 ml of ATP (40 mM in 0.4 M tris-HCl buffer, pH 7) and incubated at 37 °C in a water bath for 30 min. The enzyme activity was stopped by adding 2 ml of 10% TCA, and 0.2 ml of ATP was added and kept in ice for 20 min. The reaction mixture was centrifuged at 2500 rpm for 10 min, and the supernatant was collected. 3 ml of supernatant was treated with 1 ml of 2.5% ammonium molybdate and 0.4 ml of amino naphth sulfonic acid. The absorbance of the resultant color product was measured at 680 nm after 20 min.25

**Ionic contents (Na\(^+\) and Ca\(^{2+}\))**

Ionic contents of the serum and lens were estimated by spectrophotometric method using diagnostic kits (Labcare Diagnostics Pvt. Ltd., India).

**Statistical analysis**

The results were expressed as mean ± standard error of mean (SEM). The significant differences between multiple groups were statistically analyzed by using one-way and two-way analysis of variance (ANOVA). The data were considered statistically significant at p < 0.05. Statistical analysis was performed using Graph Pad Prism 5.0 software (GraphPad software, Inc., USA).

**Results**

**Effects on blood pressure**

During six weeks of experimental protocol, blood pressure (SBP and DBP) was found to increase significantly in a time-dependent manner in each group (except the ramipril-treated group) as compared to the normal group. CdCl\(_2\) and l-NAME models showed a significant increase in SBP (p < 0.001, p < 0.05) and DBP (p < 0.001, p < 0.001) from the second week, respectively, whereas the fructose and 2K1C models showed significant elevation in SBP (p < 0.001) and DBP (p < 0.001) from the fourth week. The ramipril-treated
group showed significant ($p < 0.001$) reduction in SBP and DBP as compared to 2K1C (Fig. 1).

**Effects on lens opacity**

The eye lenses of hypertensive animals (Fig. 2 B to E) were more opaque than the normal control (Fig. 2 A). The eye lenses of fructose, CdCl$_2$ and 2K1C hypertensive animal models were more opaque than the lenses of the L-NAME animal model. The administration of ramipril to 2K1C hypertensive animals showed substantial reduction in lenticular opacity as compared to 2K1C (Fig. 2 F).

**Effects on biochemical parameters in serum**

The significant augmentation in systemic oxidative stress was observed in all hypertensive animal models (Table 1). Fructose, CdCl$_2$, and 2K1C models showed significant ($p < 0.001$) depletion of serum enzymatic antioxidants (GPx, CAT, and SOD) in serum while L-NAME showed depletion in CAT ($p < 0.01$) and SOD ($p < 0.01$) only when compared to normal.

The level of non-enzymatic antioxidant (GSH) and an end product of lipid peroxidation, MDA were significantly reduced in fructose ($p < 0.01$, $p < 0.001$), CdCl$_2$ ($p < 0.01$, $p < 0.05$), L-NAME ($p < 0.05$, $p < 0.05$), and 2K1C ($p < 0.001$, $p < 0.01$) models, respectively, when compared to normal (Table 1). The ramipril-treated group showed significant elevation in serum GPx ($p < 0.01$), CAT ($p < 0.01$), SOD ($p < 0.001$), and GSH ($p < 0.001$) level and reduction in serum MDA ($p < 0.01$) level as compared to the 2K1C group.

**Effects on biochemical parameters in eye lenses**

The hypertensive animal models showed marked oxidative stress via attenuation in enzymatic and non-enzymatic antioxidants in rat eye lenses (Table 2). The fructose, CdCl$_2$ and 2K1C models showed significant ($p < 0.001$) reduction in GPx, CAT, SOD, and GSH levels, while L-NAME model showed significant reduction in GPx ($p < 0.05$), CAT ($p < 0.001$), and SOD ($p < 0.05$) only when compared to normal.

MDA level (Table 2) was significantly increased in fructose ($p < 0.001$), CdCl$_2$ ($p < 0.05$), 2K1C ($p < 0.001$), and L-NAME ($p < 0.01$) models. The total protein content and Ca$^{2+}$ ATPase activity in eye lenses were significantly reduced in fructose ($p < 0.001$, $p < 0.01$), CdCl$_2$ ($p < 0.001$, $p < 0.001$), and 2K1C ($p < 0.001$, $p < 0.01$) models, respectively, whereas L-NAME showed non-significant reduction as compared to normal (Fig. 3).

The significant ($p < 0.001$) elevation of serum antioxidants (GPx, CAT, SOD, and GSH) and depletion of MDA level were observed in ramipril-treated rats. Moreover, total protein content ($p < 0.001$) and Ca$^{2+}$ ATPase activity ($p < 0.01$) in eye lenses were significantly increased by ramipril in 2K1C hypertensive animal model.

**Effects on ionic (Na$^+$ and Ca$^{2+}$) contents**

Ionic contents in serum and lens were altered markedly in CdCl$_2$ and 2K1C models than fructose and L-NAME models (Fig. 4). Serum Na$^+$ level was significantly ($p < 0.01$) increased in CdCl$_2$ and 2K1C models whereas its level was unaffected in fructose and L-NAME models. Lens Na$^+$ was also significantly increased by fructose ($p < 0.01$), CdCl$_2$ ($p < 0.001$), and 2K1C ($p < 0.001$) while L-NAME failed to show the significant effect as compared to normal. Serum Ca$^{2+}$ was significantly ($p < 0.05$) reduced in CdCl$_2$ model only whereas lens Ca$^{2+}$ was significantly increased in fructose ($p < 0.05$), CdCl$_2$ ($p < 0.001$), and 2K1C ($p < 0.01$) models when compared to normal. The significant restoration of Na$^+$ ($p < 0.001$) in serum and lens, and Ca$^{2+}$ ($p < 0.01$) in the lens was observed in the ramipril-treated group.

**Discussion**

In the present study the fructose, CdCl$_2$, L-NAME, and 2K1C hypertensive animal models showed significant increase in hypertension in terms of elevation of SBP and DBP during six weeks of the experimental protocol. The mode of induction of hypertension was independent in each animal model.

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**Fig. 1.** Effects of hypertensive animal models on (A) systolic blood pressure and (B) diastolic blood pressure. Values are expressed as mean ± standard error of mean (n = 6), *p < 0.05, **p < 0.01, ***p < 0.001 when compared to normal and †p < 0.05, ‡p < 0.01, ‡‡p < 0.001 when compared to 2K1C (two-way ANOVA followed by Bonferroni's post hoc test).
Numerous studies reported that the reactive oxygen species (ROS) level is increased in the chronic hypertensive state that is indicated by depletion of serum antioxidants and elevation of MDA level. The increased blood pressure markedly produces the ROS through vascular stimulation by mechanical stretch and activation of the renin-angiotensin system and further leads to oxidative stress. In our study, we found that chronic hypertension in different animal models significantly augmented the oxidative stress as indicated by depletion of antioxidants (SOD, GPx, CAT, and GSH) and elevation of MDA level in serum. The reduced serum antioxidants via hypertension may further lead to cellular damage of several organs, including the eye lenses via modulation of antioxidant defense mechanism that is confirmed by depleting antioxidants and elevating MDA level in the eye lenses.

The increased oxidative stress and ionic imbalance in lenses are key mechanisms for the development of cataract; it can modulate the protein content and lenticular opacity. The increased ROS in the lens can cause the molecular damage of lens protein and phospholipids leading to lipid peroxidation and depletion of the antioxidant enzymes such as SOD, GPx, and CAT, resulting in elevation of lenticular opacity and cataract formation. SOD is a chain-breaking antioxidant. It converts superoxide to H$_2$O$_2$ and scavenges the superoxide anion to form hydrogen peroxide. GPx is responsible for degrading the level of H$_2$O$_2$, and CAT helps to keep the level of free radicals below toxic levels. Thus, enhanced oxidative stress in lens induced by hypertension might be responsible for the risk of cataract formation. Moreover, ramipril, an angiotensin-converting enzyme (ACE) inhibitor administered to the 2K1C-induced hypertensive animals, significantly restored the serum and lens antioxidants and MDA level concomitant with reduction of blood pressure, suggesting that high blood pressure is potentially associated with increased ROS formation in serum and lens and further leads to cataractogenic effects on eye lenses.

The elevated MDA level and reduced GSH level in hypertensive rat lenses also implicated the progression of cataractogenesis in rats. The reduced level of GSH is anticipated to cause the formation of disulphide bonds through sulphhydryl oxidation of lens crystalline and to cause the cross-linkage, which leads to cataract formation whereas increased level of MDA, a product of membrane lipid peroxidation, causes the

| Group   | GPx (µM/min/mg of Hb) | CAT (µM of H$_2$O$_2$ consumed/min/mg of Hb) | SOD (µM/min/mg of Hb) | GSH (µM/ml) | MDA (µM/ml) |
|---------|----------------------|--------------------------------------------|-----------------------|-------------|-------------|
| Normal  | 5.07 ± 0.33          | 0.53 ± 0.03                                | 2.11 ± 0.06           | 2.56 ± 0.14 | 2.63 ± 0.12 |
| Fructose| 2.40 ± 0.19          | 0.34 ± 0.02                                | 1.05 ± 0.07           | 1.65 ± 0.14 | 5.04 ± 0.38 |
| CdCl$_2$| 3.47 ± 0.27          | 0.38 ± 0.02                                | 1.35 ± 0.14           | 1.73 ± 0.11 | 3.81 ± 0.24 |
| t-NAME  | 4.75 ± 0.21          | 0.41 ± 0.01                                | 1.59 ± 0.11           | 2.33 ± 0.16 | 3.43 ± 0.26 |
| 2K1C    | 3.48 ± 0.20          | 0.36 ± 0.03                                | 1.24 ± 0.13           | 1.17 ± 0.07 | 4.21 ± 0.35 |
| Ramipril| 4.76 ± 0.13          | 0.48 ± 0.02                                | 1.89 ± 0.07           | 2.29 ± 0.13 | 2.77 ± 0.17 |

Values are expressed as mean ± standard error of mean (n = 6). $^a$p < 0.05, $^b$p < 0.01, $^c$p < 0.001 when compared to normal and $^d$p < 0.05, $^e$p < 0.01, $^f$p < 0.001 when compared to 2K1C (one-way ANOVA followed by Newman–Keuls post hoc test). CAT: catalase; GPx: glutathione peroxidase; GSH: reduced glutathione; t-NAME: N$_x$-nitro-l-arginine methyl ester; MDA: Malondialdehyde; SOD: superoxide dismutase; 2K1C: two-kidney, one clip.
development of lenticular opacity due to cross-links between membrane lipids and proteins.\textsuperscript{40}

The transparency of the lens is also dependent on protein content and intracellular ions, especially Ca\textsuperscript{2+} and Na\textsuperscript{+}. Accumulation of these ions leads to the aggregation of lens proteins and further reduces the total and soluble proteins, and increases the insoluble proteins, and causes lenticular opacity. The activation of calpain via increased Ca\textsuperscript{2+} in lens leads to enhanced proteolytic activity in lens epithelial cell that results in digestion of cytoskeletal and junctional proteins and initiates lenticular opacity.\textsuperscript{32,33} As a limitation in our study, we estimated the lens total protein content only instead of soluble/insoluble proteins, which are more relevant to the cataractogenesis. In our studies, we found that lens Ca\textsuperscript{2+} and Na\textsuperscript{+} were noticeably enhanced, and total protein was markedly reduced in fructose, CdCl\textsubscript{2}, and 2K1C animal models than the L-NAME animal model. The accumulation of Ca\textsuperscript{2+} and Na\textsuperscript{+} in the eye lenses of CdCl\textsubscript{2}-treated rats may be due to the primary toxicity of the Cd\textsuperscript{2+} that may produce oxidative damage to the vascular walls and eye lenses, and further leads to hypertension and cataractogenesis.

The cataractogenic effects of 2K1C hypertensive animal model might be related to the hyperactivity of the ocular renin-angiotensin system. It is established that 2K1C trigger the angiotensin II-mediated actions. The literature revealed that serum angiotensin-II cannot enter the eye,\textsuperscript{45} but it can modulate the action of ocular renin-angiotensin system\textsuperscript{46} and

| Group        | GPx (\(\mu\text{M/min/mg protein}\)) | CAT (\(\mu\text{M of H\textsubscript{2}O\textsubscript{2}}\) consumed/min/mg protein) | SOD (\(\mu\text{M/min/mg protein}\)) | GSH (\(\mu\text{M/mg protein}\)) | MDA (\(\mu\text{M/mg protein}\)) |
|--------------|-------------------------------------|---------------------------------|-------------------------------------|---------------------------------|---------------------------------|
| Normal       | 3.69 ± 0.24                         | 0.46 ± 0.02                     | 2.74 ± 0.05                         | 2.68 ± 0.17                     | 0.35 ± 0.023                    |
| Fructose     | 1.69 ± 0.08\textsuperscript{e}       | 0.26 ± 0.01\textsuperscript{e}  | 0.82 ± 0.02\textsuperscript{e}     | 1.54 ± 0.09\textsuperscript{e}  | 0.61 ± 0.020\textsuperscript{e} |
| CdCl\textsubscript{2} | 1.14 ± 0.07\textsuperscript{c}         | 0.30 ± 0.01\textsuperscript{c}  | 1.89 ± 0.04\textsuperscript{c}     | 1.14 ± 0.21\textsuperscript{c}  | 0.45 ± 0.02\textsuperscript{c}  |
| L-NAME       | 3.07 ± 0.16\textsuperscript{a}       | 0.25 ± 0.03\textsuperscript{a}  | 2.52 ± 0.05\textsuperscript{a}     | 2.28 ± 0.22                     | 0.47 ± 0.01\textsuperscript{a}  |
| 2K1C         | 1.69 ± 0.04\textsuperscript{a}       | 0.18 ± 0.02\textsuperscript{a}  | 1.73 ± 0.05\textsuperscript{a}     | 0.87 ± 0.04\textsuperscript{a}  | 0.64 ± 0.02\textsuperscript{a}  |
| Ramipril     | 3.42 ± 0.20\textsuperscript{d}       | 0.40 ± 0.01\textsuperscript{d}  | 2.55 ± 0.05\textsuperscript{d}     | 2.35 ± 0.20\textsuperscript{d}  | 0.38 ± 0.01\textsuperscript{d}  |

Values are expressed as mean ± standard error of mean (n = 6). \textsuperscript{a}p < 0.05, \textsuperscript{b}p < 0.01, \textsuperscript{c}p < 0.001 when compared to normal and \textsuperscript{d}p < 0.05, \textsuperscript{e}p < 0.01, \textsuperscript{f}p < 0.001 when compared to 2K1C (one-way ANOVA followed by Newman–Keuls post hoc test). CAT: catalase; GPx: glutathione peroxidase; GSH: reduced glutathione; L-NAME: N\textsubscript{o}-nitro-L-arginine methyl ester; MDA: Malondialdehyde; SOD: superoxide dismutase; 2K1C: two-kidney, one clip.

Fig. 3. Effects on (A) total protein content and (B) Ca\textsuperscript{2+} ATPase activity in rat eye lenses. Values are expressed as mean ± standard error of mean (n = 6), \textsuperscript{a}p < 0.05, \textsuperscript{b}p < 0.01, \textsuperscript{c}p < 0.001 when compared to normal and \textsuperscript{d}p < 0.05, \textsuperscript{e}p < 0.01, \textsuperscript{f}p < 0.001 when compared to 2K1C (one-way ANOVA followed by Newman–Keuls post hoc test).
trigger the angiotensin II-mediated angiogenic and inflammatory action in the eye through production of vascular endothelial growth factor, mitogen-activated protein kinases, and AGE, and further lead to the development of several blinding ocular disorders like diabetic retinopathy, glaucoma, and macular degenerations. In the present study, inhibition of ACE via ramipril showed protective effects against 2K1C-induced hypertension associated cataractogenic action. Moreover, several researchers also found that ACE inhibitors showed beneficial effects in the reduction of cataract through restoration of ionic balance (Na+/K+), free radical scavenging activity, and enhancement of enzymatic and non-enzymatic defense mechanism as well as inhibition of AGE production. Therefore, cataractogenic effects in 2K1C hypertensive animal model might be related to the action of ocular renin-angiotensin system that enhances oxidative stress and ionic imbalance in eye lenses and further leads to the development of cataract.

In the present study, L-NAME significantly elevated the blood pressure similar to other animal models but showed minor reduction of serum and lens antioxidants as compared to other animal models. It showed non-significant effects on total protein content and ions level, indicating that L-NAME produces depletion of the serum NO metabolite (NO) levels in lens causes the cataractogenesis. In addition, Ito et al reported that NO synthase inhibitors (L-NAME) prevented the development of cataract in selenite-treated rats. Therefore, it has mild cataractogenic effects and might be related to oxidative stress through hypertension rather than modulation of NO levels via L-NAME.

The risk of cataractogenesis through hypertension is dependent on its pathogenesis. Uncontrolled hypertension may modulate the oxidative stress and electrolyte homeostasis in the eye lenses and further enhance the risk of cataract formation. Fructose might exacerbate the oxidative damage, and CdCl2-induced hypertension might up-regulate Ca2+ activity and oxidative stress. The 2K1C activates the ocular renin-angiotensin system in the eye lenses via uncontrolled hypertension and further leads to cataractogenesis whereas L-NAME independently affects the lenticular alterations unrelated to hypertension. In conclusion, systemic hypertension seems to be a major risk factor that indirectly affects the lens integrity associated with cataract formation.

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![Fig. 4](image-url) Effects on (A) Na+ and (B) Ca2+ in serum and eye lenses. Values are expressed as mean ± standard error of mean (n = 6), a p < 0.05, b p < 0.01, c p < 0.001 when compared to normal and d p < 0.05, e p < 0.01, f p < 0.001 when compared to 2K1C (one-way ANOVA followed by Newman–Keuls post hoc test).
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