Role of N-Acetylcystein in Protection against Cisplatin Nephrotoxicity

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ABSTRACT

Background: Cisplatin (CDDP) is a cytotoxic therapeutic agent that causes many physiological adverse effects such as nephrotoxicity. Although N-acetylcystein (NAC) a thiol containing antioxidant, has been documented to be effective in attenuating CDDP induced renal injury, the precise mechanisms involved in its renoprotection have not been completely clarified. Methods: Four groups of albino rats were used. The first group injected by saline as control, the second by NAC (i.p.), the third by a single bolus of cisplatin (i.v.), and the last by NAC/CDDP, after 5 days from CDDP injection, investigations were conducted on the levels of serum urea and creatinine, oxidant/antioxidant status by estimation of malondialdehyde, catalase and reduced glutathione, and renal and systemic tumor necrosis factor-alpha (TNF-α). Also, renal tissues were examined histopathologically. Results: Administration of cisplatin to rats induced a significant increase in serum urea and creatinine levels in addition to severe alterations in renal tissue architecture. Cisplatin also increased malondialdehyde and decreased glutathione, and catalase in renal tissues. Also, CDDP increased both renal and systemic TNF-α. Administration of NAC markedly reduced the cisplatin-induced higher serum creatinine and urea levels and counteracted the effects of cisplatin on oxidative stress markers, protected the tissues from the cisplatin-induced lipid peroxidation, and increased TNF-α and improved renal tissue architecture. Conclusion: NAC protection is mediated by preventing the decline of antioxidant status, inhibit malondialdehyde and TNF-α and prevent necrosis and apoptosis from kidneys. These results have implications in use of NAC in human application for protecting against drug-induced nephrotoxicity.

Keywords: Cisplatin; Nephrotoxicity; Reduced glutathione; Malondialdehyde; Catalase; NAC; TNF-α; apoptosis.

INTRODUCTION

Cisplatin, Cis-diaminedichloroplatinum (CDDP), is an effective chemotherapeutic agent against multiple human cancers, including ovarian and head and neck malignancies[1]. Although the therapeutic effects of cisplatin are improved by dose escalation, high-dose therapy is limited by its cumulative nephrotoxicity[2]. Reports of accidental overdose leading to renal failure indicate that cisplatin is
sufficiently potent to be toxic to human [3].

The mechanism of cisplatin-induced nephrotoxicity is not fully understood, but inflammation and oxidative stress have been reported in cisplatin-induced nephrotoxicity [4, 5]. Hannemann and Baumann [6], reported that CDDP induced a high production of the lipid peroxidation product malondialdehyde (MDA), indirectly confirming a relationship between lipid peroxidation and platinum compound-induced nephrotoxicity. Ajith, et al. [5], reported that upon CDDP administration, reduced glutathione (GSH) content and catalase (CAT) activity are decreased.

Cytokines, particularly tumor necrosis factor-α (TNF-α), appear to contribute to cisplatin-induced renal injury and to coordinate the activation of a large network of chemokines and cytokines in the kidney following cisplatin injection [7]. So, inhibition of TNF-α production or its activity reduces cisplatin-induced renal dysfunction and structural damage [8].

A number of anti-inflammatory and antioxidant agents, such as curcumin, vitamin C, and vitamin E, provide protective effects against cisplatin-induced nephrotoxicity [5,9]. Although N-Acetyl cysteine (NAC) a thiol-containing antioxidant, has been documented to be effective in attenuating renal injury induced by CDDP, the precise mechanisms involved in its renoprotection have not been completely clarified [10].

**Aim of the work**

The present study was conducted to evaluate the protective effect of NAC on cisplatin-induced nephrotoxicity by evaluating renal functions, oxidant/antioxidant status of the kidney tissues (in the form of catalase, MDA, reduced glutathione levels), renal and systemic TNF-α and renal histopathology in albino rats.

**MATERIALS & METHODS**

**Animals**

A total of 40 male albino rats weighing 200 ± 25 g were included in the current study and divided into four groups each of 10 rats. Each group was housed in separate metal cage and provided with standard rat chow and drinking water ad libitum.

**Drugs and design of work**

Group I was given intraperitoneal injection of saline once daily (vehicle of NAC) as a control, group II was treated by intraperitoneal injection of 250 mg/kg of NAC once daily, according to Luo et al. (2008) [10], (NAC was obtained from Sedico Pharma Co. for Pharmaceuticals, in powder form and dissolved in saline). Group III received a single bolus injection of cisplatin, 5mg/kg i.v. (MERCK génériques) via the tail vein under light ether anaesthesia to induce acute renal failure [10]. Group IV treated with intraperitoneal injection of NAC (250 mg/kg) once daily, at the second day, rats of that group were injected by a single bolus of cisplatin, 5m/kg, i.e. saline and NAC were given daily for one week, while CDDP was given as one single bolus dose. After five days from cisplatin injections, rats were sacrificed by decapitation and blood samples were collected into clean tubes and allowed to stand at room temperature then centrifuged at 5000 rpm for 10
minutes to get the clear serum, kidneys were also separated from all groups for biochemical and histopathological analysis.

The experimental protocol was approved by the Ethics Review Committee for Animal Experimentation of Faculty of Medicine, Minia University.

Biochemical assay:
Serum urea was determined by using a urea enzymatic kit (Bio-Diagnostic Co., Egypt) and creatinine was assayed by creatinine kinetic kit (Bio-Diagnostic Co., Egypt).

Determination of renal MDA, GSH and CAT (oxidant/antioxidant status):
The renal cortex was separated, and subsequently homogenized in phosphate buffer saline (PBS) containing 0.05% tween-20 and protease inhibitor cocktail. The homogenate was centrifuged at 5000 rpm for 10 min at 4°C. The resulting supernatant was used for determination of MDA, GSH, catalase activity and level of TNF-α.

Catalase activity was determined by measuring the decomposition of hydrogen peroxide according to the manufacturers’ instructions of colorimetric assay kit (Biodiagnostic, Egypt). The values are expressed as U/g tissue. Each unit is defined as the amount of the enzyme required to decompose 1 μM of H₂O₂ per minute [11].

The concentration of MDA as a marker of lipid peroxidation (LPO) was determined according to the method described by Buege and Aust [12], which is based on the reaction with thiobarbituric acid, and was expressed as nmol/g tissue.

GSH level was measured in renal cortex homogenate using colorimetric assay kit (Biodiagnostic, Egypt) according to the manufacturer’s instruction. The method is based on the reduction of 5, 5′-dithiobis (2-nitrobenzoic acid) with reduced glutathione to produce a yellow colored compound. The values were expressed as mmol/g tissue [13].

Measurement of TNF-α level:
The level of TNF-α in serum and renal cortical homogenate was determined by using enzyme-linked immunosorbent assay (ELISA) kit. (KOMA BIOTECH INC. KOREA) according to the manufacturer’s instructions. A standard curve was constructed from standards provided by the manufacturer.

Histopathological examination:
One kidney of each rat was fixed in 10% formalin solution and then dehydrated in ascending grades of alcohol and embedded in paraffin. Sections at 4 μm-thickness were taken, stained with hematoxylin-eosin and evaluated by light microscope. The pathologist was blinded to treatment and evaluated the kidney sections at random for the presence or absence of lesion.

Statistical analysis
All data are presented as mean± SD. Significant difference among data were determined using one-way ANOVA and independent Student’s t test with SPSS version 10. P < 0.05 denoted the presence of statistically significant difference.
RESULTS

Biochemical assay:
Renal function of rats treated by saline and NAC were normal as evidenced by normal serum urea and creatinine (Table 1). On the other hand administration of CDDP to rats is found to induce a marked renal failure, characterized by significant increase in serum urea (172.1±2.33 mg/dl.) and CR levels (3.48±0.33 mg/dl.) as compared to normal saline (urea: 24± 3.23 and CR: 1.04±0.23 mg/dl.) and NAC groups (urea: 24.7±2.31 and CR: 1.22 ± 0.19 mg/dl.) (table 1). The concentration of serum urea and creatinine in group IV (NAC+ CDDP) were significantly reduced (urea: 61.0±0.23 and CR: 1.61±0.19 mg/dl.) compared to the CDDP group.

Table 1: Effect of NAC, CDDP and CDDP plus NAC on serum urea and creatinine levels in rats (values are expressed as mean ± SD)

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Serum urea mg/dl</th>
<th>Serum creatinine mg/dl</th>
</tr>
</thead>
<tbody>
<tr>
<td>Group I (Saline)</td>
<td>24± 3.23</td>
<td>1.04±0.23</td>
</tr>
<tr>
<td>Group II (NAC)</td>
<td>24.7±2.31</td>
<td>1.22±0.19</td>
</tr>
<tr>
<td>Group III (CDDP)</td>
<td>172.1±2.33*</td>
<td>3.48±0.33*</td>
</tr>
<tr>
<td>Group IV(NAC+CDDP)</td>
<td>61.0±0.23**</td>
<td>1.61±0.19**</td>
</tr>
</tbody>
</table>

*p<0.05 versus group I and group II
**p<0.05 versus group III

Oxidant/antioxidant status of renal tissues: Catalase activity and level of GSH were significantly decreased after CDDP administration as compared with saline or NAC groups (table 2), while the concentration of MDA was found to be elevated in the CDDP-treated group (table 2). Administration of NAC was found to significantly improve the activity of catalase and the GSH level in renal tissues induced by CDDP (table 2). Administration of NAC also inhibited the CDDP-induced increase in the MDA level (table 2).

TNF-α level: CDDP evoked a considerable increase in TNF-α abundance in the kidney and serum. Treating the rats with NAC significantly lessened the increment comparing with saline (control) and NAC groups (Fig.1-2).
Fig. 1 Effect of administration of NAC on CDDP-induced elevation in renal TNF-α level. Groups 1: Saline, 2: NAC, 3: CDDP and 4: NAC + CDDP.

Fig. 2 Effect of administration of NAC on CDDP-induced elevation in serum TNF-α level. Groups 1: Saline, 2: NAC, 3: CDDP and 4: NAC + CDDP.

Renal histopathological examination:
Histopathological analysis of kidneys of CDDP-treated rats showed various grades of tubular necrosis as well as apoptotic bodies (Fig 4 A & B), compared with kidneys of group I and II (Fig.3), a finding that denotes irreversible kidney damage. The interstitium showed inflammatory cellular infiltrate (Fig 4 C) and the cells of the renal tubules exhibited hydropic degeneration and many of their luminae contained hyaline casts.
and epithelial debris (Fig. 4: A, B, C and D). Treatment with NAC dramatically improved the CDDP nephrotoxicity and this was evidenced as mild histopathological changes that could be detected in the form of interstitial nephritis and mild hydropic degeneration, the later represents reversible cell injury (Fig. 5).

Fig. 3: Photomicrograph of normal kidney tissue of albino rat represents saline and NAC groups

Fig. 4: A, B, C and D: Photomicrographs of albino rat kidneys represent CDDP group. A & B: shows various grades of tubular necrosis as well as apoptotic bodies. C: the interstitium showed inflammatory cellular infiltrate. D: haemorrhage, hyaline casts, single cell necrosis and hydropic degeneration.
DISCUSSION

Although the exact mechanisms underlying nephropathy secondary to CDDP have not been well established, it has been postulated that cisplatin therapy induces oxidative stress, principally involving reactive oxygen species (ROS) in renal tubular cells [14]. Oxidative stress is caused by various free-oxygen radicals including superoxide anion, hydrogen peroxide and hydroxyl radical [15]. The interaction of ROS with cellular components may result in damage to DNA, proteins, and lipids [16].

A number of therapeutic agents have been evaluated experimentally and clinically against CDDP-induced nephrotoxicity, but none of them proved to be clinically effective as a complete protective agent. Nowadays, using various antioxidants for prophylaxis against toxic nephropathy is a current topic of research. Animal studies support a protective role of antioxidants against many nephrotoxic agent-induced renal dysfunction [17-20]. Furthermore, in humans, especially the use of NAC has been reported to show benefits against some drugs induced-nephropathy (radio contrast substances) [21]. NAC is a precursor of L-cystein and glutathione pathway and a scavenger of free radicals because it interacts with ROS [22]. The reactive thiol group on NAC may also directly bind to and inactivate platinum agents, as has been shown with other thiols [23]. Because of its favourable side-effect profile, low costs and positive results of randomized studies, NAC has gained favour in clinical practice as a preventive therapy.

In the present study, the role of NAC on renal functions, oxidant/antioxidant status, renal and systemic TNF-α and renal histopathology were investigated. A single bolus of intravenous administration of 5mg/kg CDDP to rats induced significant increase in serum urea and creatinine compared to saline and NAC treated animals, suggesting an acute renal failure, which confirmed by disrupted architecture of kidney tissues which...
was in the form of tubular cell necrosis, apoptosis, foamy degeneration, hyaline casts, haemorrhage and interstitial inflammation. Dicky et al. [24], also, demonstrated that CDDP treatment resulted in a consistently high serum urea and creatinine in three days and that these blood values correlated well with renal tubule damage seen histologically. Arany and Safirstein [25], reported that the kidney accumulates CDDP to a higher degree than any other organ, resulting in necrosis of the terminal portion of the proximal tubule and apoptosis in the distal nephron. CDDP induces apoptosis via activation of caspase 3[1,26,27]. Caspase-3 can be activated by caspase 9, which is activated by the release of cytochrome c from the mitochondria[28,29]. NAC at 250 mg/kg administered before CDDP treatment significantly prevented the increase of serum urea and creatinine concentrations and markedly decreased CDDP-induced renal dysfunction and organ damage as confirmed by biochemical assays and histopathological examination. This may be explained by that most thiols are electrophilic and are thought to act as free radical scavengers. Scavenging of ROS by thiols protects tubular epithelium from caspase activation and from cell death [30]. The mechanism of protection of NAC may also be due to covalent binding of the molecule to the platinum, producing an inactive complex [31,32]. Schweitzer [33], showed that sulphur-containing compounds may prevent CDDP from interacting with target molecule, displacing platinum after it is bound.

The balance between oxidants and antioxidants is crucial for the maintenance of the biological integrity of the tissues [34]. The renal antioxidant status, such as CAT and reduced glutathione concentrations in this study, are significantly decreased in cisplatin alone-treated group of animals compared to saline and NAC groups. This is in agreement with the results of Zunino, et al. [35] and Saad and Al-Rikabi [36], who suggested that the levels of GSH in the kidney are significantly reduced by cisplatin treatment. The declined antioxidant status partially explains the mechanism of nephrotoxicity induced by cisplatin. The renal accumulation of platinum and covalent binding of renal protein may also play a role in nephrotoxicity [37]. Treatment of NAC (250mg/kg) along with cisplatin could significantly prevent the depletion of that renal antioxidant system. Similar results were obtained by Luo et al., 2008 [10], who reported that NAC not only abolished but also significantly reversed the GSH decrease induced by CDDP. The decreased concentration of GSH increases the sensitivity of organs to oxidative and chemical injury [38]. Studies with a number of xenobiotics often produced GSH depletion [39]. The depletion of GSH also seems to be a prime factor that permits lipid peroxidation in cisplatin-treated group and subsequent toxicity [16]. Treatment with NAC rendered protection due to increase in GSH concentration could protect the renal cells from oxidant’s attack. Appenroth et al., 1993[40], reported that N-acetylcystein, a sulphydryl compound donor that acts by
replenishing glutathione, was shown to "completely abolish the nephrotoxic effects of CDDP".

In the current study, CAT activity is significantly decreased in cisplatin-treated rats, a similar result was also found by Maliakel, et al. [38]. A study done by Colbay, et al. [41], on nephrotoxic effect of contrast media, suggested that the decrease in antioxidant enzyme activities (catalyse) in kidney tissues was due to over usage or consumption of that enzyme due to increased oxidative stress. The decreased activity of CAT in cisplatin-treated rats, in turn may increase the lipid peroxidation [16]. Thus, the level of MDA, as a result of lipid peroxidation, increased in CDDP-treated animals. This result also coincides with that of Maliakel, et al. [38]. Hannemann and Baumann [6], reported that a high production of lipid peroxidation product MDA, indirectly confirming the relationship between lipid peroxidation and platinum compound-induced nephrotoxicity.

Pretreatment with NAC prevented lipid peroxidation (decreased MDA) by enhancing the renal CAT activity. Luo et al., [10], and Kim et al. [45], reported that reduced MDA level was responsible for prevention of CDDP nephrotoxicity. There is paucity of literatures concerning protective effect of NAC against CDDP effect on CAT and MDA. However, Colbay et al. [41], reported that NAC succeeded to lower MDA level and enhance CAT activity in an experiment studying protective effect of NAC against contrast media nephropathy. Hannemann and Baumann [6], reported that other antioxidants such as alpha tocopherol or (+)-cyanidanol prevented CDDP induced lipid peroxidation.

A crucial role for TNF-α in toxic and ischemic acute renal failure has been widely recognised [7, 8, 43, 44]. In the present study, serum and renal tissue TNF-α level increased significantly after CDDP treatment. Lee et al. [45], reported that CDDP stimulates systemic TNF-α and Ramesh et al. [46], reported that CDDP stimulates TNF-α secreted by kidney epithelial cells. Ramesh and Reeves [7], have shown that cisplatin nephrotoxicity is characterized by the activation of pro-inflammatory cytokines and chemokines, and that TNF-α plays a central role in the activation of these cytokines. Pretreatment with NAC in the current study reduced overproduction of systemic and renal TNF-α. There is a paucity of literatures concerning the effect of NAC on both systemic and renal TNF-α, but Luo et al. [10], reported that NAC inhibited TNF-α mRNA. Inhibition of TNF-α is capable of attenuating CDDP-induced renal injury [7], this also explains renoprotective effect of NAC.

Conclusion and recommendations:

It could be concluded that the antioxidant NAC preserve GSH level, arrange cellular immunity, prevent lipid peroxidation, keep catalase activity, inhibit both renal and systemic TNF-α and protect renal tissues from CDDP toxicity, hence show protective effects against nephrotoxic agents. So, it might be used for protection from CDDP nephrotoxicity after modification of its dose to fit human use as the dose used in the present study was big.
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دور الن-استيل سيستاين في الحماية من تسمم الكلوي بالسيسبلاتين

محفوظ حبيب ب"
أقسام الطب الشرعي والسوسم" و"الكيمياء الحيوية" والباتولوجي
كلية الطب- جامعة المنية

يعتبر ن-استيل سيستاين من أشهر العوامل التي تستخدم في علاج السرطان. ومع هذا فهو يسبب الكثير من الآثار الجانبية الضارة منها تسمم الكلوي. و بالرغم من أن الن-استيل سيستاين هو عقار مضاد للأكسدة (يعتبر على مجموعة الفولاذ) ومعروف أنه يحمي الكلى من التسمم من السيسبلاتين ولكن ميكانيكا هذا الدور لم يتعرف بعد.

في هذا البحث تم حقن فئران التجربة البيضاء بالمحلول المثلي كمجموعة ضابطة، ومجموعة الثانية بتغذير
الن-استيل سيستاين حكفي واحده يومياً في الغشاء البريتوني، ومجموعة الثالثة حكت بتغذير السيسبلاتين بجرعة واحدة في ورد ذاتي مسجل للأشعة الكربون الحاد. و المجموعة الأخيرة تم حقنها بعقار الن-استيل سيستاين والمجموعة. ومجموعة أخري من حقن السيسبلاتين في جميع الفئران. ثم أخذ عينات من تجوامع الدم في المجهر للتحليل. و نايلي و كريستيان و ميخالا و ميالا (قياس الأكسدة) لقياس (MDA) وميثانات البوتيل (GSH) و CAT و TNF-α، في اللذان قد ارتفعت في مجموعة متحركة. الن-استيل سيستاين قد أخفض من نسبة البوتيل والأكسدة في معدل المجموعة. وقد وجد أيضاً أن الن-استيل سيستاين قد أخفض من نسبة النتائج البوتيل والأكسدة في مجموعات متحركة. الن-استيل سيستاين قد أخفض من نسبة النتائج البوتيل والأكسدة في مجموعات متحركة. الن-استيل سيستاين قد أخفض من نسبة النتائج البوتيل والأكسدة في مجموعات متحركة.

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