Effects of Copper Nicotinate Complex on Renal Metallothionein and Metal-Responsive Transcription Factor 1 Genes Expression During 4-Dimethylaminoazobenzene Exposure in Rats

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Abstract

Background: The expression of MT genes in animal’s body is rapidly elevated in response to metal and agents which cause oxidative stress and/or inflammation. The MTF-1 plays an essential role in regulating transcription of MT gene in response to metal ions and oxidative stress. Metallothiamin complex was found to exhibit anti-tumor and anti-inflammatory activity, cytoprotective effect and reduces oxidative stress.

Objectives: The study was designed to evaluate effects of a daily dose of copper nicotinate complex (CNC) on metallothionein-III (MT-III) and metal-responsive transcription factor -1 (MTF-1) expression during 4-dimethylaminoazobenzene (DAB) exposure.

Methods: Ninety rats were divided into five groups. Group I rats were fed on standard diet and was considered the control group; group II rats were fed on standard diet containing DAB (0.06g/100g diet daily) for six months; group III rats received CNC dissolved in saline solution (1mg/kg body weight daily) for six months; group IV rats were pretreated with CNC one month before DAB; group V rats were post treated with CNC after one month of starting feeding on DAB. MT-III and MTF-1 genes expression was assayed by PCR. Renal histopathological changes were examined by light microscopy.

Results: Genes expression in all groups was statistically high at all time intervals as compared with control group, while it was decreased in groups III, IV and V as compared to cancer group (II). In group IV and V, the genes expression was statistically increased as compared with group III. MT expression in control group declined with age, while it was higher at 6th month than 2nd and 4th month in group II. MTF-1 expression was increased at 4th month followed by decrease at 6th month in all studied groups. These results were confirmed by histopathological change. There was an increase in pyknotic and necrotic nuclei in tubular epithelial cells and a mild dilatation of renal tubules.

Conclusion: Scavenging ROS during DAB-induced oxidative stress may be the major role of MT. CNC causes a partial improvement in the genes expression as well as renal tubules, so CNC may be a promising candidate used for protection against oxidative stress.

Keywords: Copper nicotinate complex, 4-Dimethylaminoazobenzene, Metallothionein, Metal-responsive transcription factor-1.

INTRODUCTION

Metallothioneins (MTs) are intracellular low molecular weight metal-binding proteins with high cysteine content (20 of 61-62 amino acids).¹ There are several MT genes in mammalian genome, which encode at least four metallothioneins. In rodents MTs are controlled by 4 genes placed on chromosome 8 (MT-I, MT-II, MT-III and MT-IV, all are functional, which code for apo-proteins MT-1, MT-2, MT-3 and MT-4, respectively), whereas human MTs are controlled by 17 genes on chromosome 16, of which 10 are functional.² In mammals, MTs have been presented largely in the cell cytoplasm, but also in lysosomes, mitochondria and nuclei. MT-1 and MT-2 are localized in most tissues; MT-3 is mainly found in heart, kidney, brain and reproductive organs, whereas MT-4 is restricted to the squamous epithelia.³ The renal cell carcinoma shows three different type of expression as MT-2A elevation, MT-1A decline and MT-1G transcripts. Expression of the MT-3 isof orm has been reported in the tubules of normal kidney as well as...
in renal cell carcinoma along with other isoform.\textsuperscript{16} The major physiological functions of MTs include homeostasis, storage and transport of essential metals Zn and Cu, protection against cytotoxicity of toxic metals and scavenging free radicals (ROS and RNS) generated in oxidative stress (anti-inflammatory, antioxidant and antiapoptotic functions).\textsuperscript{18} MT expression is induced by a multiple factors such as pro- and anti-inflammatory mediators including reactive oxygen species, antioxidants, glucocorticoids, endotoxin, and cytokines (IL-6, TNF-\(\alpha\), interferons).\textsuperscript{20}

The metal-responsive transcription factor-I (MTF-1 or MRE-binding transcription factor-I or metal regulatory transcription factor-I) is an effective transcriptional regulator involved in cellular adaptation to a variety of stresses such as oxidative stress, heavy metals exposure or hypoxia. MTF-1 is the most important activator of metallothionein genes.\textsuperscript{7}

The metalloviatamin; copper (I)-nicotinate complex [CuCl (HNA)\textsubscript{2}]\textsuperscript{+}was found to exhibit various bioactivities such as anti-tumor and anti-inflammatory activity \textsuperscript{30}, cytoprotective effect, superoxide dismutase (SOD) mimic activity. It was also found that it prevents gastric congestion and capillary damage, stimulates blood flow and reduces lipid peroxidation and oxidative stress markers such as nitric oxide.\textsuperscript{17}

So, the objective of this study was to evaluate the effect of copper (I)-nicotinate complex on rat renal MT-III and MTF-1 mRNA genes expression during 4-dimethylaminoazobenzene (DAB) exposure.

**METHODS**

Ninety male albino rats two months old with average weight of 120-150 g were used. They were housed in plastic cages at 23-24 °C with a 12-h light and dark cycle. All rats were given free access to standard chowing commercial pellets [25% protein, 11% fats, 6% fibers, 12% moisture, 8% ash (Ca, Ph, P and Mn) and 38% nitrogen free extract (starch, glucose, fibers, 12% moisture, 8% ash (Ca, Ph, P and Mn) and 38% nitrogen free extract (starch, glucose, fructose, sucrose lactose)] and tap water. All animal experiments were performed according to protocols approved by the Animal Care and use Ethics Committee of Medical Research Institute, Alexandria University, Egypt.

Rats were divided into five groups consisting of 18 animals each:
- **Group I**: rats were fed on standard diet and served as the control group.
- **Group II** (cancer group): rats were fed on standard diet containing DAB (0.06g/100g diet) daily for six months.\textsuperscript{10}
- **Group III**: rats were fed on standard diet and received a daily dose of copper nicotinate complex dissolved in saline solution (CNC, 1mg/kg body weight) by oral gavage for six months.\textsuperscript{31}
- **Group IV**: rats were fed on standard diet and pretreated orally with CNC (1mg/kg body weight/day); one month before feeding on DAB (0.06g/100g diet/day) and continued until the end of the experiment (six months).
- **Group V**: rats were fed on standard diet and post treated orally with CNC (1mg/kg body weight/day) after one month of starting feeding on DAB (0.06g/100g diet/day) and continued until the end of the experiment (six months).

In a time-course experiment, six rats of each group were randomly chosen and sacrificed under light ether anesthesia at 2, 4 and 6 months from the start of the experiment. The rats of group I represent the control group for the other groups at each time interval. The kidney tissues of rats were removed and immediately divided into two parts; the first part was used for total RNA extraction and the second part was fixed in 10% formalin for histological examination.

**Semi-quantification of MT-III mRNA and MTF-1 mRNA gene expression by PCR**\textsuperscript{15,18}

1. **RNA extraction:**

Total RNA was extracted using Gene JET\textsuperscript{TM} RNA Purification Kit (Fermentas) following the manufacturer’s instructions. The presence of RNA extract was detected by electrophoresis using a 1% agarose gel stained with ethidium bromide (0.5 mg/ml) for 40 min at 100V and the gel was observed under UV light. The eluted RNA was collected immediately, placed in ice or stored at 20°C for further processing.

2. **cDNA preparation**

cDNAs were synthesized from the mRNA by RevertAid\textsuperscript{TM} First Strand cDNA synthesis Kit (Fermentas) using a PCR thermocycler (Applied Biosystem, Foster City, CA, USA). cDNA prepared for conventional PCR was stored at 15 to 25°C.

3. **PCR:**

3a: **MT-III mRNA**

The expression level of MT-III mRNA was measured using specific primers, forward 5’-GGAAATTCGCCTGGAATAGGAC-3’ and reverse 5’-GGGTGATCCCCATAGGCTGTGG-3’ to amplify 3’-untranslated region of MT-III cDNA (bases 251-387) with simultaneous amplification of β-actin as an internal control (forward, 5’-CATGGATGACGATATCGCT-3’ and reverse 5’-CATAGGTAGTCTGTAGCGT-3’). PCR was performed with initial DNA denaturation at 95°C for 10 min and then 35 cycles of denaturation at 95°C for 1 min, annealing at 55°C for 1 min and extension at 72°C for 2 min followed by an additional elongation step at 72°C for 10 min. The amplification products were separated electrophoretically on 2% agarose gel with ethidium bromide (0.5 μg/ml) and analyzed by photography.

3b: **MTF-1 mRNA**

For semi-quantitative analysis of MTF-1 gene expression, specific primers were used (forward, 5’-TGCGAGTGCACACAAGGAGA-3’; reverse, 5’-TGAGACTGTACTGAGTGCTAAA-3’). PCR was carried out as follows: initial DNA denaturation for 2 min at 94 °C, one cycle; followed by 30 cycles of denaturation for 30 s at 94 °C, annealing for 30 s at 60 °C, and extension for 1 min at 72 °C, then an additional elongation step at 72 °C for 10 min. Aliquots of the PCR mixtures were separated on 2% agarose gel

For semi-quantitative analysis of MTF-1 gene expression, specific primers were used (forward, 5’-TGCGAGTGCACACAAGGAGA-3’; reverse, 5’-TGAGACTGTACTGAGTGCTAAA-3’). PCR was carried out as follows: initial DNA denaturation for 2 min at 94 °C, one cycle; followed by 30 cycles of denaturation for 30 s at 94 °C, annealing for 30 s at 60 °C, and extension for 1 min at 72 °C, then an additional elongation step at 72 °C for 10 min. Aliquots of the PCR mixtures were separated on 2% agarose gel for 3 min at 94 °C, one cycle; followed by 30 cycles of denaturation for 30 s at 94 °C, annealing for 30 s at 60 °C, and extension for 1 min at 72 °C, then an additional elongation step at 72 °C for 10 min. Aliquots of the PCR mixtures were separated on 2% agarose gel
stained with ethidium bromide (0.5 µg/ml) and photographed. The bands of both genes were scanned and the data were analyzed using UVPDOC-ITLS™ Image acquisition and analysis software (Ultra-Violet product, Ltd. Cambridge, UK) that analyze the relative band density to β-actin band for MT-III mRNA or to GAPDH for MTF-1 mRNA.

### Statistical analysis

Data were fed to the computer using IBM SPSS software package version 20.0. Quantitative data were described using mean and standard deviation. One-way analysis of variance (ANOVA) was used to establish the significance of differences among the different groups. Probability values \((p) < 0.05\) were considered to be statistically significant.

### RESULTS

#### Molecular results

The expression of MT-mRNA and MTF-1mRNA in all studied groups was statistically high at all time intervals as compared with control group. The genes mean values (copies) were decreased in groups III, IV and V as compared to cancer group (II). In group IV and V, the genes expression levels were statistically increased as compared with group III but did not show any significant difference when compared with each other (Tables 1 & 2).

#### Table (1): MT-mRNA gene expression (copies) in all studied groups at the second, fourth and sixth months of the experiment

<table>
<thead>
<tr>
<th>Group</th>
<th>2nd month</th>
<th>4th month</th>
<th>6th month</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control group (I)</td>
<td>0.870±0.05</td>
<td>0.742±0.03</td>
<td>0.371±0.04</td>
</tr>
<tr>
<td>Cancer group (II)</td>
<td>1.164±0.23</td>
<td>1.101±0.15</td>
<td>2.400±0.23*</td>
</tr>
<tr>
<td>CNC group (III)</td>
<td>0.903±0.08b</td>
<td>0.834±0.05b</td>
<td>0.753±0.02ab</td>
</tr>
<tr>
<td>CNC before DAB (IV)</td>
<td>0.993±0.002b</td>
<td>0.982±0.11ac</td>
<td>0.918±0.06abc</td>
</tr>
<tr>
<td>CNC after DAB (V)</td>
<td>1.003±0.002b</td>
<td>0.973±0.12abc</td>
<td>0.991±0.10abc</td>
</tr>
</tbody>
</table>

The mean difference was statistically significant at \(p < 0.05\).

ANOVA test used for comparing between the different studied groups.

\(a\) Comparing with control group (I).

\(b\) Comparing groups III, IV, V with group II.

\(c\) Comparing groups IV, V with group III.

#### Table (2): MTF-1mRNA gene expression (copies) in all studied groups at the second, fourth and sixth months of the experiment

<table>
<thead>
<tr>
<th>Group</th>
<th>2nd month</th>
<th>4th month</th>
<th>6th month</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control group (I)</td>
<td>0.189±0.01</td>
<td>0.282±0.01</td>
<td>0.152±0.01</td>
</tr>
<tr>
<td>Cancer group (II)</td>
<td>0.293±0.01a</td>
<td>0.513±0.02a</td>
<td>0.336±0.03a</td>
</tr>
<tr>
<td>CNC group (III)</td>
<td>0.243±0.03abc</td>
<td>0.454±0.04bc</td>
<td>0.233±0.02abc</td>
</tr>
<tr>
<td>CNC before DAB (IV)</td>
<td>0.274±0.01abc</td>
<td>0.503±0.02abc</td>
<td>0.269±0.02abc</td>
</tr>
<tr>
<td>CNC after DAB (V)</td>
<td>0.276±0.01abc</td>
<td>0.488±0.06abc</td>
<td>0.263±0.01abc</td>
</tr>
</tbody>
</table>

The mean difference was statistically significant at \(p < 0.05\).

ANOVA test used for comparing between the different studied groups.

\(a\) Comparing with control group (I).

\(b\) Comparing with control group (I).

\(c\) Comparing groups III, IV, V with group II.

\(d\) Comparing groups IV, V with group III.

The expression of MT-mRNA in control group declined with age, while at sixth month it was higher than second and fourth month in gp II. The MT-mRNA expression in gp III, IV and V was nearly similar at all time intervals, whereas the expression of MTF-1 was increased in all studied groups at fourth month followed by decrease at sixth month (Figure 1).
**Histopathological results**

The renal histological feature of the control group (I) showed normal glomeruli and tubules (Fig 2), whereas that of the cancer group (II) showed most degeneration of renal tubular noticed by marked dilatation of renal tubule, vesiculated and necrotic tubular epithelial cells as well as, hypertrophic glomeruli (Fig 3a). Studies of several sections of rats received CuCl$_2$ (HNA)$_2$ showed minimal urinary space, mild dilatation of renal tubules, pyknotic nuclei of the mesangial and epithelial tubular cells (Fig 3b). In group IV, regenerative tubules were noticed with mild lose their adhesion (Fig 3c). An increase in atrophied glomeruli with area of proliferating and necrotic and pyknotic tubules epithelial cell as well as mild dilatation of renal tubules was observed in group V (Fig 3d). There was no difference in the histopathological feature with the time duration in same groups (Figures 3 and 4).
Figure (2): Paraffin section micrograph of control rat kidney showing the minimal urinary space (US), normal basophilic mesangial cells (MS), and mild dilatation of normal tubules (RT)

Figure (3): Paraffin section micrograph of rat kidney at 2nd month
(a) Cancer group (II): moderate necrotic tubular epithelial cells as well as atrophy glomeruli (AG) and infiltrating lymphocytes.
(b) CNC group (III): mild adhesion of renal tubules (RT), pyknotic nuclei of the mesangial and epithelial tubular cells.
(c) CNC before DAB group (IV): normal glomeruli with mesangial cells (MC) and normal renal tubules (RT) with pyknotic nuclear epithelial cell.
(d) CNC after BAB group (V): mild atrophy glomeruli (AG) area of degenerative renal tubules with epithelial pyknotic nuclei as well as increased necrotic tubular epithelial cell.
Figure (4): Paraffin section micrograph of rat kidney at 6th month
(a) Cancer group (II): sever necrotic tubular epithelial cells as well as, hyperemic and atrophy glomeruli (AG) and infiltrating lymphocytes.
(b) CNC group (III): normal differentiated mesangial cells. Regenerative tubules were noticed with normal nuclei and contacted cytoplasm.
(c) CNC before DAB group (IV): pyknotic mesangial cell nuclei and epithelial cells of renal tubules, as well as hyaline casts of renal tubules and infiltration of lymphocytes.
(d) CNC after BAB group (V): marked dilatation of renal tubules having necrotic (NC) and pyknotic epithelia nuclei cells, glomeruli with mild urinary space and pyknotic nuclei of mesangial cells.

Table (3): Distribution of the histopathological changes of the kidney tissues in all studied groups

<table>
<thead>
<tr>
<th>Histological alterations</th>
<th>Group I</th>
<th>Group II</th>
<th>Group III</th>
<th>Group IV</th>
<th>Group V</th>
</tr>
</thead>
<tbody>
<tr>
<td>Glomerulus atrophy</td>
<td>5%</td>
<td>40%</td>
<td>11%</td>
<td>13%</td>
<td>18%</td>
</tr>
<tr>
<td>Tubular degeneration</td>
<td>+</td>
<td>++</td>
<td>+</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>Tubular necrosis</td>
<td>+</td>
<td>+++</td>
<td>+</td>
<td>+</td>
<td>++</td>
</tr>
<tr>
<td>Tubular pyknotic nuclei</td>
<td>++</td>
<td>+</td>
<td>++</td>
<td>++</td>
<td>++</td>
</tr>
</tbody>
</table>

DISCUSSION

The expression of MT genes in animal's body is rapidly elevated in response to metal and agents which cause oxidative stress and/or inflammation. The MTF-1 plays an essential role in regulating transcription of MT gene in response to metal ions and oxidative stress. The study indicated that both MT mRNA and MTF-1 mRNA expression in all groups were significantly higher than the control group at all time intervals (after 2, 4 and 6 months from the onset of the experiment). 4-dimethylaminoazobenzene (DAB) is a carcinogenic amino azo dye and metabolically demethylated by microsome cytochrome p-450 to give 4- amino azo benzene (AAB) after which N-hydroxylation and esterification (by sulfo-transferase) may occur. The mechanism of DAB carcinogenic action has been attributed principally to the formation of DNA adducts following metabolic oxidation of the amino group leading to high levels of reactive oxygen species generation which can affect the genes transcription.

The molecular mechanism of MT gene induction results from an enhanced transcription of structural genes following interaction of specific stimuli (trans-acting factors, such as metal and ROS) with response elements (RE, cis-acting elements) in the gene promoter. The MTF-1 is the major player in regulating MT gene expression in physiological conditions and in response to metal ions. ROS can activate MTF-1 indirectly, by
mobilizing Zn from intracellular stores and/or by activating some other intracellular signal pathways such as signal transduction molecules (STM). (17) Zn binds to an inactive cytoplasmic apo-MTF-1 and activates it by inducing phosphorylation of the protein. The activated complex migrates to the nucleus, where it binds to metal response element (MRE) and stimulates the synthesis of MT mRNA. (18) Furthermore, MTF-1 may be existed in an inactive form in a complex with MT (MT-MTF-1); Zn could activate MTF-1 indirectly, by binding to MT in the complex and releasing MT-Zn; the liberated apo-MTF-1 would then be further activated by Zn and bind to MRE. This may represent a basic feed-back mechanism that would control the MT production in normal and stressful conditions. (19) Moreover, the MT gene transcription via antioxidant response element (ARE) can be promoted by various ARE-binding proteins, which can be activated by metal and ROS. A synergistic activation of the MT gene expression via MRE and ARE may be important in nephrotoxicity induced by toxic metals, when the intracellular high concentrations of these metals associate with the increased production of ROS. (20)

The obtained results were denoted that the expression of both genes in Cu complex-received group (group III) was significantly increased as compared to the control group. Previous study stated that the absorption of CuCl (HNA)2 complex to different organs was as its administered form without ligand exchange and that led to the copper distribution to tissues in magnitude up to two fold of its normal content. (21) Besides, the kidney gained a considerable amount of the administered CuCl (HNA)2 and the filtered copper may undergo ligand exchange in distal tubules and that account for the participation of the kidney in copper conservation. (22) Accordingly, the high expression of both gene in kidney may be attributed to the increased accumulation of copper administration, since MT has a pivotal role in homeostasis of Zn and Cu (trapping, storage, distribution and release) (5) and with excess copper, MT may release zinc ions and bind other elements instead. (17) Furthermore, MT gene expression is induced by heavy metal ions through several mechanisms that facilitate MTF-1 transport to the nucleus and induce DNA binding, where it exerts an effect on gene transcription such as MT. (23)

The higher expression of both genes in groups IV and V than the control group as well as the CNC-received group may be due to the carcinogenic effect of DAB which generate free radicals (ROS and RNS) as well as oxidative stress. Whereas, comparing with cancer group, the genes expression was statistically low in group IV (pretreated group) and group V but still higher than that of the control group.

The improvement in renal MT mRNA and MTF-1mRNA expression in groups IV and V as compared to group II could be mostly attributed to the effect of Cu nicotinate complex. These results are in accordance with previous study which elucidate that CuCl (HNA)2 enhance cellular defense mechanisms against free radicals and lipid peroxidation by increasing utilization of copper in cells and tissues. (24) The absorbed copper complex in cells and tissues could react with apoenzymes and apoproteins leading to formation of copper-dependent enzymes and proteins respectively; these are mostly antioxidant. (25) These reactions may account to suppress copper-induced lipid peroxidation. (26) Also the pretreatment with CuCl (HNA)2 had gastroprotective effect by preventing the down regulation in the antioxidants activity whereby fighting each of peroxide elevation and subsequently lipid peroxidation. The blocked in peroxide level by the antioxidant activity of CuCl (HNA)2 emphasize its role as a potent antioxidant, since CuCl (HNA)2 affect activity of several copper enzymes controlling oxidation-reduction reactions (such as copper-zinc superoxide dismutase enzyme) as a cofactor and as a prosthetic component (27)

The present histopathological examination, revealed more changes of the renal tubules architecture, mostly appeared for the necrotic to pyknotic tubular epithelial cells as well as some vesiculated nuclei. These changes of renal nuclear epithelial cells revealed the disturbance balance of nucleic acids or nucleoprotein structure. So, it could be due the cupper binding to newly nucleoprotein synthesis. Also it revealed the more expression of the MT gene in all groups as well as the role of this gene expression caused due to the Cu nicotinate complex binding. MT expression pattern will be changed in the body when any diseases caused due to metal factor, therefore the Cu complex may trigger the expression of MT in animal’s body.

The obtained findings showed that in control group, the expression of MT-mRNA in the renal tissue declines with age. Nordberg 1998 report that the renal tissues capacity to produce MT may be decreasing in old age due to less efficient protein synthesis. (28) Furthermore, when extracellular metal concentration decreased, Zn dissociated from MT resulting in a faster rate of MT degradation. (29) On the other hand in cancer group, the expression of MT-mRNA is higher at sixth month than second and fourth month. This result may be a normal physiologic response to tumor progression that leads to increase oxidative stress and ROS.

Either after second, fourth or sixth months, the MT-mRNA expression in groups III, IV and V (CNC group and CNC-treated groups) is nearly similar that is due to the presence of CNC which is source of Cu that induce MT expression. Cu bound to MT is present as long as cells maintain the capacity to synthesize sufficient MT to accommodate Cu within cells. (30)

The expression of MTF-1 is increased in all studied groups at fourth month follow by decrease at sixth month. MTF-1 genes are transcriptionally activated by Zn load, in addition, to other inducers such as Cd, copper and hydrogen peroxide. The effect of these inducers is
indirectly proportionate by liberating Zn from intracellular stores such as metallothionein. In control group, the downregulation of MTF-1 expression is due to decline in MT expression by aging. At sixth month, the level of H2O2 and Cu is high due to DAB effect and CNC administration respectively. MTs are normally saturated with highly abundant Zn and release Zn upon exposure to copper or H2O2 leading to MTF-1 activation. The downregulation of MTF-1 following its activation is may be due to thioneine (metal-free form of MT, apo-MT); bind Zn with high affinity; which inhibit MTF-1 by Zn sequestration. Activation of MTF-1 and subsequent inactivation by newly synthesized apo-MT might be a basic mechanism to regulate MTF-1 activity upon metal load and cellular stress.

The current data on the expression of MT and MTF-1 mRNA in renal cell carcinogenesis examined emphasize that, the expression of both genes depend on a lot of stimuli. Scavenging ROS during DAB-induced oxidative stress may be the major role of MT. Cu nicotinate complex causes a partial improvement in the oxidative stress. Scavenging during DAB treatment might be a basic mechanism to regulate MTF-1 upon exposure to copper or H2O2 leading to MTF-1 activation. The downregulation of MTF-1 following its activation is may be due to thioneine (metal-free form of MT, apo-MT); bind Zn with high affinity; which inhibit MTF-1 by Zn sequestration. Activation of MTF-1 and subsequent inactivation by newly synthesized apo-MT might be a basic mechanism to regulate MTF-1 activity upon metal load and cellular stress.

**Conflict of Interest:** None to declare

**REFERENCES**


