Different Regulation of p53 Expression by Cadmium Exposure in Kidney, Liver, Intestine, Vasculature, and Brain Astrocytes

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Chronic exposure to cadmium (Cd) is known to adversely affect renal function. Our previous studies indicated that Cd induces p53-dependent apoptosis by inhibiting gene expression of the ubiquitin-conjugating enzyme (Ube) 2d family in both human and rat proximal tubular cells. In this study, the effects of Cd on protein expression of p53 and apoptotic signals in the kidney and liver of mice exposed to Cd for 12 months were examined, as well as the effects of Cd on p53 protein levels and gene expression of the Ube2d family in various cell lines. Results showed that in the kidney of mice exposed to 300 ppm Cd for 12 months, there was overaccumulation of p53 proteins in addition to the induction of apoptosis, which was triggered specifically in the proximal tubules. Interestingly, the site of apoptosis was the same as that of p53 accumulation in the proximal tubules. In the liver of mice chronically exposed to Cd, gene expression of the Ube2d family tended to be slightly decreased, together with slight apoptosis without the accumulation of p53 protein. In rat small intestine epithelial (IEC-6) cells, Cd decreased not only the p53 protein level but also gene expression of Ube2d1, Ube2d2 and Ube2d4. In human brain microvascular endothelial cells (HBMECs), Cd did not suppress gene expression of the Ube2d family, but increased the p53 protein level. In human brain astrocytes (HBASTs), Cd only increased gene expression of UBE2D3. These results suggest that Cd-induced apoptosis through p53 protein is associated with renal toxicity but not hepatic toxicity, and the modification of p53 protein by Cd may vary depending on cell type.

Key words: Cadmium, p53 protein, Ube2d family, Apoptosis, Renal toxicity

INTRODUCTION

Cadmium (Cd) is a ubiquitous toxic heavy metal emanating from industrial and agricultural sources (1). Cd exposure is considered to cause damage of vital tissues such as the kidney, liver, heart, brain and testis (2-4). The liver and kidney are especially sensitive to the toxic effect of Cd (4).

The biological half-life of Cd is so long (10–30 years) in the human body that chronic low-level Cd exposure brings about the accumulation of Cd in the kidney (4). Cd induces apoptosis in proximal tubular cells through several pathways, including the endoplasmic reticulum (ER)-mediated pathway via ER stress and calcium release, and the mitochondria-mediated pathway via direct and indirect activation of mitochondria (5). In addition, our recent studies demonstrated that Cd induced p53-dependent apoptotic cell death in renal proximal tubules both in vitro and in vivo (6) and in human proximal tubular cells (on submission). p53 is a short-lived protein, and under normal conditions is immediately degraded by the ubiquitin-proteasome system (UPS) (7-9). In MCF7 human breast carcinoma cells, ubiquitination of p53 is mediated by UBE2D2 and UBE2D3, which are members of the ubiquitin-conjugating enzyme (Ube) 2d...
family, an E2 ubiquitin-conjugating enzyme family in the UPS (10). Cd was found to suppress the gene expression of all the Ube2d family members (Ube2d1, Ube2d2, Ube2d3 and Ube2d4), and induced accumulation and phosphorylation of p53, followed by p53-dependent apoptotic cell death in renal proximal tubules both in vitro and in vivo (6). In this study, we investigated the characteristics of p53 protein expression and the occurrence of apoptosis not only in the kidney but also in the liver of mice chronically exposed to Cd for 12 months.

In addition to the liver and kidney, the brain and gastrointestinal tract are also thought to be targets of Cd (2,3). There is variation in the sensitivity of individual tissues to Cd-induced damage (11). Therefore, it was also examined whether Cd affects the protein level of p53 and gene expression of the Ube2d family, in such several cell lines as rat small intestine epithelial (IEC-6) cells, human brain microvascular endothelial cells (HBMECs) and human brain astrocytes (HBASTs).

MATERIALS AND METHODS

Animals. All animal experiments were performed in accordance with the Regulation on Animal Experimentation at School of Pharmacy, Aichi Gakuin University, Nagoya, Japan. All procedures to maintain and use mice were approved by the Animal Care and Use Committee for School of Pharmacy, Aichi Gakuin University, Nagoya, Japan. Four-week-old female C57BL/6J mice were purchased from CLEA Japan (Tokyo, Japan) and routinely bled from CLEA Japan (Tokyo, Japan). Four-week-old female C57BL/6J mice were purchased from CLEA Japan (Tokyo, Japan) and routinely bled from CLEA Japan (Tokyo, Japan). After antigen retrieval, kidney and liver sections were stained using anti-p53 antibody (Abcam, Cambridge, MA, USA). The HistoMouse®-Plus kit (Invitrogen, Grand Island, NY, USA) was used to detect p53 antibody. Apoptosis was detected using the TACS® 2 TdT-DAB in situ apoptosis detection kit (Trevingen, Gaithersburg, MD, USA). Microscopy images were obtained using a BIOR-EVO BZ-9000 microscope (Keyence, Osaka, Japan).

Cell culture and Cd treatment. IEC-6 cells were purchased from DS Pharm Biomedical (Osaka, Japan), and cultured in Dulbecco’s modified Eagle’s medium (Sigma-Aldrich, St. Louis, MO, USA), supplemented with 10% fetal bovine serum (FBS) (Gibco, Grand Island, NY, USA), 25 U/mL penicillin and 25 μg/mL streptomycin at 37°C in a humidified incubator containing 5% CO₂. IEC-6 cells were transferred to testing plates (24-well plates for Giemsa staining; 6-well plates for real-time RT PCR; 6 cm plates for western blotting) at 2.5 × 10⁶ cells/cm² and cultured for 48 hr. The culture medium was then discarded and the cells treated with Cd (CdCl₂) (Wako Pure Chemical Industries, Osaka, Japan) in serum-free culture medium for 24 hr.

HBMECs were purchased from DS Pharm Biomedical, and cultured on collagen I-coated dishes (IWAKI, Tokyo, Japan) at 37°C in a humidified incubator containing 5% CO₂. The culture medium HuMedia-EG2 (Kurabo, Osaka, Japan) consisted of the base medium (HuMedia-EB2) supplemented with 2% FBS, 10 ng/mL recombinant epidermal growth factor, 5 ng/mL recombinant basic fibroblast growth factor, 10 μg/mL heparin, and 1 μg/mL hydrocortisone. HBMECs were transferred to testing plates (24-well plates for Giemsa staining; 6-well plates for real-time RT PCR; 6 cm plates for western blotting) at 2.5 × 10⁶ cells/cm² and cultured for 48 hr. The culture medium was then discarded and cells treated with Cd (CdCl₂) in serum-containing culture medium.

HBASTs were purchased from DS Pharm Biomedical, and cultured on collagen I-coated dishes (IWAKI) at 37°C in a humidified incubator containing 5% CO₂. HBASTs were cultured in CSC complete medium kit with serum and Attachment Factor™ (Cell Systems, Kirkland, WA, USA). The HBASTs were transferred to testing plates (24-well plates for Giemsa staining; 6-well plates for real-time RT

p53 protein level. Each mouse liver was homogenized in nine volumes of low-salt RIPA buffer and the homogenate was centrifuged at 10,000 ×g for 10 min at 4°C. Protein levels of p53 in the supernatant were quantified using a p53 Enzyme Immunoassay kit (Assay Designs, Ann Arbor, MI, USA). Total protein concentration was measured using the Quick Start Protein Assay (Bio-Rad, Hercules, CA, USA), to normalize p53 levels.

Detection of apoptosis and p53 in tissue sections. The kidney and liver were fixed in 10% (v/v) neutral buffered formalin solution and embedded in paraffin. Deparaffinized serial tissue sections of 5 μm thickness were used for immunohistochemical staining of p53 and for apoptosis detection. For immunohistochemical staining of p53, antigen retrieval was conducted for 30 min at 90°C in 0.1 M boric acid, pH 7.0. After antigen retrieval, kidney and liver were sectioned in 0.3% (v/v) hydrogen peroxide for 10 min and then stained using anti-p53 antibody (Abcam, Cambridge, MA, USA). The HistoMouse®-Plus kit (Invitrogen, Grand Island, NY, USA) was used to detect p53 antibody. Apoptosis was detected using the TACS® 2 TdT-DAB in situ apoptosis detection kit (Trevingen, Gaithersburg, MD, USA). Microscopy images were obtained using a BIOR-EVO BZ-9000 microscope (Keyence, Osaka, Japan).

GOT and GPT activities. To evaluate the hepatotoxicity, the activities of glutamic oxaloacetic transaminase (GOT) and glutamate pyruvate transaminase (GPT) in the serum were examined. The automatic dry-chemistry analyzer system (Spotchem EZ SP-4430; Arkray, Kyoto, Japan) was used to determine the activities.

p53 protein level. Each mouse liver was homogenized in nine volumes of low-salt RIPA buffer and the homogenate was centrifuged at 10,000 ×g for 10 min at 4°C. Protein levels of p53 in the supernatant were quantified using a p53 Enzyme Immunoassay kit (Assay Designs, Ann Arbor, MI, USA). Total protein concentration was measured using the Quick Start Protein Assay (Bio-Rad, Hercules, CA, USA), to normalize p53 levels.

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PCR) at 4.0 × 10^4 cells/cm^2 and cultured for 48 hr. The culture medium was discarded and the cells treated with Cd in serum-containing culture medium.

**Cell morphology.** After Cd treatment, the cell layer was washed with calcium- and magnesium-free phosphate-buffered saline (PBS; Nissui Pharmaceutical, Tokyo, Japan), fixed with May-Grünwald stain solution (Merck, Darmstadt, Germany) for 3 min, and stained with Giemsa stain solution (Merck) for 30 min at room temperature. Cell morphology was examined microscopically using the OLYMPUS CK2 microscope (OLYMPUS, Tokyo, Japan) at 100× magnification.

**RNA extraction and real-time RT-PCR.** Cd-treated cells were washed twice with ice-cold PBS, and total RNA extracted with the Quick Gene RNA cultured cell kit S (Fujifilm, Tokyo, Japan). cDNA was generated from total RNA using the PrimeScript reverse transcription (RT) reagent kit (Perfect Real Time; TaKaRa Bio, Shiga, Japan). Real-time PCR was performed with the SYBR Premix Ex Taq II (Perfect Real Time, TaKaRa Bio), and a thermal cycler dice real time system (TaKaRa Bio). PCR conditions were: 10 s hot-start at 95°C followed by 40 cycles of 5 s at 95°C and 30 s at 60°C. The oligonucleotide sequences of the primers were as follows: sense, 5'-GAGTAAATGGG-GTTTGTCTTGG-3', and antisense, 5'-CCTTTCCTTTG-GATGGGTGAT-3', for the human UBE2D1 gene; sense, 5'-TGTCATCTGTTCGGGATG-3', and antisense, 5'-TCATTCGGAGCTATTCGG-3', for the human UBE2D2 gene; sense, 5'-GGTCAGGCCCCTGTCTAATG-3', and antisense, 5'-GGGCTTGTAGGTGTGTGCTAATTC-3', for the human UBE2D3 gene; sense, 5'-TGATCCGGTTTTTGAGTGAT-3', and antisense, 5'-GGCCTTGTAGGTGTGTGCTAATTC-3', for the human UBE2D4 gene; sense, 5'-GACCGGTCAAGGCTGAAC-3', and antisense, 5'-TGATCCGGTTTTTGAGTGAT-3', for the human GAPDH gene; sense, 5'-CGGGACCCGTGGGAGATGACTTGTT-3', and antisense, 5'-GACTCCACCTGTGATAGGCGCTG-3', for the rat Ube2d1 gene; sense, 5'-GCACAGTGTACC-TGACACACGA-3', and antisense, 5'-CGGAGAAGTGGTTCTCAGAC-3', for the rat Ube2d2 gene; sense, 5'-GACGGTTTTACGGGT-TCTCAGAC-3', and antisense, 5'-TCTCAGACAGCGGTCTCAGAC-3', for the rat Ube2d3

![Graph](image.png)

**Fig. 1.** Effects of chronic exposure to Cd on gene expression of the Ube2d family and on protein levels of p53 in the liver of mice. (A) Ube2d family gene expression in the liver of mice exposed to Cd for 12 months. mRNA levels were measured by real-time RT-PCR and normalized to β-actin. Data are presented as a ratio of treated groups : controls. (B) Protein levels of p53 in the liver of mice exposed to Cd for 12 months, determined using ELISA. (A,B) Values are mean ± S.D. (n = 5 or 6). *Significantly different from the control group, p < 0.05.
gene; sense, 5'AGCCTTCTCAGGACATACCC-3', and antisense, 5'TAGGTGAGGGACCTTACCC-3', for the rat Ube2d4 gene; sense, 5'CATGGGCTCCAAGGAGTGAAG-3', and antisense, 5'GTTCTGGATGGAATTGTGA-3', for the rat Gapdh gene; sense, 5'ACAACAGGACGCAAG-3', and antisense, 5'GAACGAAGGACACGGCAAAC-3', for the mouse Ube2d1 gene; sense, 5'ATGC- AAAGGTGTTGCT-3', and antisense, 5'TGCTCATTGTTCCAGGTTC-3', for the mouse Ube2d2 gene; sense, 5'CAGTAATGACCAATTGTCT-3', and antisense, 5'GGGTCTGGATGGAATTGTGA-3', for the mouse Ube2d3 gene; sense, 5'CCTAAGGCAACCGTCAG-3', and antisense, 5'AGCCATACAGGACAGCACA-3', for the mouse β-actin gene.

Fig. 2. Effects of chronic exposure for 12 months to Cd on protein expression of p53 and appearance of apoptosis in the kidney and liver of mice. (A) Representative images of p53 staining (upper panels) and TUNEL staining (lower panels) in the kidney of mice. Serial sections of kidney from control mice are shown in the left panels and those of Cd-exposed mice in the right panels. Red arrows indicate p53 expression (upper panel) and apoptotic cells (lower panel). Scale bar = 40 μm. Original magnification ×400. (B) Representative images of p53 staining (upper panels) and TUNEL staining (lower panels) in the liver of mice. Serial sections of liver from control mice are shown in the left panels and those of Cd-exposed mice in the right panels. Yellow arrows indicate apoptotic cells (lower panel). Scale bar = 40 μm. Original magnification ×400.
**Western blotting.** After Cd treatment, cells were washed twice with ice-cold PBS and harvested in sodium dodecyl sulfate (SDS) buffer (2% SDS, 50 mM Tris [pH 6.8], 10% glycerol). Protein concentrations were measured using the BCA protein assay kit (Thermo Fisher Scientific, Waltham, MA, USA). Protein was electrophoresed on SDS-polyacrylamide gels, transferred to polyvinylidene fluoride membranes (ATTO, Tokyo, Japan), and probed with antibodies against p53 (1:1000; Cell Signaling Technology, Danvers, MA, USA), GAPDH (1:2000; American Research Products, Waltham, MA, USA) or β-actin (1:1,000; American Research Products, Waltham, MA, USA). Membranes were subsequently probed with horseradish peroxidase-conjugating secondary antibodies (1:10000; GE Healthcare, Tokyo, Japan), and protein detected by Chemi-Lumi one L (Nakalai tesque, Kyoto, Japan). Chemiluminescent images were obtained using an Image Quant LAS-3000 (Fujifilm, Tokyo, Japan) device.

**Statistical analysis.** Statistical analyses were performed using one-way analysis of variance (ANOVA). When the F value was significant ($p < 0.05$), Bonferroni’s multiple $t$-test was performed for post-hoc comparisons ($p < 0.05$). Student $t$-test was performed for two-sample comparison ($p < 0.05$).

**RESULTS**

**The mouse kidney and liver.** GOT activity in the serum of Cd exposed mice was $213.7 \pm 62.6$ IU/L compared with $60 \pm 18.9$ IU/L in that of control mice. GPT activity in the serum of Cd exposed mice was $63.7 \pm 19.7$ IU/L compared with $23.6 \pm 6.1$ IU/L in that of control mice. Similarly as the mild renal toxicity in the mice exposed to 300 ppm Cd for 12 months (6), chronic Cd exposure triggers mild hepatic toxicity. The only gene expression significantly decreased by Cd exposure was $Ube2d3$ in the liver (Fig. 1A). Hepatic p53 protein levels in mice chronically exposed to Cd were the same as that in control mice (Fig. 1B).

The kidney of mice exposed to Cd for 12 months showed both apoptosis and accumulation of p53 occurring in same positions on the proximal tubules (Fig. 2A). On the other hand, the accumulation of p53 was not observed, but slightly higher levels of apoptosis were detected, in the liver of mice exposed to Cd for 12 months compared with controls (Fig. 2B). Taken together, it is suggested that p53-dependent apoptosis by chronic Cd exposure is preferred in the kidney but not in the liver of mice.

**Various cell lines.** Cd exposure of IEC-6 cells caused dose-dependent destruction of the cell layers (Fig. 3A). Gene expression of $Ube2d1$ and $Ube2d4$ was significantly decreased by as little as 2 μM Cd for 6 hr (Fig. 3B). Cd treatment also tended to decrease gene expression of $Ube2d2$ (Fig. 3B) and markedly decreased protein levels of p53 (Fig. 3C). Despite Cd treatment suppressing the gene expression of the Ube2d family, p53 protein levels in IEC cells were decreased by Cd treatment.

Giemsa staining of HBMECs treated with Cd for 24 hr showed disruption of the cell layers (Fig. 4A). Cd treat-
ment of HBMECs for 6 hr increased gene expression of *UBE2D1* significantly. Gene expression of *UBE2D4* was significantly increased by 50 μM Cd treatment (Fig. 4B). Protein levels of p53 were increased by Cd treatment (50 μM) and even at that low level, slight cell toxicity was evident (Fig. 4C). In HBMECs, Cd treatment increased p53 protein levels with the increase in gene expression of parts of the Ube2d family.

Cd toxicity was seen in HBASTs with 20 and 30 μM treatment (Fig. 5A). Cd treatment increased *UBE2D4* gene expression significantly, with no effect on *UBE2D2* gene expression (Fig. 5B). However, in HBASTs, neither gene expression of *UBE2D1* and *UBE2D3* nor protein expression of p53 was detected with the primers and antibodies described.

These data suggest that the modification of p53 protein by Cd treatment may have cell variability.

**DISCUSSION**

Our previous study demonstrated that 12-month Cd exposure decreased gene expression of Ube2d family, increased p53 protein levels and induced apoptosis in the kidney of mice (6). In this study, we examined the effects of Cd on gene expression of Ube2d family and protein levels of p53 in the liver of mice exposed to Cd for 12 months. It was also examined the effects of Cd on accumulation of p53 protein and occurrence of apoptosis with 5 μm serial sections of the kidney and liver of mice exposed to Cd for 12 months. Although gene expression of Ube2d family in the kidney of mice was significantly decreased by chronic Cd...
exposure (6), that of Ube2d family in the liver of mice was slightly decreased by chronic Cd exposure (Fig. 1A). In a similar way of changes of gene expression of Ube2d family by Cd in the kidney and liver of mice, chronic Cd exposure increased p53 protein levels in the kidney of mice (6); however, hepatic p53 protein levels of mice chronically exposed to Cd was same as that of control mice (Fig. 1B). The difference in response of the Ube2d family gene to Cd exposure in the kidney and liver of mice may have affected the level of protein expression of p53 in the two tissues.

Various proteins involved in UPS have been found to be associated with Cd toxicity in human-origin cultured cells (12,13) in addition to the Ube2d family. Therefore, UPS-related proteins other than the Ube2d family may be involved in Cd-toxicity in the liver or other tissues. This study demonstrated that chronic exposure to Cd triggered p53 dependent apoptosis in the kidney but not the liver. Indeed, p53 at gene and protein level was not regulated by Cd in HepG2 human hepatoma cell lines (14). The entire spectrum of p53 action is likely complex and cell type specific, as it has been implicated in both promoting and inhibiting apoptosis in multiple cell types (15). This study implicates that by inducing apoptosis with Cd in the liver as well as the kidney of mice, differences between the two tissues in their apoptosis-related gene response to Cd may be apparent.

Previous studies reported that Cd elicits toxic effects in microvascular endothelial cells and astrocytes of mice and rats. Cd induces apoptosis through the activation of mitogen activated protein kinase (MAPK) activity in mouse microvascular endothelial cells (16). Moreover, Cd stimulates the expression of intracellular adhesion molecule-1 (ICAM-1), an indicator of blood-brain barrier (BBB) injury in mouse microvascular endothelial cells (17). Cd induces release intracellular Ca2+ level and oxidative stress in rat astrocytes (18,19). We demonstrated that Cd increased p53 protein in human brain microvascular endothelial cells (Fig. 4C). Collectively, these findings suggest that Cd may be associated with neurological disorders.

Our recent study established that Cd changed the activities of various transcription factors in NRK-52E cells (20). A further study found that gene expression of each member of the Ube2d family is dependent on the activity of a transcription factor affected by Cd (on submission). This present study demonstrated that Cd-induced change of gene expression of the Ube2d family varies between tissues and between cell lines. Therefore, the effects of Cd on the activities of transcription factors may be responsible for differences in gene expression of the Ube2d family in different tissues or cell lines.

**CONFLICT OF INTEREST**

Authors declare that they are no conflicts of interest.

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