Effects of Co-Administration of Monomethylaminoantipyrine and Cobaltous Chloride on Hepatic Glutathione Level and Glutathione-Related Enzyme Activities in Rats

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Abstract—Concurrent administration of monomethylaminoantipyrine (MAA) and CoCl₂ caused a significant decrease of hepatic reduced glutathione and oxidized glutathione levels. Furthermore, the increase of glutathione S-transferase activity by combined treatment resulted in the decrease of Se-dependent glutathione peroxidase activity.

We previously reported that treatment of rats with MAA caused a significant increase of hepatic gamma-glutamyltranspeptidase (r-GTP) activity (1). Later, we revealed that co-administration of MAA and CoCl₂ enhanced the MAA-induced increase of hepatic r-GTP activity in rats (2). Such a marked increase of hepatic r-GTP by co-administration of MAA and CoCl₂ is thought to change glutathione level and related enzymatic activity.

The present study was therefore designed to investigate the effect of CoCl₂ alone and in combination with MAA on glutathione level and glutathione-related enzymes in rat liver.

Materials and Methods

Animals and drug administration: Male Sprague-Dawley rats weighing 140 to 150 g were used. They were housed in groups of 3 animals on wire-bottomed stainless steel cages and freely given water and commercial laboratory chow CE-2 (Clea Japan Inc., Tokyo, Japan). A 6:00 a.m.–6:00 p.m. light cycle was maintained. Rats were divided into four groups. The first group received oral administration of 5% gum arabic solution as a control. The second group received oral administration of MAA (563.7 mg/kg, equimolar to 600 mg/kg of aminopyrine) for 3 consecutive days. The third and fourth groups received CoCl₂ alone (20 mg/kg, subcutaneously) and in combination with MAA (563.7 mg/kg) for 3 consecutive days, respectively. CoCl₂ was administered to rats 2 hr prior to MAA administration. Animals were killed at 9:00 a.m. to 10:00 a.m. Animals were sacrificed by decapitation 24 hr after the last administration of MAA, and the hepatic homogenate and cytosolic fractions were prepared as reported previously (1).

Chemicals: r-Glutamyl-p-nitroanilide, GSH and GSSG were purchased from Sigma Chemical Co., St. Louis, MO, U.S.A. CoCl₂, glycylglycine, p-dimethylaminocinnamaldehyde, 1-chloro-2,4-dinitrobenzene and tris (hydroxymethyl)aminomethane were from Wako Pure Chemical Industries, Ltd., Tokyo, Japan. MAA was prepared according to the method of Morita (3). Other reagents were of analytical grade.

Assay methods: Individual contents of reduced glutathione (GSH) and oxidized glutathione (GSSG) were determined by the method of Beutler (4). Glutathione S-transferase (EC 2.5.1.18) activity was assayed by the method of Habig et al. (5). Glutathione peroxidase (GSH-Px) activity was measured by the method of Flohe and Gunzler (6). R-GTP activity was assayed by the method of Morita et al. (7). The results were expressed as mean ± S.E. Statistical significance was assessed by Student's t-test.
glutathione (GSSG) were measured by the fluorescence assay procedure of Hissin and Hilf (4). Glutathione (GSH) S-transferase activity was measured with 1-chloro-2,4-dinitrobenzene (340 nm) as a substrate, according to Habig et al. (5). Glutathione peroxidase activity of the cytosol was measured spectrophotometrically by the procedure of Reddy et al. (6). Glutathione reductase activity was measured by the method of Massey and Williams (7). γ-GTP activity of rat serum was measured by the method of Igarashi et al. (8). The activities of glutamate oxaloacetic transaminase (GOT) and glutamate pyruvate transaminase (GPT) were determined by using the commercial reagent kits of Iatron Co., Ltd., Tokyo, Japan (9, 10). Protein concentration was determined by the method of Lowry et al. (11).

Student’s t-test was used to establish significant differences in mean values between the control and treated groups. The data were subjected to 2×2 factorial analysis of variance.

Results

Effects of MAA alone and in combination with CoCl₂ on the weights of body and liver, on the activities of serum γ-GTP, GPT and GOT, and on the amounts of water and food intakes: As shown in Table 1, repeated administration of CoCl₂ alone or MAA and CoCl₂ for 3 days significantly decreased the body weight. Moreover, administration of MAA alone caused an increase in the liver weight, while administration of CoCl₂ alone led to a decrease. On the other hand, simultaneous administration of MAA and CoCl₂ resulted in a significant increase of serum γ-GTP and GOT activities when compared to the control, but not in GPT activity. In addition, repeated administration of CoCl₂ alone for 3 days resulted in a significant decrease in the amounts of water and food intakes when compared with the control, but the amounts of these intakes remained unchanged after treatment with MAA alone. Moreover, a significant decrease of the amount of food intake was observed by combined administration of MAA and CoCl₂.

Effects of MAA alone and in combination with CoCl₂ on hepatic glutathione levels of rat liver: Table 2 indicates the changes in GSH and GSSG levels of rat liver after administration of either MAA or CoCl₂ alone and combined administration of both drugs. The GSH and GSSG levels remained unchanged after treatment with MAA or CoCl₂ alone. However, a significant decreases of GSH and GSSG levels were observed by combined administration of MAA and CoCl₂.

Effects of MAA alone and in combination with CoCl₂ on the activities of GSH S-transferase and GSSG reductase: As shown in Table 3, a single administration of MAA or CoCl₂ resulted in the increase of GSH S-transferase activity. Moreover, combined administration of MAA and CoCl₂ caused a marked increase of GSH S-transferase activity. On the other hand, treatment of rats with MAA or CoCl₂ alone caused a significant increase in GSSG reductase activity. Furthermore, co-administration of MAA and CoCl₂ also increased the activity of GSSG reductase (Table 3).

Effects of MAA alone and in combination with CoCl₂ on the activities of glutathione peroxidase: Table 4 shows the effect of a single administration of MAA or CoCl₂ and combined administration of both drugs on glutathione peroxidase activities of rat liver. Glutathione peroxidase activity with H₂O₂ as a substrate was significantly decreased by combined treatment. However, the activity towards cumene hydroperoxide was not changed.

Discussion

The previous paper (2) demonstrated that the induction of hepatic γ-GTP by MAA treatment was remarkably enhanced by pretreatment with CoCl₂, by which hepatic microsomal drug metabolism and microsomal content of cytochrome P-450 were diminished.

Prohaska and Ganther (12) reported that the selenium-independent GSH peroxidase (non-Se GSH-Px) activity was associated with the GSH S-transferase and with both the Ya and Yc monomers in the B₁ and B₂ forms of the transferase (13, 14). On the other hand, selenium-dependent GSH peroxidase (Se GSH-Px) activity also is mainly present in the cytosol (6), and GSH-Px appeared to
### Table 1. Effects of treatment of rats with MAA, CoCl₂ and the combination of MAA and CoCl₂ for 3 days on the weights of body and liver, on the activities of serum γ-GTP, GPT and GOT, and on the amounts of water and food intakes

| Treatment           | Body weight (g) | Liver weight (g) | γ-GTP (mU/ml) | GPT (Karmen) | GOT (U/ml) | Water intake (ml) | Food intake (g) |
|---------------------|-----------------|------------------|---------------|--------------|------------|-------------------|-----------------|
| Control (7)         | 174.9±3.6       | 9.26±0.38        | 1.24±0.24     | 27.9±1.58    | 72.57±5.32 | 38.4±1.4          | 21.6±0.5        |
| MAA alone (7)       | 162.7±3.9*      | 11.17±0.46**     | 1.97±0.49     | 33.3±5.59    | 76.16±10.21| 42.0±1.1          | 20.4±0.1        |
| CoCl₂ alone (7)     | 131.3±2.0***    | 7.39±0.22**      | 1.87±0.25     | 26.2±1.76    | 88.21±6.78 | 26.0±1.1***       | 12.4±0.1***     |
| MAA+CoCl₂ (7)       | 130.6±3.6***    | 9.76±0.34        | 4.85±0.52***  | 39.1±5.11    | 121.79±17.53*| 38.2±2.7          | 8.7±0.3***      |

Data in the table represent the mean±S.E. Numbers in parenthesis are the numbers of animals used. *P<0.05, **P<0.01, ***P<0.001 vs. control. The statistical significance was determined by using analysis of variance (2×2 factorial): (γ-GTP), MAA, F(1,24)=27.12, P<0.01; CoCl₂, F(1,24)=27.08, P<0.01; MAA+CoCl₂, F(1,24)=9.70, P<0.01; (GPT), MAA, F(1,24)=7.18, P<0.01; CoCl₂, F(1,24)=0.34, N.S; MAA+CoCl₂, F(1,24)=1.22, N.S; (GOT), MAA, F(1,24)=3.42, N.S; CoCl₂, F(1,24)=9.11, P<0.01; MAA+CoCl₂, F(1,24)=1.94, N.S.

### Table 2. Effects of MAA alone and in combination with CoCl₂ on hepatic reduced and oxidized glutathione levels and glutathione redox ratio in rats

| Treatment           | GSH nmole/mg protein | GSH µmole/g liver | GSSG nmole/mg protein | GSSG µmole/g liver | [GSSG]/[GSH] molar ratio |
|---------------------|----------------------|-------------------|-----------------------|--------------------|-------------------------|
| Control (8)         | 39.30±1.12           | 5.77±0.33         | 5.47±0.45             | 0.81±0.09          | 0.14±0.01               |
| MAA alone (8)       | 31.46±4.30           | 4.88±0.65         | 4.04±0.52             | 0.63±0.08          | 0.13±0.01               |
| CoCl₂ alone (8)     | 43.68±2.65           | 6.54±0.28         | 5.12±0.32             | 0.77±0.05          | 0.12±0.01               |
| MAA+CoCl₂ (8)       | 15.92±1.84***        | 2.47±0.28***      | 2.55±0.37***          | 0.40±0.06**        | 0.17±0.03               |

Data in the table represent the mean±S.E. Numbers in parenthesis are the numbers of animals used. **P<0.01, ***P<0.001 vs. control. The statistical significance was determined by using analysis of variance (2×2 factorial): (GSH), (nmole/mg), MAA, F(1,28)=45.23, P<0.01; CoCl₂, F(1,28)=4.44, P<0.05; MAA+CoCl₂, F(1,28)=14.15, P<0.01; (µmole/g), MAA, F(1,28)=37.53, P<0.01; CoCl₂, F(1,28)=4.07, N.S; MAA+CoCl₂, F(1,28)=15.44, P<0.01; (GSSG), (nmole/mg), MAA, F(1,28)=24.49, P<0.01; CoCl₂, F(1,28)=5.18, P<0.05; MAA+CoCl₂, F(1,28)=1.98, N.S; (µmole/g), MAA, F(1,28)=15.02, P<0.01; CoCl₂, F(1,28)=3.49, N.S; MAA+CoCl₂, F(1,28)=1.74, N.S.
Table 3. Effects of MAA alone and in combination with CoCl₂ on the activities of glutathione S-transferase and glutathione reductase in rat liver

| Treatment     | Glutathione S-transferase | Glutathione GSSG reductase |
|---------------|---------------------------|---------------------------|
|               | μmoles/mg protein         | μmoles/g liver            | nmoles/mg protein         | μmoles/g liver |
| Control       | 0.919±0.09                | 71.26±4.71                | 50.94±7.99                | 3.98±0.62     |
| MAA alone     | 1.715±0.10***             | 129.66±6.01***            | 146.61±12.26***           | 11.06±0.91*** |
| CoCl₂ alone   | 1.217±0.09*               | 89.39±5.17*               | 119.34±11.69***           | 9.51±0.92**   |
| MAA+CoCl₂     | 2.168±0.13***             | 178.70±9.12***            | 141.36±16.27***           | 11.69±1.38*** |

Data in the table represent the mean±S.E. Numbers in parenthesis are the numbers of animals used. *P<0.05, **P<0.01 vs. control. The statistical significance was determined by using analysis of variance (2×2 factorial): (GSH S-transferase), (μmoles/mg), MAA, F(1,24)=72.91, P<0.01; CoCl₂, F(1,24)=13.50, P<0.01; MAA+CoCl₂, F(1,24)=0.58, N.S; (μmoles/g), MAA, F(1,24)=135.80, P<0.01; CoCl₂, F(1,24)=28.08, P<0.01; MAA+CoCl₂, F(1,24)=5.95, N.S; (GSSG reductase); (nmoles/mg), MAA, F(1,28)=27.78, P<0.01; CoCl₂, F(1,28)=7.99, P<0.01; MAA+CoCl₂, F(1,28)=10.87, P<0.01; (μmoles/g), MAA, F(1,28)=26.91, P<0.01; CoCl₂, F(1,28)=11.90, P<0.01; MAA+CoCl₂, F(1,28)=7.55, P<0.05.

Table 4. Effects of MAA alone and in combination with CoCl₂ on hepatic glutathione peroxidase activities in rat liver

| Treatment     | Glutathione peroxidase |
|---------------|------------------------|
|               | Hydrogen peroxide (0.25 mM) | Cumene hydroperoxide (1.2 mM) |
|               | μmoles/min/ mg protein | μmoles/min/ g liver | μmoles/min/ mg protein | μmoles/min/ g liver |
| Control       | 0.288±0.03             | 22.46±1.79          | 0.440±0.02             | 34.73±2.23        |
| MAA alone     | 0.245±0.01             | 18.43±0.85          | 0.489±0.02             | 37.06±2.00        |
| CoCl₂ alone   | 0.274±0.03             | 24.76±3.01          | 0.482±0.04             | 38.29±2.40        |
| MAA+CoCl₂     | 0.198±0.01**           | 16.39±0.95*         | 0.450±0.02             | 37.23±1.59        |

Data in the table represent the mean±S.E. Numbers in parenthesis are the numbers of animals used. *P<0.05, **P<0.01 vs. control. The statistical significance was determined by using analysis of variance (2×2 factorial): (Hydrogen peroxide), (μmoles/min/mg), MAA, F(1,28)=9.39, P<0.01; CoCl₂, F(1,28)=0.31, N.S.; MAA+CoCl₂, F(1,28)=1.75, N.S.; (μmoles/min/g), MAA, F(1,28)=14.52, P<0.01; CoCl₂, F(1,28)=0.01, N.S.; MAA+CoCl₂, F(1,28)=1.78, N.S.; (Cumene hydroperoxide), (μmoles/min/ mg), MAA, F(1,28)=0.41, N.S.; CoCl₂, F(1,28)=0.01, N.S.; MAA+CoCl₂, F(1,28)=3.45, N.S.; (μmoles/min/g), MAA, F(1,28)=0.12, N.S.; CoCl₂, F(1,28)=1.00, N.S.; MAA+CoCl₂, F(1,28)=0.83, N.S.

play a major role in the decomposition of H₂O₂ formed in the cytosol (15). As shown in Tables 3 and 4, the increase of GSH S-transferase activity by combined administration of MAA and CoCl₂ resulted in a decrease of the Se-dependent GSH-Px activity, but not the Se-independent one. Our results may indicate that generation of H₂O₂ during cytochrome P-450-mediated drug oxidation is relatively poor. This can be partly explained by the relative degradation of the hepatic Se-dependent GSH-Px activity.
which, in turn, caused a decrease of GSSG content, and concomitant increases in the activities of GSH S-transferase and GSSG reductase were observed.

Combined administration of MAA and CoCl$_2$ also caused a marked increase of serum $\gamma$-GTP activity, but no significant increase in hepatic GPT activity was observed (Table 1). Serum $\gamma$-GTP originates from liver (16), and glutathione contents in the liver is modulated by $\gamma$-GTP which is the only known enzyme capable of cleaving the $\gamma$-glutamyl bond of GSH and GSSG possessing a $\gamma$-glutamyl moiety.

As shown in Table 2, combined administration of MAA and CoCl$_2$ decreased GSH and GSSG contents. Furthermore, Table 1 shows a significant decrease of the body weight and decreased amount of food intake when MAA plus CoCl$_2$ was administered to rats. Tateishi et al. (17) reported that glutathione level in the liver is closely related to nutritional conditions. In these experiments, the depletion of hepatic GSH and GSSG appears to be partly due to the insufficient food intake in rats.

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