Effect of a Large Dose of Di (2-ethylhexyl) phthalate (DEHP) on Hepatic Peroxisome in Cynomolgus Monkeys (Macaca Fascicularis)

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Abstract: To elucidate the effect of a large dose of di (2-ethylhexyl) phthalate (DEHP), a plasticizer and peroxisome proliferator-activated receptor-α (PPARα) agonist, on hepatic peroxisomes, we orally administered 1,000 mg/kg/day, once daily, to 3 male and 4 female cynomolgus monkeys for 28 days consecutively. Light-microscopic and electron microscopic examinations of the liver were carried out in conjunction with measurement of the hepatic fatty acid β-oxidation system (FAOS), carnitine acetyltransferase (CAT) and carnitine palmitoyltransferase (CPT) activities, which are peroxisomal and/or mitochondrial enzyme activities. Electron microscopically, enlargement of the mitochondria was observed with lamellar orientation of the cristae along the major axis. Although the number of peroxisomes showed a tendency to increase when compared with those in a biopsied specimen before treatment, no abnormality in morphology was observed. A slight increase in CPT activity was noted at termination. No changes were noted in hepatic FAOS or CAT activity. In conclusion, although repeated oral treatment of cynomolgus monkeys with a large dose of DEHP induced a subtle increase in the numbers of peroxisomes with slight enlargements of the mitochondria, this low-sensitivity response to peroxisome proliferators in cynomolgus monkeys was considered to be closer to the response in humans than that in rodents. (J Toxicol Pathol 2010; 23: 75–83)

Key words: cynomolgus monkey, Di (2-ethylhexyl) phthalate (DEHP), hepatocyte, mitochondria, peroxisome

Introduction

Species differences in response to peroxisome proliferators including fibrates as hypolipidemic agents are well-known¹–⁴. Generally, rodents such as mice and rats exhibit a high susceptibility to these peroxisome proliferators with consequent hepatomegaly and hepatocarcinogenesis¹–⁴. However, nonhuman primates showed no such sensitivity⁵,⁶ according to the report of Reddy et al.; on the other hand, the fibrate derivative ciprofibrate evoked hepatic peroxisome proliferation in cynomolgus and rhesus monkeys, pigs, cats and chickens. Similarly, Hoivik et al.⁷ confirmed that treatment of cynomolgus monkeys with high doses of fenofibrate or ciprofibrate caused hepatic peroxisome proliferation in a dose-dependent manner. Therefore, it was considered that nonhuman primates would be likely to possess a proliferative reaction, although it would be a minor response.

Di (2-ethylhexyl) phthalate (DEHP), a plasticizer and its metabolites are peroxisome proliferator-activated receptor-α (PPARα) agonist, has previously been shown to induce peroxisomal proliferation in the livers of rats when administered at 100 mg/kg/day for 14 and 21 days⁸,¹⁰. In contrast, it has been reported that DEHP did not provoke such alterations in the livers of marmosets and cynomolgus monkeys when administered at 2,500 mg/kg/day for 13 weeks¹¹ and at 500 mg/kg/day for 14 or 21 days⁹,¹², respectively. Accordingly, cynomolgus monkeys were not treated with a higher dose of DEHP in these studies. In our previous study¹³, repeated oral treatment of cynomolgus monkeys with a large dose (1,000 or 2,500 mg/kg/day) of DEHP showed a tendency to increase peroxisome proliferation in hepatocytes; however, a small number of animals was used without any corresponding control. Additionally, at 1,000 mg/kg/day of DEHP, body weight
deceased by around 10%, and enlargement of the mitochondria with lamellar orientation of the cristae along the major axis was observed in two females. A dosage level of 2,500 mg/kg/day was not considered to be feasible because the body weight decreased by approximately 40% and hepatic atrophy was observed in one female. These findings implied that an appropriate evaluation could not be carried out, and 1,000 mg/kg/day was thought to be the upper limit in cynomolgus monkeys. In the present investigation, therefore, to elucidate whether a high dose of DEHP evokes peroxisomal proliferation with mitochondrial alterations in cynomolgus monkeys, we orally administered a high dose (1,000 mg/kg/day) of DEHP to both sexes for 28 days consecutively.

Materials and Methods

The present study was approved by the Institutional Animal Care and Use Committee and was conducted in accordance with the bylaws of the committee.

Chemicals

Di (2-ethylhexyl) phthalate (DEHP, Lot No. 4YNPE) was purchased from Tokyo Chemical Industry Co., Ltd. (Tokyo, Japan). All other chemicals and reagents were of the highest grade available from commercial sources, unless otherwise stated.

Animals and housing conditions

Six male and seven female cynomolgus monkeys (Macaca fascicularis) that were 3- to 5-year-olds, purpose-bred and from China were used. They were housed individually in stainless steel cages (680 mm D × 620 mm W × 770 mm H) with an artificial lighting cycle of 12 hr (06:00 to 18:00), a temperature range of 23 to 29°C, a humidity range of 35 to 75% and 15 ventilation cycles/hour. Approximately 108 g (12 g × 9 pieces) of pellet food (Teklad Global Certified 25% Protein Primate Diet, Harlan Sprague Dawley Inc., Indianapolis, IN, USA) was provided to each animal once daily. The remaining food was removed before dosing on the following day. Drinking water was available ad libitum from an automatic supply system.

During a 4-week-acclimation period, drinking water (15 mL/kg) was administered by gavage to each animal daily from 3 days before the initiation of dosing in the same manner as test article administration to acclimate the animal to the dosing procedure. The day before the initiation of dosing was designated as day minus 1 (day –1), the week before the initiation of dosing was designated as week minus 1 (week –1) and the first day of dosing was designated as day 1.

Liver biopsy

Liver biopsy for electron microscopy was conducted for all animals in the first week of the acclimation period (day –29, –26 or –25). A portion of liver tissue (approximately 0.5 g) was removed with an electric knife (MESU-150, Mizuho Ikakogyo Co. Ltd., Tokyo, Japan) under ketamine hydrochloride anesthesia (50 mg/mL, 0.2 mL/kg, Kamud Drugs Pvt. Ltd., Navi Mumbai, India).

Experimental design

DEHP was administered orally once daily to three males and four females for 28 days consecutively at a dose level of 1,000 mg/kg/day by a nasogastric catheter (Size: 10Fr, Nipro Corporation, Osaka, Japan). DEHP was dissolved in corn oil at concentrations of 200 mg/mL. Three males and three females receiving corn oil (5 mL/kg/day) in the same manner as the DEHP group served as the vehicle control (Table 1).

During the dosing period, clinical signs were observed three times daily. Body weight was measured twice a week. Food consumption was measured daily. For food consumption, weekly mean values were calculated as g/animal/day.

Clinical pathology

Hematology and blood chemistry were conducted twice during the acclimation period and once on day 26 of dosing. In blood, the erythrocyte count, leukocyte count, hematocrit value, hemoglobin concentration, reticulocyte ratio and differential leukocyte count were determined using a hematology system (ADVIA120: Siemens Healthcare Diagnostics, Tarrytown, NY, USA). In serum, aspartate aminotransferase, alanine aminotransferase, alkaline phosphatase, lactate dehydrogenase, creatine phosphokinase, total bilirubin, total protein, albumin, globulin, albumin/globulin ratio, total cholesterol, triglycerides, glucose, blood urea nitrogen, creatinine, inorganic phosphorus, calcium, sodium, potassium and chloride were quantified using an automatic analyzer (JCA-BMR: JEOL Ltd., Tokyo, Japan).

Toxicokinetics

Approximately 0.5 mL of blood was collected from the
femoral vein of each animal including those in the vehicle control group using a syringe containing sodium heparin prior to dosing and 1, 2, 4, 6 and 24 hr after dosing on day 26. The blood samples were immediately cooled on ice and centrifuged at 4°C at 1,710 × g for 15 min to obtain plasma. The plasma was stored frozen at –10°C or below until it was analyzed.

Concentrations of DEHP and its first metabolite, mono (2-ethylhexyl) phthalate (MEHP), in plasma were determined by a high-performance liquid chromatography (HPLC) system with a UV detector (Alliance#2795, Waters Corporation, Milford, MA, USA). The area under the plasma concentration-time curve until 24 hr after the 26th dosing (AUC 0-24h) for MEHP was calculated by the linear trapezoidal rule.

**Necropsy**

All animals were euthanized by exsanguination under sodium pentobarbital anesthesia (64.8 mg/mL, 0.4 mL/kg, Tokyo Chemical Industry Co., Ltd., Tokyo, Japan) 24 hr following the final dosing. At necropsy, external appearance, internal organs and tissues were examined macroscopically. The livers were removed and weighed using an electronic balance (HF-3000, A & D Co., Ltd., San Jose, CA, USA). The relative liver weight was calculated using an electronic balance (HF-3000, A & D Co., Ltd., San Jose, CA, USA). The relative liver weight was calculated from the body weight on the day of necropsy. Liver tissue was collected for measurement of the below-stated peroxisomal enzyme activities and light and electron microscopy.

**Measurement of hepatic peroxisomal and mitochondrial enzyme activities**

The right lobe of the liver was perfused with ice-cold physiological saline, frozen immediately with liquid nitrogen and stored in a freezer at –70°C or below before being assayed. All procedures described below were conducted under ice-cold conditions. After the collected liver tissue was cut into small portions with scissors, an amount of Tris buffer containing 1 mmol/L EDTA (pH 7.4) equal to nine times the liver tissue weight was added. The tissue was homogenized to prepare a 10% hepatic homogenate.

The activities of fatty acid ß-oxidation system (FAOS), carnitine acetyltransferase (CAT) and carnitine palmitoyltransferase (CPT) in the 10% hepatic homogenates were measured by a spectrophotometer (U-3010, Hitachi, Ltd., Tokyo, Japan) using a previously reported method. Each activity is reported as μmol/min/g liver.

**Light microscopy**

After excision at necropsy, the liver was fixed in 10% neutral buffered formalin and embedded in paraffin wax. Approximately 4-μm paraffin sections were stained with hematoxylin and eosin (HE) and examined microscopically.

**Electron microscopy**

The portions of liver tissue obtained by biopsy before treatment and after exsanguination at necropsy were fixed in 2.5% glutaraldehyde and 1% osmium tetroxide and embedded in epoxy resin (Quetol 812). Ultra-thin specimens (0.05 to 0.1 μm) were stained with uranyl acetate and lead citrate and examined by transmission electron microscopy (JEM-1200EX, JEOL, Ltd., Tokyo, Japan).

The numbers of peroxisomes and mitochondria in the hepatocytes in the centrilobular or periportal areas in 40 images of liver tissue were counted at 6,500 × magnification for each animal. The total area of the 40 images was approximately 3,300 μm².

**Statistical analysis**

Quantitative data were analyzed statistically to compare the vehicle control and DEHP groups. Additionally, the data from electron microscopy were analyzed to compare the values from before and after repeated treatment. They were first analyzed for homogeneity of variance by Bartlett’s test. When the variance was homogeneous, a one-way analysis of variance was applied. When the result was significant, Dunnett’s test was applied. When the variance was heterogeneous by Bartlett’s test, the data were analyzed by the Kruskal-Wallis test. When the result was significant, a Dunnnett-type test was applied. The MUSCOT statistical analysis software (Yukms Co., Ltd., Tokyo, Japan) was used for these statistical analyses at a significance level of 5%.

**Results**

**Pathological findings in the liver**

In macroscopic and light microscopic examinations, no abnormality related to DEHP treatment was observed.

In regard to organ weights, no statistically significant differences were noted the liver; however, slight increases both in absolute and relative weights were noted in two females (Animal Nos. 10 and 13) that showed no body weight loss in the DEHP group (Table 2). The increase ratios were 22 or 24% for the relative weight when compared with the mean values in the corn oil group. The relative weight of 1 other female (Animal No. 11) in the DEHP group increased; however, the absolute weight did not increase, and body weight loss was observed in this animal.

Electron-microscopically, enlargement of the mitochondria with lamellar orientation of the cristae along the major axis was observed in one female (Animal No. 13) in the DEHP group (Fig. 1). Some mitochondria showed elongated enlargement.

In the DEHP-treated females, the numbers of peroxisomes in both centrilobular and periportal areas increased significantly (***p<0.01) at the end of the dosing period compared with those before treatment (Fig. 2). The individual increase ratios were 33.3% to 111.4% and 38.5% to 81.3% in the centrilobular and periportal areas, respectively; however, no significant changes were noted in mitochondria (Fig. 3). In the DEHP-treated males, no statistically significant differences in the numbers of peroxisomes or mitochondria were noted; however, the
Peroxisome Proliferation in the Liver Induced by DEHP

numbers of peroxisomes in the periportal area showed a tendency to increase; i.e., the mean value was higher than those before treatment or in the corn oil group (Fig. 2, 3).

**Hepatic peroxisomal and mitochondrial enzymes**

The activity of the mitochondrial enzyme CPT increased significantly in the males treated with DEHP, at 1.6 fold higher (2.043 units for corn oil versus 3.313 units for DEHP). No statistically significant differences were observed in the FAOS or CAT activities (Fig. 4).

**Toxicokinetics**

Extremely low concentrations (2.0–9.0 μg/mL) of the parent compound DEHP in plasma were detected sporadically at 1 to 6 hr after dosing in two males and two females in the DEHP group. Meanwhile, its first metabolite, MEHP, increased with time after dosing in all animals in the DEHP group. The plasma MEHP concentration reached a maximum at 4 to 6 hr after dosing. The individual Cmax values ranged from 39.2 to 70.5 μg/mL, and the mean Cmax value in the DEHP group was 826.6 ± 271.2 μg·hr/mL.

**Clinical observations and laboratory tests**

In regard to clinical signs, as abnormalities related to DEHP treatment, soft stool and/or diarrhea were observed in all males and females almost every day. Vomiting was observed in one male and one female on day 3 or 26.

In regard to body weight, a decrease was observed in two males and one female (Animal Nos. 8, 9 and 11) in the DEHP group at the end of the dosing period when compared with the first day of dosing (Fig. 5). The decrease ratios in these animals were 10.0%, 15.5% and 24.2%. No abnormalities were observed in any other animal.

In regard to food consumption, the mean values of food consumption throughout the dosing period were decreased in one male and one female (Animal Nos. 9 and 11) in the DEHP group. The decrease ratios of the mean values in these animals were 43% and 44%, respectively, when compared with the mean values in weeks –2 to –1. No abnormalities were observed in any other animal.

In regard to blood chemistry, serum lipids (total cholesterol or triglyceride) did not decrease in the DEHP group (Table 3). An increase in triglycerides was noted in 1

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**Table 2.** Organ Weights of the Livers of Cynomolgus Monkeys Orally Administrated Corn Oil or DEHP at 1,000 mg/kg/day for 28 Consecutive Days

| Group | Male | Female |
|-------|------|--------|
|       | Animal No. (g) | Animal No. (g) |
| Corn oil |   |   |
| 1 | 70.6 | 4 | 59.4 |
| 2 | 70.4 | 5 | 44.7 |
| 3 | 79.4 | 6 | 61.5 |
| Mean ± SD | 73.47 ± 5.14 | Mean ± SD | 55.20 ± 9.15 |
| DEHP |   |   |
| 1 | 81.1 | 10 | 66.2 |
| 8 | 65.3 | 11 | 57.2 |
| 9 | 68.0 | 12 | 48.9 |
| – | – | 13 | 71.9 |
| Mean ± SD | 71.47 ± 8.45 | Mean ± SD | 61.05 ± 10.11 |

Not significantly different from the corn oil group.

| Group | Male | Female |
|-------|------|--------|
|       | Animal No. (g/kg) | Animal No. (g/kg) |
| Corn oil |   |   |
| 1 | 16.2 | 4 | 22.0 |
| 2 | 17.8 | 5 | 19.0 |
| 3 | 20.7 | 6 | 18.9 |
| Mean ± SD | 18.23 ± 2.30 | Mean ± SD | 19.98 ± 1.75 |
| DEHP |   |   |
| 7 | 19.7 | 10 | 24.4 |
| 8 | 20.2 | 11 | 25.8 |
| 9 | 21.1 | 12 | 20.0 |
| – | – | 13 | 24.7 |
| Mean ± SD | 20.34 ± 0.71 | Mean ± SD | 23.74 ± 2.53 |

Not significantly different from the corn oil group.
female (Animal No. 11) in the DEHP group. In regard to hematology, no abnormality related to DEHP treatment was observed.

**Discussion**

Statistically significantly increased numbers of peroxisomes in the centrilobular and periportal areas were observed in the DEHP-treated females compared with before treatment. In the DEHP-treated males, no statistically significant differences were noted; however, the numbers of peroxisomes in the periportal area were higher than those before treatment or in the corn oil group. These results suggested that a large dose of DEHP induced an increase in the numbers of peroxisomes. On the other hand, no significant differences were observed when compared with the corn oil group in the numbers of peroxisomes in the males or females. No differences were observed in the peroxisomal enzyme activities in liver homogenates between the DEHP and corn oil groups. These results suggested that the peroxisome proliferation was subtle. In our previous study\textsuperscript{13}, we could not reach a conclusion as to whether or not DEHP induced peroxisome proliferation. We added the
biopsy before treatment and a vehicle control and increased the number of animals in the present study. As a consequence, we were able to conclude that DEHP induced a subtle change in hepatic peroxisome proliferation in the cynomolgus monkeys.

In our previous study, morphological changes of mitochondria, such as enlargement and lamellar orientation of the cristae along the major axis, were observed in 2 female cynomolgus monkeys whose body weights decreased at ratios of 10.7% or 11.7% at 1,000 mg/kg/day of DEHP, and the possibility that malnutrition could have caused the mitochondrial changes was suggested. In the present study, mitochondrial abnormalities were observed in one female; however, body weight loss was not noted in this animal. Mitochondrial changes caused by peroxisome proliferators have previously been reported as follows: DEHP induced increases in both the size and number of mitochondria in hepatocytes in rats, and other PPARα agonists (ciprofibrate and fenofibrate) induced an increase in number...
and elongation of mitochondria in hepatocytes in cynomolgus monkeys. Additionally, peroxisome and mitochondria are closely associated with each other in PPARα agonist-related metabolism. Consequently, there is a high possibility that the mitochondrial changes observed in the present study were caused by effects of DEHP on the hepatocytes. The toxicological significance and mechanism of the mitochondrial changes were unclear in the present study, and other approaches from the standpoint of molecular biology would seem to be required.

CPT is widely known to be a fatty acid oxidation rate-limiting enzyme in the mitochondria. Hepatic CPT activity increased in males treated with DEHP but not in females, and morphological changes of the mitochondria were observed only in females. Thus, the relation between the morphological changes of the mitochondria and increased activity of CPT was unclear. The result in the present study differed from the previously reported results in rats. It is possible that factors other than testosterone had an influence in the cynomolgus monkeys, but this was uncertain.

A number of experiments have suggested that there are species differences in the expression of PPARα and response to PPARα agonists (including differences in receptor activation, peroxisome proliferation and induction of target genes), which are higher in rodents than in other species. Additionally, there is a wide range of receptor

| Group                        | Animal No./Day | Total cholesterol (mg/dL) | Triglyceride (mg/dL) |
|------------------------------|----------------|---------------------------|----------------------|
|                              |                | Day –5*                   | Day 26*              | Day –5*                   | Day 26*              |
| Control (Male)               |                |                           |                      |                      |
| Corn oil                     | 1              | 122                       | 111                  | 59                     | 33                   |
|                              | 2              | 76                        | 79                   | 35                     | 39                   |
|                              | 3              | 137                       | 137                  | 48                     | 40                   |
| Mean                         |               | 111.7                     | 109.0                | 47.3                   | 37.3                 |
| SD                           |               | 31.8                      | 29.1                 | 12.0                   | 3.8                  |
| DEHP (Male) 1,000 mg/kg/day  | 7              | 93                        | 84                   | 26                     | 22                   |
|                              | 8              | 92                        | 93                   | 43                     | 37                   |
|                              | 9              | 74                        | 103                  | 14                     | 34                   |
| Mean                         |               | 86.3                      | 93.3                 | 27.7                   | 31.0                 |
| SD                           |               | 10.7                      | 9.5                  | 14.6                   | 7.9                  |
| Control (Female)             |                |                           |                      |                      |
| Corn oil                     | 4              | 153                       | 157                  | 34                     | 18                   |
|                              | 5              | 129                       | 117                  | 44                     | 49                   |
|                              | 6              | 126                       | 126                  | 29                     | 49                   |
| Mean                         |               | 136.0                     | 133.3                | 35.7                   | 38.7                 |
| SD                           |               | 14.8                      | 21.0                 | 7.6                    | 17.9                 |
| DEHP (Female) 1,000 mg/kg/day| 10             | 156                       | 136                  | 33                     | 35                   |
|                              | 11             | 175                       | 195                  | 52                     | 285†                 |
|                              | 12             | 136                       | 123                  | 27                     | 41                   |
|                              | 13             | 139                       | 137                  | 24                     | 25                   |
| Mean                         |               | 151.5                     | 147.8                | 34.0                   | 96.5                 |
| SD                           |               | 18.0                      | 32.1                 | 12.6                   | 125.8                |

Not significantly different from the corn oil group.

*: The day before initiation of dosing was designated as day –1, and the first day of dosing was designated as day 1.
affinities for different PPARα ligands. The lower reactivity of cynomolgus monkeys to DEHP may be explained by these differences. MEHP is the first metabolite of DEHP and is a PPARα agonist. The first stage in metabolism of DEHP is hydrolysis to MEHP and 2-ethylhexanol. Hydrolysis to MEHP rapidly takes place in the intestine, and further hydrolysis takes place after absorption. Additionally, hydrolysis to MEHP by plasma lipase takes place in blood samples. The DEHP and MEHP concentrations in plasma might not have shown pharmacokinetically true values because there is a possibility that hydrolysis to MEHP took place during the procedure for obtaining plasma and/or its preservation. However, the increased plasma MEHP level revealed sufficient intestinal absorption of DEHP in the present study.

In the present study, we concluded that DEHP induced hepatic peroxisome proliferation in cynomolgus monkeys; however, the degree of increase was very low, hepatomegaly or hepatic proliferation was not observed and the exposure level was extremely high (1,000 mg/kg/day). In contrast, peroxisome proliferation was observed from 100 mg/kg/day in rats. At the same time, our results reconfirmed that peroxisome proliferation was observed from 100 mg/kg/day level was extremely high (1,000 mg/kg/day). In contrast, however, the degree of increase was very low, hepatomegaly hepatic peroxisome proliferation in cynomolgus monkeys; that the primate may be refractory to PPAR-induced proliferation, and there was no remarkable increase in the rodent; that is to say, there was no indication of cell response to fibrates in a manner that is different from the animal.

The response to peroxisome proliferators in cynomolgus monkeys was considered to be closer to that in humans than the response in rodents. However, further investigation into the causes of species differences appears to be necessary for accurate assessments of the risk of peroxisome proliferators for humans from the results in cynomolgus monkeys.

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