Monoester Formation by Hydrolysis of Dialkyl Phthalate Migrating from Polyvinyl Chloride Products in Human Saliva

Tatsuhiro Niino,^{*,a} Tohru Ishibashi,^a Takeshi Itho,^a Senzo Sakai,^a Hazimu Ishiwata,^b Takashi Yamada,^c and Sukeo Onodera^d

^aTokyo Kenbikyo-in Foundation, Center of Food & Environmental Sciences, 44-1 Nihonbashi Hakozaki-cho, Chuo-ku, Tokyo 103-0015, Japan, ^bNational Institute of Health Sciences, 1-18-1 Kamiyoga, Setagaya-ku, Tokyo 158-8501, Japan, ^cJapan Food Additives Association, Nihonbashi-San-Ei-Bldg., 3F 1-3-9 Nihonbashi-Horidome-cho Chuo-ku, Tokyo 103-0012, Japan, and ^dFaculty of Pharmaceutical Sciences, Tokyo University of Science, 12 Ichigaya-Funagawara-machi, Shinjuku-ku, Tokyo 162-0826, Japan

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The migration dialkyl phthalate was tested in volunteers who chewed polyvinyl chloride (PVC) toy products under controlled conditions. The PVC toy samples consisted of ball A containing 100 and 185 mg/g di-*n*-butyl phthalate (DBP) and di-2-ethylhexyl phthalate (DEHP) respectively, and ball B, containing 256 mg/g diisononyl phthalate (DINP). The migration of dialkyl phthalate into simulated saliva was also tested *in vitro* by shaking toy samples. The migration rates of DBP, DEHP and DINP from balls A and B were 11.7, 44.4 and 78.0 $\mu\text{g}/10\text{ cm}^2/\text{hr}$, respectively, *in vivo*, and 339, 315 and 535 $\mu\text{g}/10\text{ cm}^2/\text{hr}$, respectively, *in vitro*. The presence of mono-*n*-butyl phthalate (MBP) and mono-2-ethylhexyl phthalate (MEHP) in saliva collected after chewing ball A was confirmed by GCMS-SIM. Human saliva which collected from volunteers incubated with DBP and DEHP at 37°C over 60 min, hydrolyzed these compounds to their monoesters.

Key words — dialkyl phthalate, monoester, hydrolysis, polyvinyl chloride toy, migration

INTRODUCTION

Dialkyl phthalates (PAEs) are widely used as plasticizers to impart softness and flexibility to normally rigid plastics such as polyvinyl chloride (PVC). Medical devices and toys are often made of PVC, containing the predominant plasticizers, di-2-ethylhexyl phthalate (DEHP) or diisononyl phthalate (DINP).^{1,2)} Some PAEs have induced testicular toxicity and other effects on the male and female

reproductive tract at high dosages in rats and other animals.^{3,4)} Jobling *et al.* showed that di-*n*-butyl phthalate (DBP) binds to the estrogen receptor, displaces the natural ligand from the receptor, then acts as an estrogen agonist rather than an antagonist.⁵⁾ The DEHP and DINP consist of many structurally dissimilar compounds known as peroxisome proliferators that induce tumors in the rodent liver.⁶⁾

Orally-ingested PAE is hydrolyzed to its monoester by endogenous esterases of many tissues, such as the small intestine.^{7,8)} Monoesters are thought to be more hepatocarcinogenic than their parent compound.⁹⁾

The rate of DINP release from PVC toys chewed by children has been measured using two general migration tests. An *in vivo* test in which human volunteers chew PVC toys under controlled conditions was developed by Meuling and Rijk in the Netherlands¹⁰⁾ and by Chen for the U.S. Consumer Product Safety Commission (USCPSC).¹¹⁾ In addition, the effect of shaking or impacting PVC toys in simulated saliva *in vitro* has been tested. The release rates *in vivo* are higher than those obtained *in vitro*. However, the *in vivo* and *in vitro* migration of PAE has not been simultaneously studied except for DINP. Therefore, we examined the *in vivo* and *in vitro* migration of PAE from PVC toys containing DBP, DEHP and DINP. In addition, we examined monoester formation by the hydrolysis of PAE that migrated into saliva from PVC toys.

MATERIALS AND METHODS

Reagents — DBP and DINP were purchased from Kanto Chemical Co., Inc. (Tokyo, Japan) and DEHP from Wako Pure Chemical Ind. (Osaka, Japan). The purity of DBP, DEHP and DINP was over 99.7, 99.5

*To whom correspondence should be addressed: Tokyo Kenbikyo-in Foundation, Center of Food & Environmental Sciences, 44-1 Nihonbashi Hakozaki-cho, Chuo-ku, Tokyo 103-0015, Japan. Tel.: +81-3-3663-9684; Fax: +81-3-3663-9682; E-mail: tatsu-n@jc4.so-net.ne.jp

and 97.9%, respectively. Mono-*n*-butyl phthalate (MBP) and mono-2-ethylhexyl phthalate (MEHP) of over 90% purity were purchased from Tokyo Kasei Kogyo Co., Ltd. (Tokyo, Japan). Trimethylsilyldiazomethane (TMSD) was purchased from Aldrich Chemical Co. (U.S.A.). All other solvents and reagents were of analytical grade, and confirmed as being PAE free. The composition of the simulated saliva according to the British Standard Specification for Safety Harnesses was: 4.5 g sodium chloride, 0.3 g potassium chloride, 0.3 g sodium sulfate, 0.4 g ammonium chloride, 0.2 g urea and 3.0 g lactic acid dissolved in 1000 ml distilled water adjusted to pH 6.5 to 7.0 with 5 M sodium hydroxide.¹²⁾ PAE and monoesters were solid-phase extracted using Oasis hydrophilic lipophilic balance (HLB) cartridges (3 cc) from Waters Co. (Milford, Mass., U.S.A.). Prior to use, their cartridges were washed with 10 ml of methanol, followed by 10 ml of methanol • 0.1% acetate (1 : 9). Glassware was heated at temperatures over 230°C for at least 5 hr before use.

HPLC Analysis — The HPLC conditions were as follows: apparatus, LC-10A (Shimadzu, Kyoto, Japan); column, Inertsil C8-3 (0.46 mm ϕ $\times$ 250 mm); column oven temperature, 40°C; mobile phase, (A) water/acetic acid 200 : 2(w/w), (B) acetonitrile, A : B = 3 : 7; detector, UV (254 nm); injection volume, 10 μ l.

GC/MS Analysis — The gas chromatography (GC) conditions were as follows: column, CBP5 (0.25 mm i.d. $\times$ 50 m, film thickness 0.25 μ m); carrier gas, helium at 2.0 ml/min; column temperature, 90°C(1 min) $\rightarrow$ (40°C/min $\rightarrow$ 290°C; injection port and interface temperature, 290°C. The MS conditions for electron impact ionization of monoalkyl phthalate were as follows: ion energy, 70 eV; ion source temperature, 270°C; selected ion, *m/z* 163 and 149.

PAE Migration Test from PVC Toys — The PVC toy products were ball A that contained 100 mg/g DBP and 190 mg/g DEHP, and ball B that contained 257 mg/g DINP. The balls were punched over a 2.5 $\times$ 3.0 cm area (total surface area was approximately 15 cm²). These are referred to below as toy samples A or B.

Migration test *in vitro*: Rotary shaking migration was examined using a rotary shaker, VR-36 [TAITEC Co. (Saitama, Japan)]. Toy samples were placed in 50 ml glass centrifuge tubes containing 30 ml of simulated saliva. Tubes were placed on the rotary shaker in a incubator at 35°C, and shaken for

15 min at a rate of 300 rpm. The solutions (5 ml) were sampled, and the rest was discarded. Four consecutive 15 min periods of this operation was repeated. The collected samples were mixed with 5 ml of acetonitrile. Migrated PAE in the samples was quantified by HPLC.

Migration test *in vivo*: One female and three male volunteers gently chewed toy samples or control polypropylene (PP) disks PAE-free for 15 min. They repeated this process for four 15 min intervals on toy sample A (four sessions), and were a 5 min break before actively sucking the sample. During each 15 min period, all produced saliva was collected in labeled 50 ml glass centrifuge tubes.^{10,11)} The total volume and pH of produced saliva in each tube were recorded. All saliva samples were diluted to 10 ml with distilled water, mixed with 10 ml of acetonitrile, then sedimented by centrifugation at 3000 rpm for 10 min. This procedure was repeated by the same volunteers on toy sample B 1 week later. Migrated DBP and DEHP in the supernatant was quantified by HPLC.

The amount of monoalkyl phthalate in saliva samples was measured as follows. The supernatant was mixed with 15% sodium chloride and rendered alkaline with 0.1 N sodium hydroxide, then PAE was extracted with *n*-Hexane. Monoesters in the aqueous acetonitrile layer acidified with 0.01 N hydrochloric acid were extracted with 20% dichloromethane • *n*-hexane. The extracted monoesters were methyl-esterified with TMSD as described by Ohfuji et al.¹³⁾ Methyl-esterified monoesters were quantified by GC/MS.

Incubation Test of PAE with Human Saliva — Saliva (0.5 ml) collected from volunteers who chewed PP disks was added to the same volume of simulated saliva and incubated with 50 nmol of DBP or DEHP in 10 μ l of dimethylsulfoxide at 37°C for 5, 15, 30 or 60 min, respectively. The incubated mixture was adjusted to pH 4 with 0.01 N hydrochloric acid, mixed with acetonitrile (1 ml), then centrifuged at 3000 rpm for 10 min. The supernatants were mixed with 8 ml of 0.1% acetic acid and passed through Oasis HLB cartridges. Organic compounds remaining in the column were eluted with 20 ml of acetonitrile. The eluates were evaporated to near dryness, then PAE and monoesters were analyzed by HPLC.

Table 1. Migration Rates of Dialkyl Phthalate (PAE) from PVC Toy Samples

Toy sample	PAE	Contents ^{a)} (mg/g)	Migrated PAE ($\mu\text{g}/10\text{ cm}^2/\text{hr}$)		B/A
			<i>in vivo</i> ^{b)} (A)	<i>in vitro</i> ^{c)} (B)	
Toy ball A	DBP	100	11.7 $\pm$ 9.8	339 $\pm$ 6.9	28.9
	DEHP	185	44.4 $\pm$ 12.3	315 $\pm$ 25.0	6.6
Toy ball B	DINP	256	78.0 $\pm$ 28.9	535 $\pm$ 37.7	6.9

a) PAE contents in toy sample were measured by extraction with acetone using rotaty shaker at 300 rpm for 3 hr. b) These release rates were obtained by *in vivo* migration tests after 4 volunteers chewed toy sample for 60 min (15 min $\times$ 4 sessions). Values are means $\pm$ S.D. ($n = 4$). c) These release rates were obtained by *in vitro* migration tests of simulated saliva shaken at 300 rpm at 35°C for 60 min (15 min $\times$ 4 sessions). Values are means $\pm$ S.D. ($n = 5$).

RESULTS AND DISCUSSION

In Vitro and *In Vivo* Migration of PAE

Table 1 shows the release rates of PAE into human saliva (*in vivo*) and into simulated saliva (*in vitro*) from toy samples. The rates of DBP and DEHP migration into the simulated saliva were 339 and 315 $\mu\text{g}/10\text{ cm}^2/\text{hr}$, respectively (654 $\mu\text{g}/10\text{ cm}^2/\text{hr}$ at the sum total PAEs), and 11.7 and 44.4 $\mu\text{g}/10\text{ cm}^2/\text{hr}$ into the human saliva, respectively (56.1 $\mu\text{g}/10\text{ cm}^2/\text{hr}$ at the sum total PAEs). Those of DINP that migrated into the simulated and human saliva were 535 $\mu\text{g}/10\text{ cm}^2/\text{hr}$ and 78.0 $\mu\text{g}/10\text{ cm}^2/\text{hr}$, respectively. The release rates into saliva simulant were of the order DBP > DEHP > DINP. Although migration into saliva *in vivo* seemed to increase with shorter alkyl chains, about 70% less DBP and DEHP than DINP migrated. The amounts of DBP, DEHP and DINP that migrated *in vivo* were 28.9, 6.6 and 6.9-fold on the ratio of *in vitro* migration. Very little DBP migrated into saliva than DEHP and DINP.

The migration of DINP from PVC products *in vivo* has been tested in the Netherlands¹⁰⁾ and in the U.S.A.¹¹⁾ The Dutch consensus study indicated a mean DINP release rate of 108 $\mu\text{g}/10\text{ cm}^2/\text{hr}$ from samples containing 430 mg/g of DINP which was considerably less than the American CPSC values of 219 $\mu\text{g}/10\text{ cm}^2/\text{hr}$. Generally, release rate and DINP content of the sample did not correlate.¹⁴⁾ Less DINP migrated from toy sample B than that indicated by the Dutch consensus study. Rates of saliva produced by chewing toy samples A and B were 37–92 ml/hr and 38–45 ml/hr, and pH of the saliva was 7.0–7.3 and 6.7–7.6.

MBP and MEHP in saliva collected during the *in vivo* test of the toy sample A were methyl-esterified with TMSD, and analyzed by GC/MS. The methyl-esterified monoesters have a stable base peak at m/z 163 and 149 when analyzed by electron-impact MS. The chemical structure of these ions is

shown in Fig. 1. Selective monitoring ions were used to identify monoesters in saliva extracts. The SIM chromatograms of the extracted saliva in Fig. 2 show the presence of MBP and MEHP indicating their formation through PAE conversion in saliva produced while chewing the toy sample.

Hydrolysis of DBP and DEHP by Saliva to Form

We examined the recovery of PAE and the monoester during the incubation test. Standard DBP, DEHP, MBP and MEHP added to simulated saliva

Fig. 1. GC/MS-SIM Trace of Methyl Esterified MBP and MEHP

1, mono-*n*-butyl phthalate (MBP); 2, mono-2-ethylhexyl phthalate (MEHP). Amount of injected MBP and MEHP after methyl-esterification were 100 ng each.

Fig. 2. GC/MS-SIM Trace of Methyl Esterified Saliva Extract
Saliva was alkalified with NaOH and extracted with *n*-hexane. The lower layer was then acidified with HCl and extracted with 20% dichloromethane • *n*-hexane.

were removed at rates of 96.8%, 83.1%, 99.4% and 99.6% for DBP, DEHP, MBP, MEHP, respectively.

Figures 3 and 4 show the formation of MBP and MEHP by of saliva containing 50 nmol of standard DBP and DEHP. The amounts of DBP and DEHP in the saliva decreased to 95% and 12%, respectively over 60 min at 37°C. MBP was formed from the added DBP within 5 min. The MBP formation accounted for 87% of the added DBP within 60 min. On the other hand, the amount of MEHP slightly increased over 60 min.

In general, PAE is thought to be metabolized to monoesters by esterases in the wall of the small intestine and by pancreatic lipases.^{7,8)} DEHP is rapidly absorbed from the rodent gut, mostly in the form of the monoester because of the rapid hydrolysis of the DEHP by gut lipases.^{15,16)} The present studies showed that MBP and MEHP were formed by the monohydrolysis of DBP and DEHP in human saliva. We surmised that the hydrolysis of PAE in human saliva differs according to the length of the alkyl chains, because more MBP was formed from DBP than MEHP from DEHP. Kayano *et al.* reported that the hydrolysis of PAE by the mouse hepatic esterase

Fig. 3. The Relationship between MBP Formation by DBP Hydrolysis and Incubation Time in the Human Saliva

Saliva obtained from volunteers who had chewed PP disk was mixed with 50 nmol DBP and incubated at 37°C for 60 min. Values are means of duplicate assays. ◇, Amount of DBP. ●, Amount of MBP.

Fig. 4. The Relationship between MEHP Formation by DEHP Hydrolysis and Incubation Time in the Human Saliva

Saliva obtained from volunteers who had chewed PP disks was mixed with 50 nmol DEHP and incubated at 37°C for 60 min. Values are means of duplicate assays. ◇, Amount of DEHP. ●, Amount of MEHP.

has substrate specificity for chained alkyl groups.¹⁷⁾ The MEHP compounds are thought to be more hepatocarcinogenic and teratogenic than DEHP at high dosages.⁹⁾ In conclusion, the results of the present study indicated that infants and toddlers can be exposed to PAE by chewing PVC toys, and that the monoester may be formed through hydrolysis by saliva in the oral cavity.

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