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Assessing the genotoxicity of two commonly occurring byproducts of water disinfection: chloral hydrate and bromal hydrate

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Highlights

- Genotoxicity of chloral hydrate and bromal hydrate investigated *via 3 in vitro* assays
- Chloral hydrate not statistically exhibiting significant genotoxic effects
- Mutagenic activity of bromal hydrate in the *Salmonella* strain TA100
- Significant DNA lesions in CHO cells induced by bromal hydrate
- Higher genotoxicities of brominated DBPs in comparison to chlorinated species

Abstract

Water disinfection treatments result in the formation of disinfection byproducts (DBPs) that have been linked to adverse human health outcomes including higher incidence of bladder and colorectal cancer. However, data about the genotoxicity of DBPs is limited to only a small fraction of compounds. Chloral hydrate (CH) and bromal hydrate (BH) are two trihaloacetaldehydes commonly detected in disinfected waters, but little is known about their genotoxicity, especially BH.

We investigated the genotoxicity of CH and BH using a test battery that includes three *in vitro* genotoxicity assays.

We conducted the Ames test using *Salmonella* bacterial strains TA97a, TA98, TA100 and TA102, and the alkaline comet assay and the micronucleus test both using Chinese hamster ovary cells. We carried out the tests in the absence and presence of the metabolic fraction S9 mix.

CH did not exhibit statistically significant genotoxic effects in any of the three assays. In contrast, BH exhibited mutagenic activity in the *Salmonella* strain TA100 and induced statistically significant DNA lesions in CHO cells as appeared in the comet assay. The genotoxic potential of BH in both assays decreased in the presence of the metabolic fraction S9 mix. BH did not induce chromosomal damage in CHO cells.

Our results show that BH exhibited genotoxic activity by causing mutations and primary DNA damage while CH did not induce genotoxic effects. Our findings highlight concerns about the higher genotoxicity of brominated DBPs in comparison to their chlorinated analogues.

Introduction

One of the most significant public health advances of the twentieth century was the adoption of drinking water disinfection in many countries [1]. This practice has sharply reduced the incidence of infectious diseases such as cholera, typhoid, and dysentery [2,3]. After this dramatic success, disinfection practices have been introduced into swimming pools and other recreational water venues to ensure the elimination of pathogenic microorganisms and the prevention of waterborne disease outbreaks [4]. However, disinfection treatments result in the undesirable formation of chemical contaminants known as disinfection byproducts (DBPs), in consequence

to reactions taking place between disinfectants and organic matter present in water [5,6]. Exposure to DBPs in humans can take place through ingestion of drinking water or inhalation and dermal absorption during showering or swimming [7-10]. Many studies have suggested associations between exposure to DBPs and adverse health effects. Increased incidence of asthma [11], bladder cancer [12,13], and colorectal cancer [14] have been reported. Adverse pregnancy outcomes such as spontaneous abortions [15], stillbirth [16], and fetal growth restriction [17] have also been noted. To date, more than six hundred DBPs including trihalomethanes, haloacids, halonitriles, haloaldehydes, halo ketones, halonitromethanes, haloamines, haloamides, haloalcohols, and halobenzoquinones have been identified in disinfected waters [9,18-23]. Several laboratory-controlled studies have been conducted to evaluate potential toxicities of DBPs providing evidence about cytotoxic, genotoxic, carcinogenic and teratogenic potentials [20, 24-28]. However, the toxicological data are limited to only a small fraction of identified DBPs. In consequence, many DBPs that have been detected in disinfected waters remain with unknown toxicological profiles. Chloral hydrate and bromal hydrate, the hydrated forms of trichloroacetaldehyde and tribromoacetaldehyde respectively, belong to the chemical class of haloacetaldehydes. This class of DBPs has been reported to be one of the most abundant DBP classes by weight [19,25,29,30]. Occurrence studies have shown that the predominant trihaloacetaldehyde in chlorinated waters is chloral hydrate, while bromal hydrate is the predominant trihaloacetaldehyde in chlorinated waters containing high levels of bromide [19,31]. In a recent study, BH was detected as one of the degradation byproducts of benzophenone-3, a UV filter commonly used in sunscreens, in chlorinated swimming pools filled with seawater [32].

Toxicokinetic studies have shown that CH is rapidly absorbed after oral administration, and enters the liver where it undergoes extensive metabolism in rodents [33,34] and in humans [35,36]. Studies of the potential carcinogenicity of CH in mice have demonstrated that it is able to induce hepatocellular adenomas and carcinomas, and exposure to CH has been associated with increases in malignant lymphoma and adenoma of the pituitary gland [37-39]. However, there was still no persuasive evidence to connect chloral hydrate exposure and the development of cancers in humans [40]. CH was also found to induce significant aneugenic effects in mice [41]. Furthermore, micronuclei were produced in germ cells of male mice treated intraperitoneally with CH [42]. CH was also reported to be able to lead to chromosomal loss in mouse spermatids [43] and in human lymphocytes [44]. Nevertheless, most of the investigations incorporated only one or two *in vitro* assays [25,45] and results from genotoxicity assessment of CH remain inconclusive. Concerning BH, although little is known about its toxicity, the U.S. Environmental Protection Agency (EPA) included this compound to the list of priority DBPs to be monitored in a nationwide occurrence study [46] due to anticipations of potential toxicity based on alarming structure-activity relationships [47]. To address this scarcity of data, we analyzed the genotoxicity of CH and BH using a battery of three genotoxicity assays, namely the Ames test, the comet assay, and the micronucleus assay. The use of a test battery is critical since no single genotoxicity test is capable of detecting all genotoxic mechanisms [48]. We performed the three assays in the absence and presence of the metabolic activation fraction S9 mix to assess the effects of metabolic reactions on the toxicity of the two compounds.

Methods

Chemicals

The identifiers and structures of CH and BH are shown in Table 1. CH (crystallized, $\geq 98\%$) was obtained from Sigma-Aldrich (China). BH was prepared by adding tribromoacetaldehyde (bromal, Sigma-Aldrich, UK, 97% purity), to ultrapure water and then recrystallizing the product from a small volume of water. Ultrapure water was produced from a Millipore water system (resistivity = 18.2 M Ω .cm). Before toxicological analyses, stock solutions were prepared in dimethyl sulfoxide (DMSO, Chromasolv plus, $\geq 99.7\%$) obtained from Sigma-Aldrich (UK) and were immediately stored in amber glass vials at -80 °C.

Metabolic activation mixture (S9 Mix)

The metabolic activation mixture was a 9000 g centrifuged supernatant of a liver homogenate (S9). It was prepared from male OFA rats (Charles River Laboratories, France). Five days before sacrifice, the rats were treated with a single injection of Aroclor 1254 (500 mg/kg body weight). The final protein concentration of the S9 mix was 26 mg/mL as determined by the method of Lowry et al. [49]. In the Salmonella mutagenicity assay, the composition of the metabolic mixture (S9 mix) included 4% S9, 10 mM glucose-6-phosphate (G6P) and 8 mM nicotinamide adenine dinucleotide phosphate (NADP) (De M \acute{e} o et al. 1996). In the comet and micronucleus assays, the S9 mix contained 10% S9, 5 mM G6P, 4 mM NADP, 33 mM KCL and 8 mM MgCl₂ diluted in 0.15 M saline phosphate buffer [50].

Cell cultures

The comet and the micronucleus assays were performed using Chinese Hamster Ovary cells (CHO-K1, ATCC). Cells were grown in McCoy's 5A medium (Sigma-Aldrich, St Quentin

Fallavier, France) supplemented with 10% fetal calf serum, 1 mM glutamine and penicillin-streptomycin (100 U/mL and 10 µg/mL), and incubated in a humidified atmosphere containing 5% carbon dioxide (CO₂).

Salmonella/microsome mutagenicity assay

The *Salmonella typhimurium* tester strains TA97a, TA98, TA100 and TA102 used in the Ames test were supplied by Prof. B.N. Ames (Berkeley, CA, USA). These strains were used to detect different types of mutations in agreement with the recommendations of Maron and Ames [51]. The strain TA100 is able to detect base-pair substitution mutations. The strains TA97a and TA98 are able to detect frameshift mutations. The strain TA102 detects cross-linking mutagens [52]. The strains were stored at -80 °C and regularly checked for genetic markers. The mutagenicity assay was carried out according to Maron and Ames [51], with a modified version of the liquid-incubation technique [53]. *Salmonella* strains were grown in Oxoid Nutrient Broth N° 2 with ampicillin (25 µg/mL) for 12 h at 37 °C with gentle shaking. After the incubation period, various volumes of solutions of the test substances (four test doses per compound), not exceeding 10 µL (0.5%, v/v) to avoid toxicity, were added to 0.1 mL of the overnight culture and 0.1 mL of PBS or S9 Mix. The mixtures were incubated either for 60 min in the dark. Then, 2 mL of melted top agar containing 0.045 mM histidine and biotin were added, and the mixtures were poured onto Vogel-Bonner (VB) minimal plates. For each series of experiments, negative controls with 5 or 10 µL of DMSO were included to determine the number of spontaneous revertants/plate. Positive controls were also included to ensure the performance of the tester strains: 0.002 µg/plate IRC191 for TA97a, 0.002 µg/plate 2,4,7-trinitrofluorenone for TA98, 5 µg/plate NaN₃ for TA100, 0.002 µg/plate mitomycin C for TA 102. After a 48-h incubation period, revertants were counted with an automatic counter (Scan 1200, Interscience, Saint Nom La Bretèche, France).

A two-step analysis was performed to interpret the data. The Dunnett test [54] was primarily performed to determine a significant difference between the mean number of induced revertants and the mean number of spontaneous revertants. If the Dunnett test was positive for at least one dose, a nonlinear regression analysis was carried out using an arbitrary model as described previously [54] with TableCurve 2D software (version 5.01, Jandel Scientific Software, San Rafael, CA):

$$Rev/plate = (a + b \times D) \times 2^{-c \times D^2}$$

where Rev/plate is the number of revertants by plate, D is the dose, and a, b, and c are calculated coefficients.

Model significance was based on two criteria: (i) model probability (P) being < 0.05 and (ii) error probability (P_E) being > 0.05 . The mutagenic activity (MA, rev/ μ g) was defined as the maximal slope of the ascending part of the dose-response curve and was calculated as the first derivative at the origin.

Cytotoxicity assay: WST-1 test

The cytotoxicities of CH and BH in CHO cells were analyzed using the WST-1-based cytotoxicity assay. The latter is a colorimetric assay that allows the quantitation of cellular cytotoxicity and proliferation based upon the reduction of the tetrazolium salt WST-1 to formazan by cellular dehydrogenases. A total of 50 000 cells were seeded in 96-well plates and incubated overnight at 37 °C with 5% CO₂ in a humidified incubator. A range of concentrations of BH and CH was incorporated in triplicate cultures (final DMSO concentration less than 0.2%), and cells were incubated at 37 °C for 2 h. At the end of the incubation period, cultures were submitted to three successive washes with PBS and incubated in 10% WST1 (Roche Diagnostics, Meylan, France) in culture medium for 30 min. Cell viability was evaluated by the

assessment of WST1 absorbance at 450 nm in a microplate reader (Infinite M200 Pro, Tecan, Lyon, France).

Alkaline comet assay

The alkaline comet assay was performed as described by Tice et al. [55] with slight modifications [56]. A total of 50 000 CHO cells were plated 24-well plates and incubated for 24 h at 37 °C in a humidified atmosphere containing 5% CO₂. Two sets of experiments were performed. In the first set of experiments, cells were rinsed and incubated into culture medium containing BH or CH solutions in duplicate. In the second set of experiments, cells were rinsed and incubated into culture medium containing 10% S9 mix, and BH or CH solutions. Two negative controls were added to determine the background DNA-damage levels in CHO cells: the culture medium control and a 1% DMSO control, corresponding to the maximal concentration of DMSO incorporated into cell cultures. Two positive controls with the well-known genotoxic compound Methyl methanesulfonate (MMS) with and without S9 Mix were also added.

After a contact period of 2 h, the cells were embedded in low-melting-point agarose. Lysis, DNA unwinding, and electrophoresis were performed as described previously [57]. Following the electrophoretic run, the slides were neutralized with 0.4 M Tris-HCl (pH 7.4), rinsed with ultrapure water, dipped into 100% methanol (HPLC-grade purity), and dried at room temperature. Staining was performed with ethidium bromide solution (2 µg/mL), and the slides were examined at 250 × magnification using a BX53 fluorescence microscope (Olympus France, Rungis, France). Image analysis was performed using the Komet software (version 6.0, Andor Technology, Belfast, Northern Ireland) on 100 randomly selected cells (50 cells for each of two replicate slides). DNA damage was expressed as Olive tail moment (OTM, arbitrary units) [50].

The calculated OTM values were distributed into 40 classes between the minimal and the maximal OTM values. A nonlinear regression analysis was performed on the normalized distribution frequencies using a χ^2 function with TableCurve 2D (version 5.01, Systat Software, Erkrath, Germany). The calculated degree of freedom (n) for the function, named OTM χ^2 , was assumed to be a quantitative measure of the level of DNA damage in the sample [58]. The test was considered positive when a dose-response relationship could be established between the OTM χ^2 and the concentrations of the compound and when one concentration at least induced a significant increase of OTM χ^2 by the comparison of means ($p < 0.05$). The genotoxicity was calculated by nonlinear regression analysis with table Curve 2 D using an arbitrary model

$$G = \frac{aCo}{b + Co} + c$$

where G: genotoxicity, Co: concentration (mg/mL) and a, b, and c: calculated coefficients. The maximal genotoxic activity was defined as the first derivative of the model at the maximal tested dose (1 $\mu\text{g/mL}$).

Micronucleus assay

A total of 50,000 CHO-K1 cells was plated in chamber slides and incubated for 24 h at 37 °C in a humidified atmosphere containing 5% CO₂. Various concentrations of BH and CH solutions were incorporated into duplicate cell cultures. Two negative controls were added to determine the background DNA-damage levels in CHO cells: the culture medium control and a 1% DMSO control, corresponding to the maximal concentration of DMSO incorporated into cell cultures. Two positive controls with well-known genotoxic compounds were also included to ensure the sensitivity of the assay: these were 0.06 $\mu\text{g/mL}$ mitomycine C without S9 Mix and 5 $\mu\text{g/mL}$ benzo[a]pyrene (BaP) with S9 Mix.

After a 3-h exposure, cells were rinsed with PBS and incubated in fresh medium containing cytochalasin B (3 $\mu\text{g}/\text{mL}$) for an additional 22 h period to stop cytokinesis. At the end of the incubation period, cells were submitted to two successive washes with PBS and fixed with methanol (HPLC purity grade solvent). The slides were air dried and stained with 5% Giemsa stain in Milli-Q water for 15 min.

The Cytokinesis Blocked Proliferation Index (CBPI) was used to select adequate concentrations for the assessment of micronuclei, as it has been considered a measure of cytotoxicity [58]. CBPI was determined by scoring the number of mononucleated (M1), binucleated (M2), and trinucleated (M3) cells among 500 Giemsa-stained cells with well-preserved cytoplasm: $\text{CBPI} = (\text{M1} + 2 \times \text{M2} + 3 \times \text{M3}) / 500$.

The Proliferative Index (CI%), i.e. the percentage of cell replication inhibition, was calculated using the following formula: $\text{PI}\% = 100 - \{100 \times (\text{CBPI}_{\text{test material}} - 1) / (\text{CBPI}_{\text{DMSO control}} - 1)\}$. After this step, only the doses inducing a decrease of less than 50% of PI% as compared to the negative control were taken into account for counting micronuclei. Micronuclei were identified according to the morphological criteria previously defined by Kirsch-Volders et al. [58]. Statistical differences between negative controls and treated samples were determined using the χ^2 test. The micronucleus assay was considered positive when a dose-response relationship could be established between micronucleated cells rates and concentrations, and when at least one concentration induced a significant increase of micronucleated cells as compared to the DMSO-control culture.

Results

The Ames test

We evaluated the capacity of CH and BH to induce mutations in DNA using the Ames test in four *Salmonella* tester strains: TA97a, TA98, TA100, and TA102. The assay was carried out in the absence and presence of exogenous metabolic activation (S9 mix). As shown in Table 2, CH did not exhibit mutagenic effects in the tested strains, unlike BH, which induced mutagenic activity in the strain TA100. The results of the regression analyses for the dose-response relationship of BH in the presence and absence of S9 mix are displayed in Figure 1. The mutagenic activity calculated for BH was 19.5 Rev/ μg (5.8 Rev/nmol) and 7.9 rev/ μg (2.4 Rev/nmol) in the absence and presence of the metabolic fraction S9 mix, respectively. Thus, the presence of the metabolic activation system induced about 60% decrease in the mutagenicity of BH.

Cytotoxicity assay on CHO cells

We examined the cytotoxicity of CH and BH in CHO using the WST-1-based assay. Figure 2 displays the cellular viability as a function of concentration for CH and BH. CH induced a slight decrease in the viability of CHO cells at concentrations up to 1.0 mg/mL. In contrast, BH exhibited cytotoxic activity at levels as low as 5.0 $\mu\text{g}/\text{mL}$.

The comet assay

We examined the capacity of CH and BH to induce DNA damage in CHO cells using the comet assay. CHO cells were exposed to CH and BH at different concentrations for 2 h in the absence and in the presence of the metabolic mixture S9 mix. Since BH displayed cytotoxicity in CHO cells as assessed by the WST-1 test, the used concentrations were selected to maintain cell viability above 95%. The results of the comet assay are reported in Figure 3A for OTM values and in Figure 3B for OTM χ^2 values. Parametric analyses were limited to OTM χ^2 values as OTM values were not normally distributed (Fig 3A, box plot). CH did not induce any significant

DNA strand breaks in the absence or in the presence of S9 mix up to 1 mg/mL. On the contrary, BH induced statistically significant DNA lesions at the three tested concentrations in the absence of metabolic activation only. Modeling of the dose-response curve for BH is displayed in Figure 4. The genotoxicity (G) was 3.68 OTM χ^2 units/ μg ($r^2 = 0.96$, $p < 10^{-5}$, $P_E = 0.07$).

The micronucleus assay

We examined the ability of CH and BH to induce chromosomal damage by conducting the micronucleus assay in CHO cells. Cells were exposed for 3 h to a range of concentrations of CH and BH in the absence and the presence of the metabolic activation fraction S9 mix. Table 3 shows the proliferative index (PI), the mean number of micronucleated binucleated cells (MBCs), and the p-values. PI gives an estimation of the antiproliferative/cytotoxic activity of compounds. PI values beyond 50% mean that evaluation of chromosomal damage would not be possible at the concentrations in question. These critical concentrations were 50 and 25 $\mu\text{g/mL}$ for CH and 2.5 and 5.0 $\mu\text{g/mL}$ for BH without and with the S9 mix, respectively. The number of MBCs was obtained by counting micronucleated binucleated cells per 1000 cells. No significant increases in micronucleated cell rates could be observed for both compounds. Thus, CH and BH did not exhibit any clastogenic or aneugenic effects against CHO cells.

Discussion

Several epidemiologic studies have suggested that exposure to DBPs is associated with increased incidence of cancer, particularly bladder and colorectal cancers [12-14]. The analysis of genotoxicity of DBPs allows identifying the chemical species that could be responsible for carcinogenicity [48]. Genotoxicity testing detects carcinogens that are thought to act primarily via a mechanism involving direct genetic damage [48]. CH has been frequently detected as a predominantly occurring DBP in chlorinated drinking and swimming pool waters [19,25,59]. In

the presence of relatively high levels of bromide, brominated DBPs are formed upon disinfection with chlorine [60]. BH has been detected as the predominant trihaloacetaldehyde in chlorinated seawater swimming pools where high bromide contents are found [19]. BH has been also detected as the major haloacetaldehyde in chloraminated drinking water originating from bromide-rich source water [25,46]. Although some studies have investigated the genotoxicity of CH, findings have not been consistent [25,45,61,62]. This could be due the equivocal responses exhibited by CH leading to different interpretations (as slightly genotoxic or not genotoxic) and therefore equivocal conclusions. In addition, very little is known about the toxicity of BH for which two genotoxicity investigations have reported positive genotoxic effects in the comet assay [25,45]. However, additional studies incorporating a battery of genotoxicity assays seem necessary to obtain further knowledge about the toxicity of CH and BH and the mechanisms of action involved in any eventual genotoxicity. We tested the potential of CH and BH to induce genetic damage using the Ames test, the comet assay, and the micronucleus assay. As reviewed by Claxton et al. [63], the Ames test serves as a reliable tool in screening for mutagenic agents. Extensive reviews have shown that many compounds that are mutagenic in the Ames test are rodent carcinogens [48]. The comet test allows measuring DNA single- and double-strand breaks and detecting alkali labile sites [64,65]. The induction of micronuclei has been demonstrated to be an effective biomarker of cancer risk [66]. The *in vitro* micronucleus assay detects reliably chromosomal breaks and aneuploidy [67-68]. The combination of the three assays could provide insights into the mechanism of action involved in genotoxicity.

As far as we know, this study evaluates for the first time the mutagenicity of BH using the *Salmonella* Ames test and the genotoxicity of both CH and BH using the micronucleus test in CHO cells. To take into account the effects of metabolic reactions on the genotoxicity of the two

compounds, we carried out the tests in the presence and absence of metabolic activation fraction S9 mix.

According to our findings, CH did not exhibit genotoxic activity in any of the three assays. These results are consistent with previous studies reporting the lack of genotoxicity indications for CH in the comet assay using CHO cells [25, 61]. Similarly, in an evaluation of the mutagenicity of CH *in vitro* and *in vivo*, authors reported the absence of indications for a mutagenic potential [62]. In the same way, negative results have been reported in the micronucleus test where CH failed to induce micronuclei or showed very weak effects in human lymphocytes [69], mouse lymphoma cells [70], TK6 cells and human peripheral blood lymphocytes [45]. Some investigations reporting positive genotoxicity findings have been reported in the literature for CH. Using the comet assay, Liviak et al. [45] reported genetic damages caused by CH in TK6 cells. In the micronucleus assay, CH induced aneuploidy in CHO cells at high doses [71] and induced micronuclei in mouse spermatocytes [41-43], and human fibroblasts [72].

Concerning BH, our findings show that it induced mutagenic activity in the Ames test, caused genetic damage in CHO cells in the comet assay, but failed to induce micronuclei in the CHO cells. In the Ames test, positive results were limited to the TA100 strain. The hisG46 marker in TA100 results from the substitution of a leucine (GAG/CTC) by a proline (GGG/CCC). This mutation is reverted to the wild-type strain by mutagens that cause base-pair substitution mutations at one of the GC pairs. Among them, brominated reactive byproducts may combine with G to form a small adduct and thus can revert the mutation. The genotoxicity of BH decreased considerably when metabolic activation was incorporated suggesting the formation of less toxic metabolites. Our findings are consistent with previous investigations reporting genotoxic activity for BH detected using the comet assay in CHO cells [25], human lymphocytes

and TK6 cells [45]. Moreover, a negative response has been previously reported for BH using the micronucleus test in human lymphocytes and TK6 cells [45]. Overall, the results of the three assays imply that the mechanism of toxicity of BH involves the induction of mutations and primary DNA lesions but not chromosomal damage.

The present study corroborates previous investigations highlighting the higher genotoxic potential for brominated DBPs in comparison to their chlorinated analogs [20,25,27,73,74]. This trend has been reported for several chemical classes of DBPs including THMs, HAAs, and HANs. The difference in genotoxic potentials between brominated and chlorinated species has been attributed to differences in the leaving tendency of the halogens in the alkyl halide [25, 27,28,75]. These findings highlighting the toxicity of brominated DBPs are of relevance to public health since epidemiologic investigations have associated statistically significant increases in risks of birth defects with residence in areas where water supply had high levels of brominated DBPs [76].

One of the limitations of this study was the lack of data about the pharmacokinetics of BH, so its absorption and fate once introduced to biological media remain unknown. Also, the present study is limited to genotoxicity assessment, yet there are other events that could lead to carcinogenesis besides genotoxic effects. In addition, the study is limited to a genotoxicity endpoint, but DBPs could cause other toxicological effects and be implicated in some pathological conditions. Despite these limitations, the *in vitro* assays used here are reliable tools to detect genotoxic compounds such as BH and hence to highlight the importance of limiting the exposure of individuals to DBPs.

Conclusions

Our findings show that CH did not induce any genotoxic effects using the Ames test, the comet assay, and the micronucleus test, while BH was mutagenic in the *Salmonella* assay and caused genomic damage in the comet assay but not in the micronucleus test. These results imply that the mechanism of genotoxicity of BH involves inducing mutations and DNA damage but not chromosomal aberrations. The genotoxicity of BH decreased in the presence of metabolic fractions suggesting the formation of less genotoxic metabolites. Furthermore, our findings highlighting the toxicity of brominated DBPs are of relevance to public health since epidemiologic investigations have associated statistically significant increases in risks of birth defects with residence in areas where water supply had high levels of brominated DBPs [76]. This study providing evidence of the genotoxic potential of a commonly occurring DBP emphasize the importance of reducing the levels of DBPs, particularly brominated species, in disinfected waters. Further research examining the genotoxicity of other DBPs is warranted to identify genotoxic DBPs that could be responsible for the reported adverse health effects.

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Figure Legends

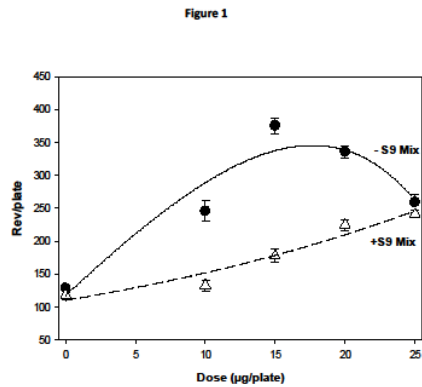


Figure 1. Dose-response relationship of BH on the tester strain TA100 with and without the S9 mix. Non-linear regression analyses were performed as described in the methods section.

In the absence of the S9 mix (dark circles, black line), $r^2 = 0.90$, $P < 10^{-5}$ and $P_{\text{error}} = 0.65$

In presence of the S9 mix (white triangles, dashed line), $r^2 = 0.90$, $P < 10^{-5}$ and $P_{\text{error}} = 0.07$

Figure 2

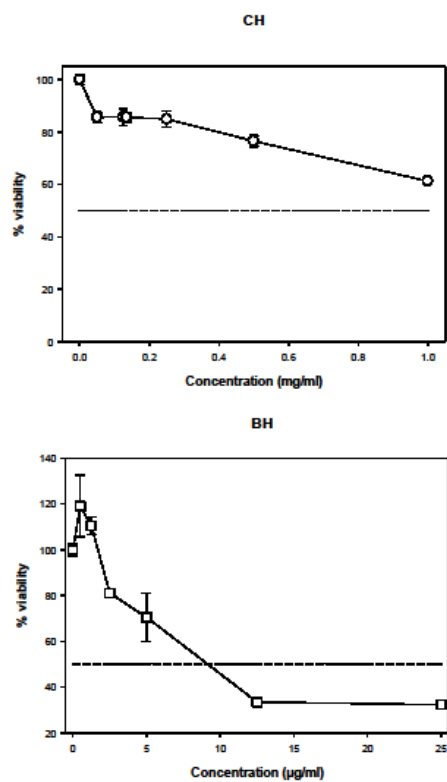


Figure 2. Cytotoxicity assay (WST-1) on CHO cells for CH and BH. The dashed line represents the 50% viability

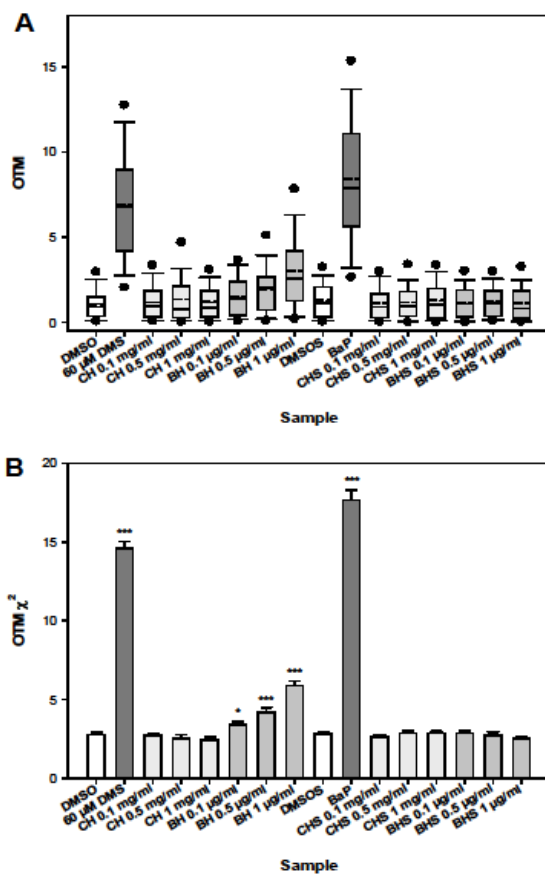


Figure 3. Results of the Comet assay on CHO cells for CH and BH.

- A.** Box-and-Whisker plot of the distributions of OTM values. The graphs display the 25th and 75th percentiles (lower and upper sides of boxes), the 5th and 95th percentiles (error bars), the lowest and highest OTM values (black circles), the median (black line), and the mean (dashed line). Each condition includes 100 OTM values.
- B.** Histogram of OTM χ^2 values.
- DMSO: dimethylsulfoxide; DMS: dimethylsulfate; BaP: benzo[a]pyrene; BH: bromal hydrate; CH: chloral hydrate; S: treatment with S9 mix
- *: $P < 0.05$; ***: $P < 0.001$

Figure 4

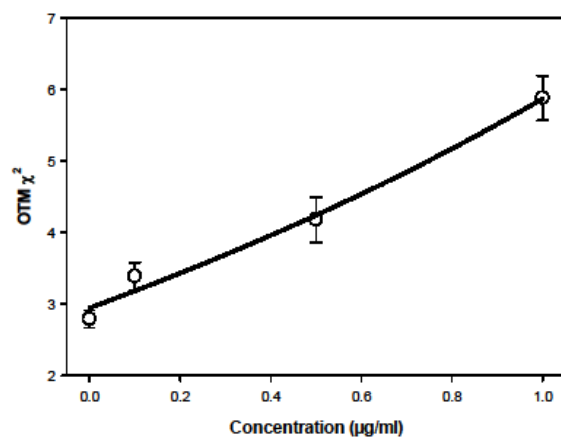


Figure 4. Dose-response relationship of BH on CHO cells with the comet assay.

A non-linear regression analysis was performed as described in the methods section $r^2 = 0.96$, $P < 10^{-5}$ and $P_{\text{error}} = 0.07$. The maximal calculated genotoxic activity was $3.68 \text{ OTM } \chi^2/\mu\text{g}$.

Table Legends**Table 1.** Names, CAS numbers, and chemical structures of CH and BH

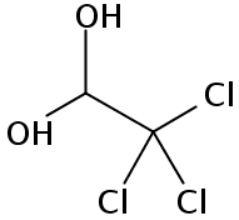
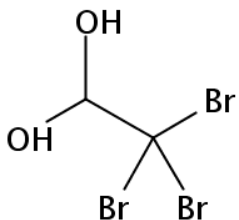
Name	CAS Number	Structure
Chloral Hydrate 2,2,2-Trichloro-1,1-ethanediol Trichloroacetaldehyde hydrate	302-17-0	 <chem>OCC(Cl)(Cl)Cl</chem>
Bromal Hydrate 2,2,2-Tribromo-1,1-ethanediol Tribromoacetaldehyde hydrate	507-42-6	 <chem>OCC(Br)(Br)Br</chem>

Table 2. Complete results of the Ames test for CH and BH with and without the S9 mix

Sample	Dose ^a	Strain (rev/plate)							
		TA97a		TA98		TA100		TA102	
		-S9	+S9	-S9	+S9	-S9	+S9	-S9	+S9
CH									
	0	151±4	151±16	22±3	36 ± 4	178±12	173±12	521±14	504±16
	20	131±4	162±6	29±5	29±3	179±13	177±15	500±6	441±4
	30	136±12	145±11	23±3	29±4	179±11	182±10	495±19	468±12
	40	141±14	154±10	30±3	25±4	174±3	170±7	521±4	502±14
	50	144±2	144±11	20±6	26±4	177±8	178±17	518±6	502±20
Pos. Ctrl ^b		258±10	1198±112	94±7	642±18	2119±13	1312±1	1939±30	863±33
BH									
	0	139±13	153±23	20±4	26±5	126±4	118±11	521±10	494±26
	20	168±11	168±3	24±1	34±2	245±20	133±10	480±18	400±28
	30	163±21	213±7	33±5	28±4	375±14	177±12	502±4	426±14
	40	164±6	206±16	32±4	37±1	336±12	224±10	506±6	432±49
	50	120±8	169±51	32±6	33±3	259±14	241±3	521±8	407±20
Pos. Ctrl ^b		199±1	1039±50	147±20	639±10	1677±6	1516±39	3617±152	906±20

Pos. Ctrl: positive control

All data are the means ± SD of three consecutive independent experiments.

Bold characters: statistically significant difference with the control by the Dunnett's test at $P < 0.01$

^aDose in µg/plate

^bPositive controls used in the absence of S9 mix were Acridine Mutagen (ICR191; 0.002 µg/plate), 2,4,7-Trinitro-9-fluorenone (TNF; 0.002 µg/plate), sodium azide (NaN₃; 10 µg/plate), Mitomycin C (0.002 µg/plate) in TA97a, TA98, TA100, and TA102 strains, respectively. In the presence of S9 mix, benzo[a]pyrene (BaP; 1 µg/plate) was used as a positive control in the four tester strains.

Table 3. Complete results of the micronucleus assay on CH and BH with and without the S9 mix

Sample	Concentration (µg/ml)	Without S9 Mix			With S9 Mix		
		PI (%)	Mean MBC (‰)	p*	PI (%)	Mean MBC (‰)	p*
Neg. Ctrl	0	100	9.5 ± 0.7		100	10.5 ± 0.7	
Pos. Ctrl	0.05 ^a , 5 ^b	98	30 ± 2.8	<0.001	98	23.5 ± 2.1	<0.01
DMSO	1%	95	8.5 ± 0.7	NS	99	9.5 ± 0.7	NS
BH	0.1	95	10 ± 1.4	NS	97	10 ± 1.4	NS
	0.25	86	10 ± 2.8	NS	95	10.5 ± 0.7	NS
	0.5	82	10.5 ± 0.7	NS	95	11.5 ± 0.7	NS
	1	75	11 ± 1.4	NS	84	12.5 ± 0.7	NS
	2.5	toxic	-	-	72	14 ± 2.8	NS
	5	toxic	-	-	toxic	-	-
CH	0.5	99	10.5 ± 0.7	NS	97	10.5 ± 0.7	NS
	1.25	96	11.5 ± 0.7	NS	94	11 ± 1.4	NS
	2.5	92	10.5 ± 0.7	NS	89	10 ± 2.8	NS
	5	88	10.5 ± 2.1	NS	85	10.5 ± 0.7	NS
	12.5	79	10 ± 1.4	NS	71	13 ± 1.4	NS
	25	72	10.5 ± 0.7	NS	toxic	-	-
	50	toxic	-	-	toxic	-	-

Neg. Ctrl: negative control

Pos. Ctrl: positive control

NS: non-significant

PI: proliferative Index expressed in %, when less than 50% the cytotoxicity impairs the evaluation of genotoxicity (denoted as toxic in the table)

Mean MBC: the mean number of micronucleated binucleated cells in 1000 binucleated cells

^aconcentration of Mitomycin C, the positive control in the absence of S9 mix ^bconcentration of BaP, the positive control in the presence of S9 mix

*compared with negative controls (chi-squared test)