

genotype (having the *GSTT1-1* gene). This finding, along with mechanistic studies, highlights the emerging importance of dermal/inhalation exposure to the THMs, or possibly other DBPs, and the role of genotype for risk for drinking-water-associated bladder cancer. More than 50% of the total organic halogen (TOX) formed by chlorination and more than 50% of the assimilable organic carbon (AOC) formed by ozonation has not been identified chemically. The potential interactions among the 600 identified DBPs in the complex mixture of drinking water to which we are exposed by various routes is not reflected in any of the toxicology studies of individual DBPs. The categories of DBPs described here, the identified data gaps, and the emerging role of dermal/inhalation exposure provide guidance for drinking water and public health research.

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Contents

| | |
|---------------------------------------------------------------------------------------|-----|
| 1. Introduction | 180 |
| 2. Overview of DBP regulations in the United States | 183 |
| 3. Summary of epidemiology studies of cancer and drinking water | 185 |
| 4. Occurrence, genotoxicity, and carcinogenicity of the regulated DBPs | 185 |
| 4.1. Trihalomethanes (THMs) | 185 |
| 4.1.1. Occurrence | 185 |
| 4.1.2. Genotoxicity | 186 |
| 4.1.3. Carcinogenicity | 188 |
| 4.2. Haloacetic acids (HAAs) | 191 |
| 4.2.1. Occurrence | 191 |
| 4.2.2. Genotoxicity | 191 |
| 4.2.3. Carcinogenicity | 193 |
| 4.3. Bromate | 193 |
| 4.3.1. Occurrence | 193 |
| 4.3.2. Genotoxicity | 193 |
| 4.3.3. Carcinogenicity | 194 |
| 4.4. Chlorite | 194 |
| 4.4.1. Occurrence | 194 |
| 4.4.2. Genotoxicity | 195 |
| 4.4.3. Carcinogenicity | 195 |
| 5. Summary of the occurrence, genotoxicity, and carcinogenicity of the regulated DBPs | 195 |
| 5.1. Summary of the occurrence of the regulated DBPs | 195 |
| 5.2. Summary of the genotoxicity of the regulated DBPs | 195 |
| 5.3. Summary of the carcinogenicity of the regulated DBPs | 195 |
| 5.4. Overall summary of the regulated DBPs | 196 |
| 6. Emerging unregulated DBPs | 197 |
| 6.1. Halonitromethanes | 197 |
| 6.1.1. Occurrence | 197 |
| 6.1.2. Genotoxicity | 198 |
| 6.1.3. Carcinogenicity | 199 |
| 6.2. Iodo-acids and other unregulated halo-acids | 199 |
| 6.2.1. Occurrence | 199 |
| 6.2.2. Genotoxicity | 202 |
| 6.2.3. Carcinogenicity | 202 |
| 6.3. Iodo-THMs and other unregulated halomethanes | 202 |
| 6.3.1. Occurrence | 202 |
| 6.3.2. Genotoxicity | 203 |
| 6.3.3. Carcinogenicity | 204 |
| 6.4. MX and BMX compounds (halofuranones) | 205 |
| 6.4.1. Occurrence | 205 |
| 6.4.2. Genotoxicity | 205 |
| 6.4.3. Carcinogenicity | 207 |

