#### MODELING DISSOLVED OZONE AND BROMATE ION FORMATION IN OZONE CONTACTORS

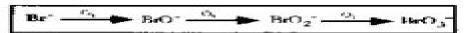
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(Received 6 September, 1996; accepted 20 September, 1997)

Abstract. Many drinking water utilities that use source waters with significant bromide ion levels are currently seeking disinfection regimes that will minimize the formation of brominated disinfection byproclacus while providing adequate disinfection. While occuration appears to be a promising option for achieving those goals, the uncertainty of future drinking water regulations has developed a need for predicting actual disinfection byproduct formation prior to the costly insentment for appracing cuisting treatment her lifties. The models developed in this paper provide comparisons of consistion application methods, providing a basis for minimizing becomes and aiding in future design considerations. Theoretical and empirical models for the determination of coone transferred, dissolved occure concentrations, and bromate formation have been derived and compared with pilot-scale and full-scale data; a good agreement has been observed between the actual data and the predicted data, showing the validity of these models. True-batch bromate formation more closely simulated pilot-scale and full-scale data. Bromate formation in one stage or two stage ocone contactors and different reactor configurations have been compared. Oxone gas phase concentration appears to have an effect on bromation have been compared. Oxone gas phase concentration appears to have an effect on bromatic formation as well.

#### 1. Introduction

Ozone is a strong oxidant and it oxidizes bromide iron Br<sup>-</sup> in water to bromate ion BrO<sub>3</sub><sup>-</sup> through two different pathways. BrO<sub>3</sub><sup>-</sup> can form through both a molecular ozone (O<sub>3</sub>) pathway and a hydroxyl radical (OH.)\* pathway, depending on the dissolved organic carbon (DOC), br<sup>-</sup> content, and pH of the source waters (Richardson et al., 1981 and Yates and Strenstrom, 1993). By the molecular ozone pathway, Br<sup>-</sup> is first oxidized by dissolved ozone to hypobromite ion (OBr<sup>-</sup>) which is then further oxidized to BrO<sub>3</sub><sup>-</sup> as shown below.



This reaction is pH-dependent since  $BrO_3^-$  is in equilibrium with hyprobromous acid (HOBr). The molecular ozone theory suggests that  $BrO_3^-$  formation is directly driven by dissolved ozone (DO<sub>3</sub>) and the OH indical theory indicates that dissolved ozone plays only an indirect role by decomposing O<sub>3</sub> to produce radicals which further seact with bromine species to produce  $BrO_3^-$ :

OH radicals are produced as a result of the decomposition of molecular ozone (O<sub>2</sub>) at pH>8.0, O<sub>2</sub> is highly selective as an oxident whereas OH radicals are non-selective. The reaction rates of OH radicals with beomine species are many orders of magnitude faster than with O<sub>2</sub>.

Water, Air, and Soil Pollistion 108: 1-32, 1998.

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#### **Table of Contents Modeling Dissolved Ozone Behavior In Ozone Contactors**

- 1. Understanding the eBook Modeling Dissolved Ozone Behavior In Ozone Contactors
  - The Rise of Digital Reading Modeling Dissolved Ozone Behavior In Ozone Contactors
  - Advantages of eBooks Over Traditional Books
- 2. Identifying Modeling Dissolved Ozone Behavior In Ozone Contactors
  - Exploring Different Genres
  - Considering Fiction vs. Non-Fiction
  - Determining Your Reading Goals
- 3. Choosing the Right eBook Platform
  - Popular eBook Platforms
  - Features to Look for in an Modeling Dissolved Ozone Behavior In Ozone Contactors
  - User-Friendly Interface
- 4. Exploring eBook Recommendations from Modeling Dissolved Ozone Behavior In Ozone Contactors

- Personalized Recommendations
- Modeling Dissolved Ozone Behavior In Ozone Contactors User Reviews and Ratings
- Modeling Dissolved Ozone Behavior In Ozone Contactors and Bestseller Lists
- 5. Accessing Modeling Dissolved Ozone Behavior In Ozone Contactors Free and Paid eBooks
  - Modeling Dissolved Ozone Behavior In Ozone Contactors Public Domain eBooks
  - Modeling Dissolved Ozone Behavior In Ozone Contactors eBook Subscription Services
  - Modeling Dissolved Ozone Behavior In Ozone Contactors Budget-Friendly Options
- 6. Navigating Modeling Dissolved Ozone Behavior In Ozone Contactors eBook Formats
  - o ePub, PDF, MOBI, and More
  - Modeling Dissolved Ozone Behavior In Ozone Contactors Compatibility with Devices
  - Modeling Dissolved Ozone Behavior In Ozone Contactors Enhanced eBook Features
- 7. Enhancing Your Reading Experience
  - Adjustable Fonts and Text Sizes of Modeling Dissolved Ozone Behavior In Ozone Contactors
  - Highlighting and Note-Taking Modeling Dissolved Ozone Behavior In Ozone Contactors
  - Interactive Elements Modeling Dissolved Ozone Behavior In Ozone Contactors
- 8. Staying Engaged with Modeling Dissolved Ozone Behavior In Ozone Contactors
  - Joining Online Reading Communities
  - Participating in Virtual Book Clubs
  - Following Authors and Publishers Modeling Dissolved Ozone Behavior In Ozone Contactors
- 9. Balancing eBooks and Physical Books Modeling Dissolved Ozone Behavior In Ozone Contactors
  - Benefits of a Digital Library
  - Creating a Diverse Reading Collection Modeling Dissolved Ozone Behavior In Ozone Contactors
- 10. Overcoming Reading Challenges
  - Dealing with Digital Eye Strain
  - Minimizing Distractions
  - Managing Screen Time
- 11. Cultivating a Reading Routine Modeling Dissolved Ozone Behavior In Ozone Contactors
  - Setting Reading Goals Modeling Dissolved Ozone Behavior In Ozone Contactors
  - Carving Out Dedicated Reading Time
- 12. Sourcing Reliable Information of Modeling Dissolved Ozone Behavior In Ozone Contactors

- Fact-Checking eBook Content of Modeling Dissolved Ozone Behavior In Ozone Contactors
- Distinguishing Credible Sources
- 13. Promoting Lifelong Learning
  - Utilizing eBooks for Skill Development
  - Exploring Educational eBooks
- 14. Embracing eBook Trends
  - Integration of Multimedia Elements
  - Interactive and Gamified eBooks

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