Modular Advanced Oxidation Process Enabled by Cathodic Hydrogen Peroxide Production

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Supporting Information

ABSTRACT: Hydrogen peroxide (H₂O₂) is frequently used in combination with ultraviolet (UV) light to treat trace organic contaminants in advanced oxidation processes (AOPs). In small-scale applications, such as wellhead and point-of-entry water treatment systems, the need to maintain a stock solution of concentrated H₂O₂ increases the operational cost and complicates the operation of AOPs. To avoid the need for replenishing a stock solution of H₂O₂, a gas diffusion electrode was used to generate low concentrations of H₂O₂ directly in the water prior to its exposure to UV light. Following the AOP, the solution was passed through an anodic chamber to lower the solution pH and remove the residual H₂O₂. The extent of transformation of trace organic contaminants was a function of the current density and the concentrations of HO· scavengers in the source water. The electrical energy per order (EEO) ranged from 1 to 3 kWh m⁻³, with the UV lamp accounting for most of the energy consumption. The gas diffusion electrode exhibited high efficiency for H₂O₂ production over extended periods and did not show a diminution in performance in any of the matrices.

INTRODUCTION

Distributed water treatment systems offer a potential means of exploiting alternative water sources, including municipal wastewater effluent, roof water, stormwater, and water from shallow aquifers.¹ Unfortunately, alternative water sources often contain trace concentrations of organic contaminants (e.g., pesticides, solvents, pharmaceuticals, disinfection byproducts).² As a result, distributed treatment is often seen as an impractical means of providing potable water. Previous attempts to develop point-of-use treatment systems capable of removing trace organic contaminants prior to nonpotable reuse have employed electrochemical processes³,⁴ but these systems suffer from limitations including the production of toxic byproducts, an inability to remove recalcitrant compounds and high cost of treatment.⁵

Trace organic contaminants can be removed from water by exposure to hydroxyl radicals (HO·) in advanced oxidation processes (AOPs).⁶⁻⁹ In full-scale potable water reuse systems, trace organic contaminants are frequently removed by addition of a modest concentration of H₂O₂ (e.g., 3 mg/L) followed by exposure to ultraviolet (UV) light. This approach offers numerous benefits over other AOPs in terms of energy consumption, reliability, and production of toxic byproducts.¹⁰ Although UV/H₂O₂ is a well established technology in centralized treatment facilities, challenges associated with the transport and storage of H₂O₂ make it an impractical solution for distributed treatment systems.¹¹ Electrochemical production of H₂O₂ from O₂ is an attractive alternative means of producing H₂O₂ if it can be achieved without the consumption of large amounts of energy or the formation of toxic byproducts.¹²

Electrochemical production of low concentrations of H₂O₂ can be achieved by several different approaches. Systems in which oxygen is bubbled into a solution prior to reduction on an electrode surface consume considerable amounts of energy due to the low solubility of oxygen and the need to ensure that it reaches the electrode surface. Bubbling air or pure oxygen into a solution is also an impractical approach for H₂O₂ production in decentralized systems because it requires pumps and controllers. Furthermore, the yield of H₂O₂ from reduction of O₂ is often quite low, which greatly increases electricity consumption.¹¹,¹²,¹³ Recently, gas diffusion electrodes have been used to generate H₂O₂ without a need to bubble air or oxygen into a solution.¹¹,¹³,¹⁴ Most of the research in this

Received: March 11, 2015
Revised: May 12, 2015
Accepted: May 19, 2015
Published: June 3, 2015
area has been focused on producing concentrated H$_2$O$_2$ solutions by using highly conductive solutions or organic solvents.\textsuperscript{12,13} Cathodic production of H$_2$O$_2$ for the removal of organics has typically been used for electro-Fenton treatment; however, differences in pH needed for the optimal kinetics of the two reactions (i.e., production of H$_2$O$_2$ is most efficient at basic pH values and Fenton’s reaction is more effective at acidic pH values) results in inefficient oxidation of trace organics if pH correction is not performed.\textsuperscript{11,15} To produce low concentrations of H$_2$O$_2$ in water immediately prior to an AOP, the cathode must be capable of producing H$_2$O$_2$ at a low ionic strength, poorly buffered solution at circumneutral pH values. Furthermore, increases in pH that occur in the cathode chamber due to consumption of protons must be compensated for by a subsequent treatment process if the water is to be sent into a water distribution system.

The purpose of this study was to evaluate a system that combines in situ electrochemical production of H$_2$O$_2$ followed by UV irradiation and anodic pH adjustment as a cost-effective means of removing trace organic contaminants from water. This system, which also inactivates waterborne pathogens and transforms photolabile contaminants through exposure to UV light, can be controlled by varying the production of H$_2$O$_2$ through adjustment of the applied current. To provide insight into the performance of the system under conditions likely to be encountered in distributed water treatment systems, three representative source waters (i.e., synthetic surface water, synthetic groundwater, and municipal wastewater effluent) were tested and compared to an electrolyte solution consisting of dilute sodium chloride. The performance of the system was investigated in terms of contaminant removal and energy consumption.

\section*{MATERIALS AND METHODS}

\textbf{Materials.} All experiments were performed at room temperature ($23 \pm 2$ °C) with chemicals of reagent grade or higher (Sigma-Aldrich, St. Louis, MO). The composition of the waters used is summarized in Table S1, Supporting Information.

\textbf{Electrochemical Cell and UV Reactor.} Experiments were carried out in a two-chambered parallel plate electrochemical cell consisting of two square Perspex frames (internal dimensions: $8 \times 8 \times 1.9$ cm$^3$) separated by a cation exchange membrane (Ultrex CMI-7000, Membranes International Inc., Ringwood, NJ). The frames were bolted together between two square Perspex side plates, creating anode and cathode compartments that each had effective volumes of 122 mL (Figure S1, Supporting Information). A solid plate was used for the anode frame, while the cathode chamber was bolted with a hollow side plate allowing for one side of the gas diffusion cathode to be exposed to air. A Ti mesh electrode coated with an Ir mixed-metal oxide was used as the anode (dimensions: 7.8 × 7.8 cm; 1 mm thickness; specific surface area 1.0 m$^2$ m$^{-2}$; Magneto Special Anodes, Netherlands). The anode and cathode had projected electrode surface areas of 64 cm$^2$. The UV reactor consisted of a 1 L brown glass bottle (V$_{\text{effective}}$ = 925 mL) containing a low-pressure UV lamp (arc length = 16.5 cm, optical path length = 4.3 cm) used typically for swimming pool disinfection (G23 Odyssey Pool Lamp, 9W, Odyssey Aquarium Appliance Co., Ltd., Guangdong, China; Figure S1, Supporting Information).

\textbf{Gas Diffusion Cathode Fabrication.} The gas diffusion cathode was created by modifying carbon fiber paper (AvCarb P7ST, $10 \times 10$ cm$^2$, Fuel Cell Store, College Station, TX) with a conductive, hydrophobic support layer and a carbon catalyst.\textsuperscript{14} The air-facing side of the cathode was prepared by coating a mixture of 60 wt % PTFE and 30 wt % graphite powder (200 mesh, Alfa Aesar, Ward Hill, MA) onto one side of the carbon base layer. The cathode was then air-dried at room temperature, followed by sintering at 350 °C for 40 min. The liquid-facing side was prepared by applying a mixture of 3 mL of propanol with 150 mg of carbon black (Cabot Black Pearls 2000, Cabot, Boston, MA) and 50 mg of PTFE onto the other side of the carbon base layer. The cathode was again air-dried at room temperature, followed by sintering at 350 °C for 40 min.

\textbf{Experimental Approach.} Electrolysis experiments were performed at fixed currents controlled by a multichannel potentiostat (Gamry Instruments Inc., Warminster, PA). Water entered the cathode compartment operating in a flow-through mode with hydraulic residence times (HRT) ranging from 1.5 to 5.0 min (120–35 L d$^{-1}$). Cathode effluent was supplied to the UV reactor and then passed through the anode of the electrochemical cell. The applied charge density ($\rho_q$, C L$^{-1}$) was expressed as a product of the current density ($I$, A m$^{-2}$), electrode surface area ($A$, m$^2$), and the hydraulic residence time ($t$, s) normalized by the half chamber reactor volume ($V$, L):

$$\rho_q = \int \frac{I(t) \times A}{V} \times dt = \frac{IAt}{V}$$

(1)

Source waters were amended with a mixture of ten test compounds each at a concentration of 10 µg L$^{-1}$. For each experiment, samples were collected prior to the electrochemical cell, after passing through the cathode chamber, after the UV reactor, and after passing through the anode. At least 3.5 L of the test solution was passed through the system prior to collection of a sample. Samples were analyzed for H$_2$O$_2$, trace organic compounds, and pH. 1.9 mL subsamples to be analyzed for trace organic compounds were mixed with 0.1 mL of methanol to quench radical reactions that could occur prior to analysis. H$_2$O$_2$ and pH were measured within 5 min, whereas trace contaminants were stored for a maximum of 8 h. Experiments quantifying H$_2$O$_2$ production in the varying waters were performed with applied cathodic current densities from 0 to 30 A m$^{-2}$ under varying flow regimes (35–120 L d$^{-1}$). To assess the long-term cathode performance, 6000 L of 5 mM Na$_2$SO$_4$ in tap water (alkalinity = 0.34 mM, [Ca$^{2+}$] = 0.2 mM) was run through the cell continuously at an applied current density of 15 A m$^{-2}$ at a fixed flow rate of 120 L d$^{-1}$. Samples were collected daily and analyzed for H$_2$O$_2$. The effects of dissolved oxygen concentration on H$_2$O$_2$ production were evaluated by sparging source water with N$_2$ to remove O$_2$. The rate of production of H$_2$O$_2$ remained unchanged under N$_2$-sparged conditions.

\textbf{Analytical Methods.} H$_2$O$_2$ and free chlorine were measured with a Shimadzu UV-2600 spectrophotometer with the titanium(IV) sulfate method at 405 nm and the N,N-diethyl-p-phenylenediamine (DPD) method at 515 nm, respectively.\textsuperscript{16,17} Determination of free chlorine was performed in the absence of H$_2$O$_2$ to eliminate the positive interference of H$_2$O$_2$ with DPD.\textsuperscript{18} The UV absorbance of the four source waters was measured with a Shimadzu UV-2600 spectrophotometer. Dissolved organic carbon and dissolved inorganic carbon (DIC) were measured using a Shimadzu TOC-V analyzer. NO$_3^-$, Cl$^-$, and SO$_4^{2-}$ were analyzed using a Dionex DX-120 ion chromatograph with an AS19G column.
K⁺, Na⁺, Ca²⁺, and Mg²⁺ were analyzed using a Dionex ICS-2000 ion chromatograph with a CS12A column. Fluence rate values were determined by chemical actinometry using 10 μM atrazine as an actinometer (ε₂₅₄ = 3860 M⁻¹ cm⁻¹), buffered at pH = 8 using a borate buffer. Conductivity was measured with an Ultrameter II 4P (Myron L Company, Carlsbad, CA). Test compounds were quantified in multiple reaction monitoring (MRM) mode with an Agilent 1200 series HPLC system coupled to a 6460 triple quadrupole tandem mass spectrometer (HPLC-MS/MS), as described previously.²¹ Analytical details and compound specific parameters are provided in the Supporting Information text and Tables S2 and S3, respectively.

**Electrical Power Calculations.** The gas diffusion electrode was polarized cathodically against a Ag/AgCl reference electrode (±0.197 V vs SHE; BASI, USA). The full cell potential between the working (i.e., cathode) and counter (i.e., anode) electrodes was measured in a two-electrode setup. The total system power (P_total, W) is a combination of the UV lamp power (P_lamp, W) and the electrochemical cell power, which can be expressed as a product of the current density (I, A m⁻²), cell potential (V_cell), and the electrode surface area (A, m²):

\[
P_{\text{total}} = I \times A \times V_{\text{cell}} + P_{\text{lamp}}
\]

**RESULTS**

**Hydrogen Peroxide Production as a Function of Current in Varying Source Waters.** Hydrogen peroxide concentrations (Figure 1A) and production rates (Figure S2, Supporting Information) were determined for an array of charge densities that were achieved from combinations of current densities (0–30 A m⁻²) and cathode chamber retention times (1.5–5 min). A linear relationship between H₂O₂ production and applied charge density was observed at charge densities greater than 50 C L⁻¹, independent of how the charge was obtained (i.e., high current density and short retention times or low current density and long retention times):

\[
[H_2O_2] = 0.0048 \times \rho_q \quad (50 \leq \rho_q \leq 375 \text{C L}^{-1})
\]

where ρ_q is the specific charge density applied in C L⁻¹ and [H₂O₂] is the hydrogen peroxide concentration in mM. For all of the waters tested, the Coulombic efficiency of O₂ reduction to H₂O₂ averaged 88.8 ± 1.8% at charge densities greater than 50 C L⁻¹ (Figure 1B). At lower charge densities, lower Coulombic efficiencies were observed (note the deviation from the linear fit in Figure 1A at charge densities below 50 C L⁻¹). The observed increased Coulombic efficiencies at higher charge densities agreed with previously published data for electrochemical synthesis of H₂O₂ using a gas diffusion electrode composed of a fluorocarbon binder and activated carbon catalyst fed with conductive, alkaline solutions.¹¹,¹²,²²

H₂O₂ production was independent of the type of source water used despite the substantial variability in the composition of the matrices (Table 1). This suggests that H₂O₂ production was not affected by pH, the presence of natural organic matter (NOM), dissolved ions, or conductivity over the range of applied charge densities studied. Although H₂O₂ production was not influenced by influent water quality, the cell potential, and therefore energy consumption, was affected by the conductivity of the source waters. Higher ionic strength waters exhibited lower ohmic resistances and therefore operated at lower total system powers (Figure 1C).
lower cell potentials, thus decreasing their energy consumption. For a given applied charge, the power required decreased with increased conductivity (synthetic surface water > synthetic groundwater > wastewater effluent ≡ electrolyte) (Figure S3, Supporting Information). Even for low conductivity surface water, however, the energy consumption for hydrogen peroxide production at a flow rate of 120 L d⁻¹ was still relatively low (0.018–0.31 kWh m⁻³ for 5 < I < 30 A m⁻²), indicating that in situ H₂O₂ production required much less energy than operation of the UV lamp (1.8 kWh m⁻³; calculations provided in the Supporting Information).

The H₂O₂ production rate increased linearly with applied current density, with a maximum of between 14.4 and 14.8 mg H₂O₂ L⁻¹ min⁻¹ at 30 A m⁻² for all of the waters tested (Figure S2, Supporting Information). In full-scale AOP systems (e.g., the Orange County Water District’s Groundwater Replenishment System), 3 mg H₂O₂ L⁻¹ (0.09 mM) is typically applied. This concentration can be obtained with the gas diffusion electrode at an applied current density of only 4.14 A m⁻² at a hydraulic residence time of 1.5 min. Under these conditions, this benchtop system could process approximately 120 L of water per day while consuming approximately 1.7 Wh to produce H₂O₂.

Trace Organic Contaminant Removal by Electro-generated H₂O₂ and UV Irradiation. The removal of electro-contaminants involved direct photolysis, reactions with HO⁺ produced by photolysis of H₂O₂ in the UV chamber, and direct oxidation of contaminants on the anode. At the fluence employed in the UV chamber (F₀ ~ 3000 mJ cm⁻²), direct photolysis only removed those compounds that exhibited high quantum yields and strong light absorbance at 254 nm (Figure S4, Supporting Information). Among the compounds tested, carbamazepine exhibited the lowest tendency for direct transformation by UV light (<30% removal for all matrices), while more photoreactive compounds, such as sulfamethoxazole, propranolol, and atrazine, displayed higher removals (55–99%). Unlike the variability of the compounds with respect to direct photolysis, the suite of trace organics all reacted with HO⁺ at near diffusion controlled rates (10⁻⁸–10⁻¹⁰ M⁻¹ s⁻¹; Table S3, Supporting Information). As a result of its low reactivity with UV light, carbamazepine removal in the presence of H₂O₂ and UV light provided useful information on the transformation of organic contaminants by HO⁺.

The extent of removal of carbamazepine varied among the different matrices (Figure 2). The presence of HO⁺ scavengers explained much of the variability. In the absence of current (and therefore H₂O₂), the variability of carbamazepine transformation was predominately influenced by the screening of light, as accounted for by the water factor:²⁰²⁴

$$\text{water factor} = \frac{1 - 10^{-1.2\alpha z}}{(2.3)(1.2)\alpha z}$$  (4)

where z is the mixed water body depth (m) and α is the attenuation coefficient of the water body. The water factor, which was primarily influenced by the amount of NOM, followed the trend: electrolyte (0.998) > groundwater (0.985) > synthetic surface water (0.958) > wastewater effluent (0.508). Transformation of carbamazepine solely in the presence of UV light varied from 22 ± 5% in the electrolyte solution to 15 ± 4% in the wastewater effluent. The observed removal of carbamazepine in the surface water and wastewater effluent, however, was greater than suggested from the water factor. This may be explained by the generation of HO⁺ and DOM⁺ produced from NOM sensitization by UV light, which can be significant at UV fluences employed in AOPs.²⁵²⁹

The rate of transformation of trace contaminants increased with current density due to additional HO⁺ production that occurred at higher H₂O₂ concentrations (Tables S4–S7, Supporting Information). Complete carbamazepine transformation was observed after the UV treatment chamber at an applied current density of 5 A m⁻² for the electrolyte. For the three representative source waters, carbamazepine transformation increased to 98.7 ± 0.6% for groundwater, 93.3 ± 1.0% for surface water, and 78.5 ± 1.9% for wastewater effluent as the current increased to 25 A m⁻². Organic compounds that have lower reaction rate constants with HO⁺ and are not susceptible to direct photolysis will require higher current densities to achieve a similar level of treatment.

The fraction of HO⁺ that reacted with the contaminants can be estimated by considering the concentrations and rate constants for reactions of different solutes with HO⁺:

$$\text{fraction HO⁺ to contaminants} = \frac{\sum k_{\text{OM}⁺,\text{cont}}[\text{cont}]}{\sum k_{\text{HO}⁺,\text{OM}}[S]}$$  (5)

where $k_{\text{OM}⁺,\text{S}}$ and $k_{\text{HO}⁺,\text{cont}}$ are the second order reaction rate constants of scavengers and contaminants with HO⁺, respectively, and [S] is the concentration of the scavenger (e.g., HCO₃⁻, CO₃²⁻, NOM, H₂O₂; Tables 1 and S3, Supporting Information).

Effect of H₂O₂ on Treatment Efficiency. At increasing H₂O₂ concentrations, a trade-off exists between additional transformation of trace organics by HO⁺ produced from H₂O₂ photolysis and greater radical scavenging and light screening by H₂O₂.²⁷ Therefore, despite linear increases in H₂O₂ production with current density, there is a diminishing benefit to the treatment. As H₂O₂ increased from 0.09 mM (4.14 A m⁻²) to 0.54 mM (25 A m⁻²), the fraction of HO⁺ reacting with contaminants decreased by 20%, 21%, and 10% for the surface water, groundwater, and wastewater effluent, respectively. At 0.54 mM H₂O₂ (25 A m⁻²), there was a 4.0%, 4.7%, and 3.8% reduction in direct photolysis rates of contaminants from additional light screening by H₂O₂ for the surface water, groundwater, and wastewater effluent, respectively (details of

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**Figure 2.** Removal of carbamazepine as a function of current density for the four types of source waters (WWTP: wastewater treatment plant).
branching ratio and direct photolysis calculations are included in the Supporting Information).

**Effect of pH on Treatment Efficiency.** At pH 8, approximately 6.7%, 6.5%, and 2.9% of HO• reacted with the organic contaminants in the UV reactor at an initial H₂O₂ concentration of 3 mg L⁻¹ (0.09 mM) for the surface water, groundwater, and wastewater effluent, respectively. At pH 10, the fraction of HO• reacting with trace organic contaminants decreased to 0.9%, 0.6%, and 0.4% for the three source waters, respectively. The significant decrease in HO• reacting with the trace organic contaminants was due to scavenging by carbonate at the higher pH values. The product of this reaction, CO₃²⁻, can play a significant role in the transformation of certain organic compounds (e.g., propranolol and sulfamethoxazole; Table S3, Supporting Information).²⁸ As a result of differences in alkalinity of the different source waters, the importance of carbonate scavenging and CO₃⁻ reactions depends on the source water composition and the applied current density (i.e., higher applied currents result in greater pH increases in the cathode chamber). Although nitrite is an effective scavenger of HO• (*k*₅(NO₂⁻, HO•) = 6 × 10⁹ M⁻¹ s⁻¹), less than 0.3% of the generated HO• would be scavenged by nitrite at concentrations typically found in nitrified wastewater effluent (i.e., ~0.1 mg L⁻¹). The formation of halogen radicals from reactions between HO• and halide ions (chloride and bromide) should only be significant at low pH values and therefore will have a negligible impact on contaminant transformation at the circumneutral and basic pH values observed in this system.²⁹

**Anodic pH Adjustment.** The generation of H₂O₂ by the cathode consumed protons and increased the solution pH (reaction 6). In the anode, oxidation reactions produced protons and lowered the solution pH (reaction 7):

O₃ + 2e⁻ + 2H⁺ ⇌ H₂O₂ (cathode)  

H₂O ⇌ 1⁄2O₂ + 2e⁻ + 2H⁺ (anode)  

2Cl⁻ ⇌ Cl₂ + 2e⁻ (anode)  

Cl₂ + H₂O ⇌ HOCl + H⁺ + Cl⁻ (9)  

For the production of 3 mg L⁻¹ (0.09 mM) of H₂O₂ at a current density of 4.14 A m⁻², approximately 0.20 mequiv L⁻¹ of protons should have been consumed or produced at the cathode and anode, respectively. To maintain electroneutrality, a net migration of protons occurred from the anode chamber to the cathode chamber via the cation exchange membrane. In addition to protons, cations that were present at higher concentrations (e.g., Na⁺, Ca²⁺, Mg²⁺) also carried ionic charge through the membrane satisfying electroneutrality in the cathode chamber while creating a proton deficit in the cathode chamber.³⁰

As a result of differences in buffering among matrices, the solution pH should have increased more in the cathode chamber for waters with low alkalinity. The pH in the cathode chamber increased for each of the waters as the current increased from 0 to 25 A m⁻² (Figure 3). The magnitude of pH increase was most pronounced for the electrolyte and surface water (alkalinity = 0 and 2.45 mM, respectively) with post-cathode pH values ranging from 10 to 10.5, while the pH never exceeded 9.9 and 9.2 in the groundwater (alkalinity = 3.89 mM) and the municipal wastewater effluent (alkalinity = 4.97 mM), respectively. The pH increases following the cathode resulted in supersaturation with respect to calcite (CaCO₃(s)) in the groundwater (log SI > 1.31), surface water (log SI > 1.58), and wastewater effluent (log SI > 1.09) beginning at a current density of 5 A m⁻². This reaction could result in scaling on the cathode or the ion exchange membrane that might eventually affect system performance. Loss of CaCO₃(s) from the system could also result in an overall decrease in pH as water passed...
through the treatment system. For example, if the surface water solution reached equilibrium at the current density needed to produce 3 mg L$^{-1}$ (0.09 mM) H$_2$O$_2$, 0.74 mmol (74 mg) of calcite would precipitate for each liter of water treated and the pH would have dropped from 9.82 to 7.23. On the basis of the observed pH values, it is evident that equilibrium was not achieved. However, additional research is needed to assess the importance of calcite precipitation to scaling and pH control.

To readjust the solution pH, water leaving the UV reactor was passed through the anode chamber. If no mineral precipitation occurred in the cathode and UV chambers, the final pH should have been equal to the influent pH. A slight decrease in pH was observed in all solutions to which current was applied, with greater pH decreases at higher current densities occurring in the least buffered of the three environmental matrices (i.e., surface water). Under the conditions that would likely be used for treatment (i.e., $30-100$ A m$^{-2}$ and short hydraulic residence times), the final pH was approximately equal to the initial pH.

### Anodic Quenching of Residual Hydrogen Peroxide.

Due to the relatively low molar absorptivity of H$_2$O$_2$ at 254 nm ($\varepsilon_{254} = 18.6$ M$^{-1}$ cm$^{-1}$) and the limited residence times in the UV reactor ($\tau = 660$ s), much of the H$_2$O$_2$ passed through the UV chamber without undergoing photolysis. For solutions with relatively low light screening (i.e., electrolyte, synthetic groundwater, and synthetic surface water), between 40% and 50% of the H$_2$O$_2$ was photolyzed at current densities ranging from 5 to 25 A m$^{-2}$ (Figure 4 and Figure S5, Supporting Information). As expected, less H$_2$O$_2$ photolysis occurred in municipal wastewater effluent due to light screening.

![Graph showing production of H$_2$O$_2$ in the cathode, residual H$_2$O$_2$ after the UV cell, and residual H$_2$O$_2$ after the anode for four types of source waters at applied current density of 25 A m$^{-2}$](image)

**Figure 4.** Production of H$_2$O$_2$ in the cathode, residual H$_2$O$_2$ after the UV cell, and residual H$_2$O$_2$ after the anode for four types of source waters at applied current density of 25 A m$^{-2}$. See Figure S5, Supporting Information, for data on the production and removal of H$_2$O$_2$ over the full range of current densities (5–25 A m$^{-2}$) (WWTP: wastewater treatment plant).

In practice, many centralized treatment plants employ reducing agents (e.g., bisulfite), chlorine, or activated carbon to remove residual H$_2$O$_2$ before distribution.$^{31}$ The use of activated carbon or the addition of chemicals, however, may be impractical in a distributed treatment system. Partial removal of H$_2$O$_2$ occurred when the solution passed through the anode, especially in the electrolyte solution (Figure 4 and Figure S5, Supporting Information). Anodic removal of H$_2$O$_2$ increased with increasing current density (Figure S5, Supporting Information). In the NaCl electrolyte solution, up to 0.12 mM H$_2$O$_2$ was removed at 25 A m$^{-2}$. For the three source waters, however, the anode only removed about 25% of the amount removed in the electrolyte.

Removal of H$_2$O$_2$ in the anode was attributable to a combination of direct anodic oxidation and reactions with oxidants produced on the anode surface. For example, oxidation of chloride can result in the production of hypochlorous acid (HOCl; $pK_a = 7.6$) (Reactions 8 and 9). Hypochlorite reacts rapidly with hydrogen peroxide under alkaline conditions with the bimolecular rate constant increasing from $196$ to $7.5 \times 10^7$ M$^{-1}$ s$^{-1}$ from pH 6 to 9 (see the Supporting Information for calculation of the pH-dependent bimolecular rate constant).$^{32-34}$

$$\text{OCl}^- + \text{H}_2\text{O}_2 \rightarrow \text{Cl}^- + \text{O}_2 + \text{H}_2\text{O} \quad (10)$$

Although Ti-IrO$_2$ electrodes have a high electrocatalytic activity with respect to chlorine evolution, only modest concentrations of chlorine were produced in control experiments at varying chloride concentrations due to the short hydraulic residence times and relatively low current densities applied$^{35-37}$ (Table 2). To separate the effects of reactive halogen species from direct electrode oxidation on the removal of H$_2$O$_2$, experiments were repeated using an inert electrolyte (i.e., Na$_2$SO$_4$) (Figure S6, Supporting Information). H$_2$O$_2$ removal was independent of applied current density with 37 ± 2 μM H$_2$O$_2$ removed from 5 to 25 A m$^{-2}$; a concentration equivalent to the observed H$_2$O$_2$ removal in the anode for the three source waters in Figure S5, Supporting Information. Given the low chloride concentrations (<1 mM) of the simulated surface water and groundwater, it is not surprising that OCl$^-$ production was low and only a small quantity of H$_2$O$_2$ was removed in the anode.

Despite the electrolyte and municipal wastewater effluent having roughly the same concentration of chloride, H$_2$O$_2$ removal was significantly higher in the electrolyte control than in the wastewater effluent, suggesting that reactive halogen species did not play an important role in H$_2$O$_2$ removal in the wastewater effluent matrix. NOM is an effective sink of HOCl/OCl$^-$; however, under the experimental conditions used in this study, the half-life of HOCl/OCl$^-$ with respect to its reaction with H$_2$O$_2$ was much shorter than that predicted for NOM (i.e., 1.39 and 49.3 s, respectively, as described in the Supporting Information).$^{38}$ As a result, NOM is only a minor sink for HOCl/OCl$^-$ in the presence of H$_2$O$_2$. This was consistent with observations from experiments in which H$_2$O$_2$ removal decreased by less than 40% (58 μM) when NOM was added to a solution containing a fixed concentration of chlorine at a current density of 25 A m$^{-2}$, suggesting that NOM is only

### Table 2. Chloride-Chlorine Electrochemical Oxidation

| current density (A m$^{-2}$) | [Cl$^-$ (mM)] |
|-----------------------------|---------------|
|                             | 2.5 | 5  | 10 | 15 | 25 |
| 0.5                         | 0   | 0  | 0  | 0  | 0  |
| 5                           | 0   | 0  | 0.79 | 17 | 34 |
| 10                          | 0   | 0  | 3.9 | 25 | 41 |
| 15                          | 0   | 0  | 7.9 | 32 | 61 |

$^{a}$Free chlorine production (as μM Cl$^-$) in the anode chamber as a function of applied current density and chloride concentration. Experiments were performed with a stainless steel cathode to prevent H$_2$O$_2$ formation, which interferes with the chlorine measurement.
partially responsible for the difference in $\text{H}_2\text{O}_2$ removal between the two solutions (Figure S6, Supporting Information).

The apparent discrepancy between municipal wastewater effluent and the sodium chloride solution may be partially attributable to differences in pH values in the anode chamber.36 At higher pH values, like those found in the municipal wastewater effluent after anodic treatment, there is a larger driving force for oxygen evolution compared to $\text{Cl}_2$ production:

$$\text{O}_3 + 4\text{H}^+ + 4e^- \rightarrow \text{H}_2\text{O} \quad E_{\text{H}^+}^0 = 1.23\text{V} \quad (11)$$

$$\text{Cl}_2 + 2e^- \rightarrow 2\text{Cl}^- \quad E_{\text{Cl}^-}^0 = 1.36\text{V} \quad (12)$$

Experiments conducted in the anode chamber at different pH values confirmed that chlorine production increased substantially as pH dropped from 9 to 7 (Figure S7, Supporting Information). Due to the presence of bicarbonate in the municipal wastewater effluent, the anode pH was considerably higher (i.e., ∼8) during treatment than in the unbuffered NaCl electrolyte, where pH decreased to approximately 6 during anodic treatment. As a result, considerably more HOCl/OCl$^-$ was produced in the anode chamber when the electrolyte was treated.

Total hydrogen peroxide removal in the system (i.e., photolysis and anodic loss) was 48 ± 5% for the simulated groundwater, 49 ± 3% for the simulated surface water, and 25 ± 3% for the wastewater effluent for the array of current densities tested. At the current density required to produce 3 mg L$^{-1}$ (0.09 mM) $\text{H}_2\text{O}_2$ water leaving the treatment system effluent contained 1.5–2.3 mg L$^{-1}$ $\text{H}_2\text{O}_2$. Although $\text{H}_2\text{O}_2$ does not pose a health risk at these concentrations, its presence in tap water may be undesirable. In a point-of-use water treatment system, it might be possible to remove the excess $\text{H}_2\text{O}_2$ by passing it through activated carbon or a high surface area catalyst such as metal oxide or silver. Alternatively, the efficiency of chloride oxidation in the anode chamber might be improved through the use of three-dimensional or porous electrodes that reduce mass transfer limitations or through the use of more catalytic anode materials.5

**Anodic Transformation of Trace Organic Contaminants.** Despite only accounting for a small fraction of the total removal observed in the treatment system, the anode did transform some of the compounds. Experiments conducted in the absence of UV exposure with solutions amended with 10 $\mu$g L$^{-1}$ of trace organic contaminants indicated that direct anodic oxidation resulted in the removal of up to 20% of certain trace organics (e.g., propranolol) at a current density of 25 A m$^{-2}$ (Figure S8, Supporting Information). Oxidation of organic compounds on the anode could be increased through the use of inactive anodes (e.g., boron-doped diamond, doped-SnO$_2$, PbO$_2$).5,37 These electrodes, however, have higher capital and operating costs than Ti-IrO$_2$ electrodes.

Although carbamazepine is relatively unreactive with hypochlorite, other compounds (e.g., propranolol and sulfamethoxazole) react with HOCl (Table S3, Supporting Information). However, the presence of $\text{H}_2\text{O}_2$ reduced the importance of reactions between trace organics and chlorine species generated at the anode because chlorine preferentially reacts with $\text{H}_2\text{O}_2$. As a result, nearly all of the observed loss of the test compounds in the anode chamber was due to direct oxidation on the anode surface. This observation was consistent with experiments comparing anodic removal of trace organics in the presence and absence of $\text{H}_2\text{O}_2$ (Figure S9, Supporting Information).

**Long-term Cathode Performance.** The performance of gas diffusion electrodes can decrease over time due to clogging of the pores by precipitates, fouling with NOM as well as charge transfer resistance attributable to the loss of conductive graphite paste.42 In a long-term trial, cathode performance (i.e., $\text{H}_2\text{O}_2$ production at a fixed current density) decreased by less than 2% after 6000 L of tap water amended with 5 mM Na$_2$SO$_4$ was passed through the system at an applied current density of 15 A m$^{-2}$ (Figure S10, Supporting Information). Calcium carbonate scaling due to the elevated pH and migration of calcium ions in the tap water into the cathode chamber was observed on the interior of the cathode. Nonetheless, $\text{H}_2\text{O}_2$ production was unaffected during this 50 day test. Additional experiments are needed to assess the importance of scaling and the efficacy of simple descaling approaches (i.e., polarization reversal) over longer time periods and more realistic operating conditions.

**System Energy Consumption.** The treatment system used electricity to produce the oxidant (i.e., $\text{H}_2\text{O}_2$) and to convert it into HO$^*$ (i.e., the UV lamp). Electrical energy per order ($E_{\text{EO}}$) is a useful figure of merit for comparing the efficiency and cost of the treatment system with other AOPs. $E_{\text{EO}}$ is the electrical energy (in kWh) required to reduce a contaminant concentration by 1 order of magnitude in 1 m$^3$ of water:43–45

$$E_{\text{EO}} = \frac{P}{Q \log(C_0/C)} \quad (13)$$

where $P$ (kW) is the electrical power for the electrochemical cell and UV lamp, $Q$ (m$^3$ h$^{-1}$) is the system flow rate, and $C_0$ and $C$ (M) are the initial and final contaminant concentrations. Without the production of $\text{H}_2\text{O}_2$, $E_{\text{EO}}$ values ranged from 16.8 ± 0.3 to 28.1 ± 0.2 kWh m$^{-3}$ order$^{-1}$, with larger amounts of energy needed to transform contaminants in waters that contained high concentrations of HO$^*$ scavengers and chromophores (i.e., municipal wastewater effluent and synthetic groundwater) (Figure 5). A substantial decrease in $E_{\text{EO}}$ occurred when current was applied to the electrochemical cell (i.e., from 0 to 5 A m$^{-2}$). As current density increased from 5 to 25 A m$^{-2}$, the $E_{\text{EO}}$ decreased by less than 11%. These data

![Figure 5](Image 340x101 to 549x258)
indicate that the UV–H₂O₂ AOP system is much more efficient than the use of UV alone and that there is a marginal benefit associated with the production of higher H₂O₂ concentrations in the cathode because the reactions become less efficient at higher concentrations of H₂O₂. Only 8–14% of the energy demand was attributable to the electrochemical production of H₂O₂, with the majority of the energy required for the operation of the low-pressure UV lamp. At a current density of 25 A m⁻², an energy requirement of 1.08 ± 0.1 to 2.84 ± 0.1 kWh m⁻³ order⁻¹ was observed, which was similar to results from previous studies of the transformation of trace organic compounds by UV/H₂O₂ in different source waters.¹⁰

The high Coulombic efficiency and low current densities result in a low-cost means of H₂O₂ production even for treatment of water with low conductivity. For comparison, electrochemically produced H₂O₂ costs between 0.1 and 0.3 $ kg⁻¹, while H₂O₂ produced by the anthraquinone process typically costs between 1 and 2 $ kg⁻¹.¹⁶ When considering the lack of a need to transport, store, and handle H₂O₂ as well as modest capital and operational costs, the modular AOP treatment system can be a competitive technology for point-of-use treatment at a household and community level or even for wellhead treatment of trace organic contaminants present in potable water sources. As H₂O₂ production was directly proportional to current density, the treatment system could be scaled up by using faster flow rates accompanied by higher applied current densities or by increasing surface area of the cathode. To obtain the equivalent contaminant removal after scale-up, optimization of the UV reactor geometry would be required. Additional research is needed to assess long-term system performance under realistic operating conditions.

### ASSOCIATED CONTENT

#### Supporting Information

Details of the calculations for fluence rate, photolysis rates, H₂O₂ half-life, oxidant branching ratios, EEO, and methods for test compound analysis in addition to supporting tables, schematics, and figures referenced in this work. The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acs.est.5b01254.

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**Notes**

The authors declare no competing financial interest.

### ACKNOWLEDGMENTS

This study was supported by the U.S. National Institute for Environmental Health Sciences (NIEMS) Superfund Research Program (Grant P42 ES004705) and the Superfund Research Center at University of California, Berkeley. T.H. was also supported by a postdoctoral fellowship from the Research Foundation Flanders (FWO Vlaanderen). The authors thank Dr. Oskar Modin for his guidance with the fabrication of the gas diffusion electrode and Dr. Jelena Radjenovic for her insightful comments.

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