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## Delineating the Effects of Molecular and Colloidal Interactions of Dissolved Organic Matter on Titania Photocatalysis

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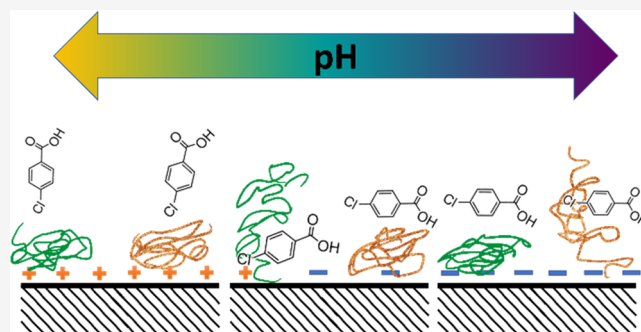


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**ABSTRACT:** In the face of significant challenges to practical applications of photocatalysis for water treatment, recent reports revealed a potential route to overcome a problem posed by dissolved organic matter (DOM). These studies showed that inhibition of photocatalytic processes by DOM is driven largely by competition for active surface sites on  $\text{TiO}_2$  or other catalysts, and controlling the type of DOM present in solution could significantly mitigate DOM fouling. Whether or not control of solution parameters could achieve the same preventative action is not known. Here, a series of DOM isolates, including humic acid (HA) and transphilic (TPI), hydrophobic (HPO), or colloidal fractions of organic matter from a membrane bioreactor mixed liquor supernatant, were tested for inhibitory activity under a range of pH values (3, 5, 7, and 9) and ionic compositions ( $\text{NaCl}$ ,  $\text{CaCl}_2$ , and  $\text{Al}_2(\text{SO}_4)_3$  with ionic strengths (IS) ranging from 0 to 3 M). The resulting  $\text{TiO}_2$ -DOM agglomerates were monitored for size and  $\zeta$ -potential. Inhibitory profiles were generated using *para*-chlorobenzoic acid (*pCBA*) as probe with varying concentrations of inhibitory DOM for each solution condition to discern the extent of surface-phase quenching of radicals. Manipulation of pH clearly impacted inhibition, and the effect varied by DOM type; for example, interference occurred at all pHs for HA, at neutral or basic pHs for TPI, and only at pH 7 for HPO. Particle sizes did not correlate with inhibitory action of DOM. Increases in ionic strength induced growth of  $\text{TiO}_2$  and  $\text{TiO}_2$ -DOM agglomerates, but again, particle sizes did not correlate to inhibition by DOM. The changes to IS, regardless of ion type, were not affected by the presence of TPI or HPO. Since particle stability did not correlate directly with photocatalytic activity, we suggest that surface-based quenching reactions arise from site-specific adsorption rather than generalized particle destabilization and aggregation.



### INTRODUCTION

Access to clean and safe drinking water is a critical challenge for people around the world, especially those who live in rural areas.<sup>1,2</sup> Photocatalytic technologies may become a sustainable water treatment solution in some situations, if certain hurdles are overcome.<sup>3,4</sup> The well-studied  $\text{TiO}_2$  offers an attractive option owing to its wide availability and high photocatalytic activity,<sup>3,5–8</sup> but  $\text{TiO}_2$ -based photocatalysis has yet to be utilized in practical water treatment systems.<sup>7</sup> One of the primary barriers to commercialization is the mass transfer limitation of reactive oxygen species (ROS) to target contaminants,<sup>5</sup> especially in complex water matrices.<sup>9</sup> In fact, a 2019 study showed that less than 5% of generated  $\cdot\text{OH}$  radical in  $\text{TiO}_2$  photocatalysis is available for the degradation of target pollutants in natural waters due to interference of different solutes.<sup>8</sup> Dissolved organic matter (DOM) is one of the most detrimental constituents because it inhibits photocatalytic processes competitively reacting with ROS and adsorbing to active surface sites for surface-phase reactions.<sup>3,7,10</sup> Even in pure water, only a small portion of ROS produced by  $\text{TiO}_2$  diffuse away from the surface into the bulk solution,<sup>11</sup> so the surface interactions between DOM and

targeted pollutants are critically important. For successful application of photocatalysis, control over the type of DOM present in the process water is essential.

Several recent studies aimed to better understand the inhibitory mechanisms of DOM in photocatalytic systems. In 2015, Brame et al. experimentally validated an analytical model for accurately simulating photocatalytic performance of two different photoactive materials ( $\text{TiO}_2$  and  $\text{Si-C}_{60}$ ) when inhibited by either bulk or surface-phase scavenging; they concluded that the adsorption of DOM on the surface of the photocatalyst caused the most substantial reductions in performance; just 5 mg/L of Suwannee River natural organic matter was sufficient to reduce the  $\text{TiO}_2$ -photodegradation rate for two probe molecules by half or more.<sup>3</sup> They also noted that

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the adsorption affinity of DOM predicted inhibition when using a combined surface (Langmuir) and bulk inhibition model.<sup>3</sup> This dynamic was later shown to be important in understanding the inhibitory effects of DOM in wastewater treatment;<sup>12</sup> effluent organic matter from clean membrane decreased the photocatalytic efficiency by 100%, against 33% for the fouled membrane. This difference was explained by a higher permeation of colloidal DOM through the clean membrane, compared to the fouled membrane.<sup>4</sup> In 2018, Luo et al. supplied an example of this phenomenon by showing that more humic acid (HA) adsorbed onto a TiO<sub>2</sub> photocatalyst surface than fulvic acid (FA), due to higher hydrophobicity of HA, and consequently caused more inhibition.<sup>13</sup> Similarly, we demonstrated that fractionated organic matter from activated sludge exhibited distinct inhibitory profiles, with the colloidal fraction causing more inhibition than transphilic (TPI) or hydrophobic (HPO) isolates.<sup>4</sup>

The progress in understanding photocatalyst quenching based on the nature of DOM is promising for addressing unwanted inhibitory reactions, but obvious questions remain regarding the roles of solution chemistry and cosolutes. At a basic level, simple colloidal physics could explain DOM's inhibitory effects; for example, elevated ionic strength could induce aggregation of TiO<sub>2</sub> particles with DOM via compression of the electrical double layer, which could increase surface quenching reactions. The situation may be more complex, however, when examined at the level of molecular-scale interactions. In a 2014 report, researchers showed that on positively charged surfaces, DOM (humic and fulvic acids) forms irreversibly sorbed, nanometer-scale adlayers.<sup>14</sup> These DOM layers likely form on positively charged TiO<sub>2</sub> below its isoelectric point, typically between 6 and 6.5.<sup>7</sup> However, the impact of pH on DOM–TiO<sub>2</sub> interactions is also complicated by the wide array of moieties constituting DOM, with numerous different pK<sub>a</sub> values for the acidic groups.<sup>15</sup> Alternately, DOM can serve as a stabilization agent for metal oxide nanoparticles via steric hindrance.<sup>16,17</sup> In addition, ionic cosolutes may induce morphological changes in DOM; researchers have shown that the macromolecular structure of HA depended significantly on the type of cation in solution, with divalent cations exerting a stronger effect on size and conformation than monovalent cations.<sup>18</sup> Little is known about how these ionic effects, whether particle stability or DOM conformational changes, impact photocatalysis in complex matrices. Control of intermolecular interaction dynamics may provide tools to manipulate the adsorption dynamics of DOM onto TiO<sub>2</sub> by simple operational changes (e.g., pH adjustments or ion exchange processes) to mitigate surface-phase inhibition of photocatalysis.

The question of whether colloidal physics can predict the detrimental surface-phase quenching of photocatalysts by DOM remains outstanding. Here, particle stability experiments and inhibitory profiles of TiO<sub>2</sub> photoactivity are used as tools to observe how changes to pH, ionic strength, or specific ionic constituents affect DOM–TiO<sub>2</sub> adsorption and the consequent impacts on photocatalysis. Parameters are tested across colloidal stability thresholds to intentionally induce aggregation and co-precipitation of DOM and TiO<sub>2</sub> in solution.

## MATERIALS AND METHODS

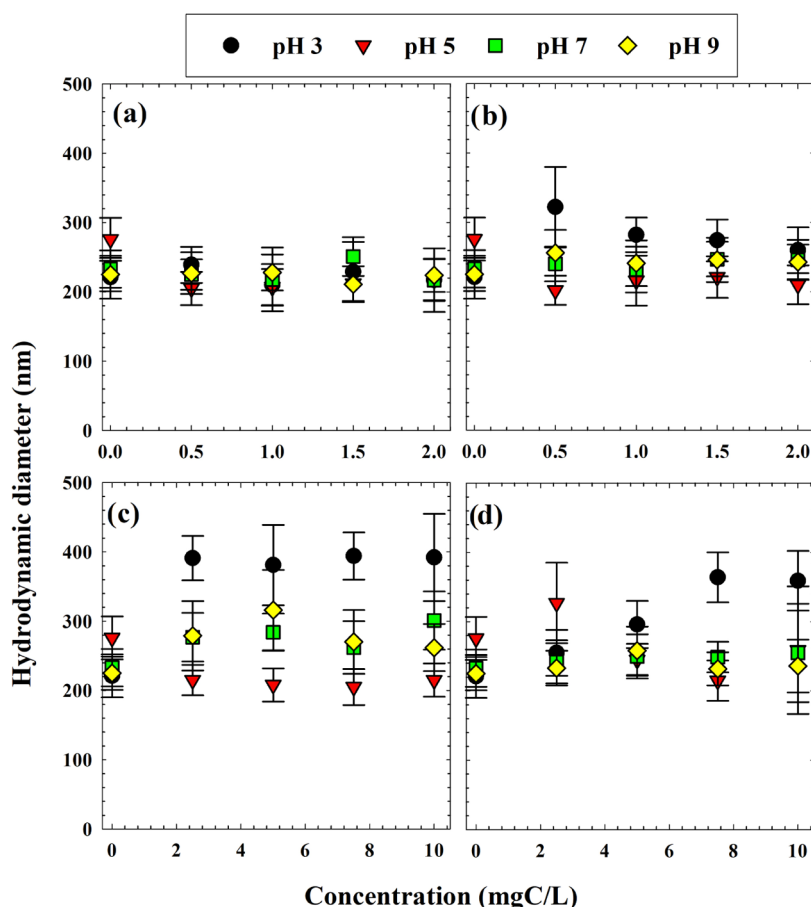
**Chemicals.** Humic acid, 4-chlorobenzoic acid (*p*CBA), and titanium dioxide (99.9% Anatase) with a nominal particle size of 32

nm and surface area of 45 m<sup>2</sup> g<sup>−1</sup> were obtained from Alfa Aesar (Haverhill, MA). Hydrochloric acid (HCl), sodium hydroxide (NaOH), magnesium sulfate (MgSO<sub>4</sub>), aluminum sulfate (Al<sub>2</sub>(SO<sub>4</sub>)<sub>3</sub>), sodium chloride (NaCl), and calcium chloride (CaCl<sub>2</sub>) were all purchased from VWR (Radnor, PA). A Nanopure Infinity system (Thermo Fisher Scientific, Inc., Waltham, MA) was used to produce ultrapure water (>18.2 MΩ cm). HPLC solvents including acetonitrile and phosphoric acid were purchased from Alfa Aesar, and all of them were of HPLC grade.

**Activated Sludge Organic Matter Collection.** Three DOM fractions were used in this study including colloidal (C), transphilic (TPI), and hydrophobic (HPO) fractions. These isolates were obtained by processing the bulk supernatant samples collected from a full-scale membrane bioreactor wastewater treatment plant (La Grande Motte, France), as described previously.<sup>19</sup> The DOM fractionation process was performed on the bulk supernatant obtained from a prefiltered activated sludge (1 μm) and concentrated with reverse osmosis membranes (TW30 Filmtech membranes). Then, the DOM fractions were isolated through dialysis bags (molecular weight cutoff, 3.5 kDa), XAD8 and XAD4 resins to collect colloidal, HPO, and TPI fractions, respectively.<sup>4,20</sup> All fractions were finally freeze-dried for further use; thorough characterization of each has been performed previously.<sup>19</sup>

**Photochemical Experiments.** Photodegradation experiments were performed in an enclosed UV cabinet equipped with a UV LED (LG Innotek UVC 6868, South Korea) with an emission peak at 278 nm (UV<sub>278</sub>) placed 20 cm from the reaction vessel, which was placed on a magnetic stirrer. To achieve quasi-collimated irradiation, a black tube was used to isolate rays from one of several lamps within the cabinet and mitigate reflection. An intensity value of 722 μW·cm<sup>−2</sup> was measured for the UV LED using a BLUE–Wave UVNB-25 Spectrometer (StellarNet, Inc., Tampa, FL). The UV emission spectrum for the LED was reported previously.<sup>21</sup> The vessel contained 10 mL solution, 5 mg L<sup>−1</sup> TiO<sub>2</sub> with 10 μM *p*CBA as a probe compound, which has a known reaction rate constant with •OH.<sup>22</sup> Inhibitory profiles were constructed by determining *p*CBA degradation rate constants across a series of DOM concentrations for each DOM sample (HA or the fractionated activated sludge isolates). Solution pHs were adjusted to 3, 5, 7, or 9 using HCl or NaOH when needed. The ionic strengths of solutions were adjusted from 0 to 3 M by adding salts with cations of varying valencies, NaCl, CaCl<sub>2</sub>, or Al<sub>2</sub>(SO<sub>4</sub>)<sub>3</sub>, when evaluating the effects of ionic strength or ion type on photodegradation rates. Samples were withdrawn at fixed time points for *p*CBA quantification via HPLC (Agilent Technologies, Inc., 1260 infinity). A C18 (125 mm) column was used in the analysis, and acetonitrile and 10 mM phosphoric acid were the mobile phase solvents (40:60) with a flow rate of 0.5 mL·min<sup>−1</sup>. The HPLC instrument had a variable-wavelength detector, which was adjusted to 234 nm for the detection of *p*CBA.<sup>23</sup> *p*CBA degradation rates were calculated for all of the experiments using linear regression in the plot of the natural log of *p*CBA concentration versus fluence, which followed first-order kinetics as demonstrated previously.<sup>4,12</sup> Fluence values were calculated from intensity measurements according to Bolton and Linden,<sup>24</sup> and each value accounted for reductant transmission factors: Petri, water, divergence, and reflection.

**Aggregate Characterization.** The hydrodynamic diameter and ζ-potential of pristine TiO<sub>2</sub> and (presumed) TiO<sub>2</sub>–DOM aggregates were measured using dynamic light scattering (DLS) and phase analysis light scattering (PALS) using a Malvern Zetasizer Nano ZS90 (Malvern Instruments, Worcestershire, U.K.). TiO<sub>2</sub> solutions were sonicated for 5 min before adding to a given solution and mixed with an experimental solution for 2 min before taking DLS or PALS measurements. PALS measurements were performed in a folded capillary cell. A refractive index of 2.524 was assigned to TiO<sub>2</sub> nanoparticles.<sup>25</sup> Error values for size and ζ-potential represent standard errors computed by the Malvern Zetasizer Software, based on a minimum of 10 measurement runs per sample and two samples per datum. Aggregate size and ζ-potential measurements were compared across experimental conditions using three-factor ANOVA tests using SigmaPlot v. 14.0 software (Systat Software,



**Figure 1.** Hydrodynamic diameters of TiO<sub>2</sub> particles at pH 3, 5, 7, and 9 in the presence of (a), HA (b), colloids (c), TPI, and (d) HPO. Error bars indicate standard error.

Inc., San Jose, CA) with significance defined as  $p$ -value < 0.05. Three ANOVA tests were performed: one for the size of particles based on changes in DOM concentration, DOM type, and pH; second under the same categories for  $\zeta$ -potential; and third for the size of particles with changing IS, ion type, and DOM types.

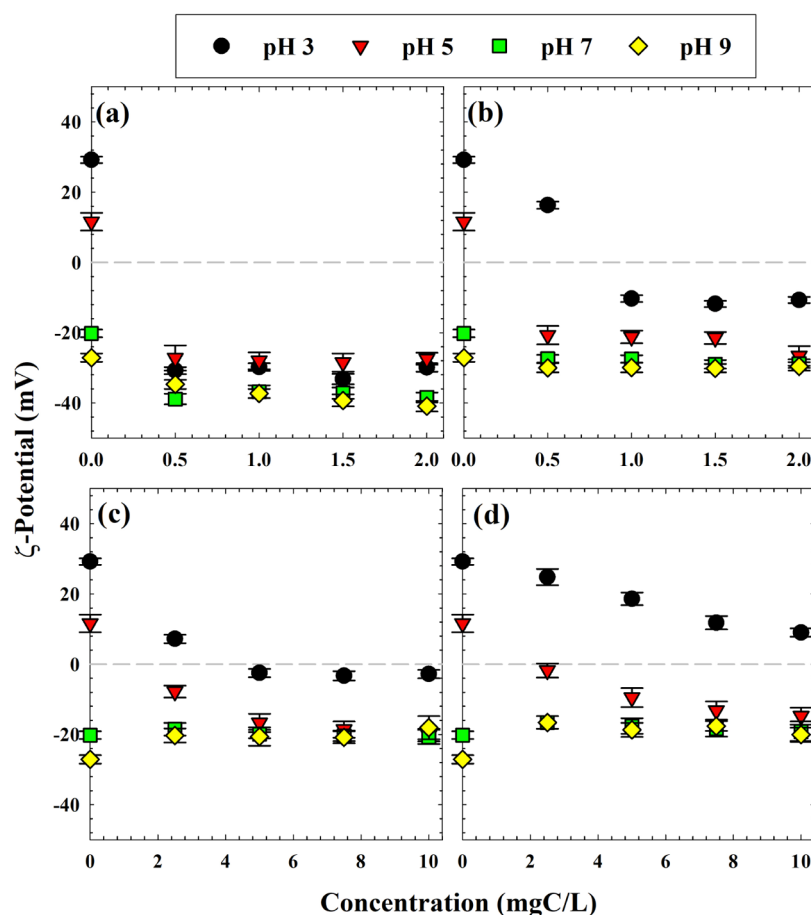
## RESULTS AND DISCUSSION

**Impacts of DOM and pH on Particle Size and  $\zeta$ -Potential Variations.** To determine the role of pH in the aggregation of TiO<sub>2</sub> in the absence or presence of DOM fractions, the hydrodynamic diameter and  $\zeta$ -potential of TiO<sub>2</sub> were measured at various pHs. Particle  $\zeta$ -potential measurements were conducted across DOM concentrations found to be meaningful to photocatalytic quenching in our previous work, up to 2.0 mgC/L for HA and C or 10 mgC/L for TPI and HPO (these concentrations were categorized as lowest, low, high, and highest for ANOVA purposes).<sup>4</sup> Particle size data are shown in Figure 1 for each DOM isolate across pH values. In general, hydrodynamic diameters varied little with DOM concentration; mean sizes are recorded in Table S1, with corresponding ANOVA results shown in Tables S2–S4. DOM type and pH were both found to significantly ( $p$ -value < 0.05) alter aggregate size, while DOM concentration did not (Table S2). Pairwise comparisons between factor groups revealed (Table S3) that particle sizes for HA differed significantly from both TPI and HPO but not from colloids. TPI was also found to be statistically different compared to colloids. At pH 3, TiO<sub>2</sub>-DOM particle size differences were significant (>40 nm differences in means), but no significant

difference was found for any other pH pairing (<12 nm differences in means).

Particle sizes (Figure 1) were generally inversely correlated with  $\zeta$ -potential magnitudes, which are plotted in Figure 2 and enumerated in Table S5. Corresponding ANOVA results are shown in Tables S6–S8. The isoelectric point of colloidal TiO<sub>2</sub> is typically between pH 6.0 and 6.5,<sup>7</sup> so the proximal pH cases (5 and 7) without DOM had smaller absolute  $\zeta$ -potential values (+11.6 and −20.2 mV, respectively) than pH 3 (+29.2 mV) or pH 9 (−27.1 mV). Three-factor ANOVA analysis revealed that each of the three factors (pH, DOM type, and DOM concentration) affected the particle  $\zeta$ -potential significantly (Table S6). HA yielded surface  $\zeta$ -potentials significantly more negative than each of the other DOM isolates (Table S8), according to pairwise comparisons (Table S7). In order of negativity of  $\zeta$ -potentials across the different DOM concentrations and pH values, HA was most negative, followed by colloids, TPI, and then HPO. The presence of DOM, compared to the zero-DOM case, caused significant changes to  $\zeta$ -potential no matter the DOM concentration, but no differences were statistically different when comparing different levels of DOM content. This observation suggests that small amounts of DOM, relative to impacts on photocatalytic outcomes,<sup>4</sup> exerted meaningful changes to particle surface characteristics. Adjustments to solution pH were also found to significantly impact  $\zeta$ -potential across each pairing, except for pH 7 with pH 9 (Table S7).



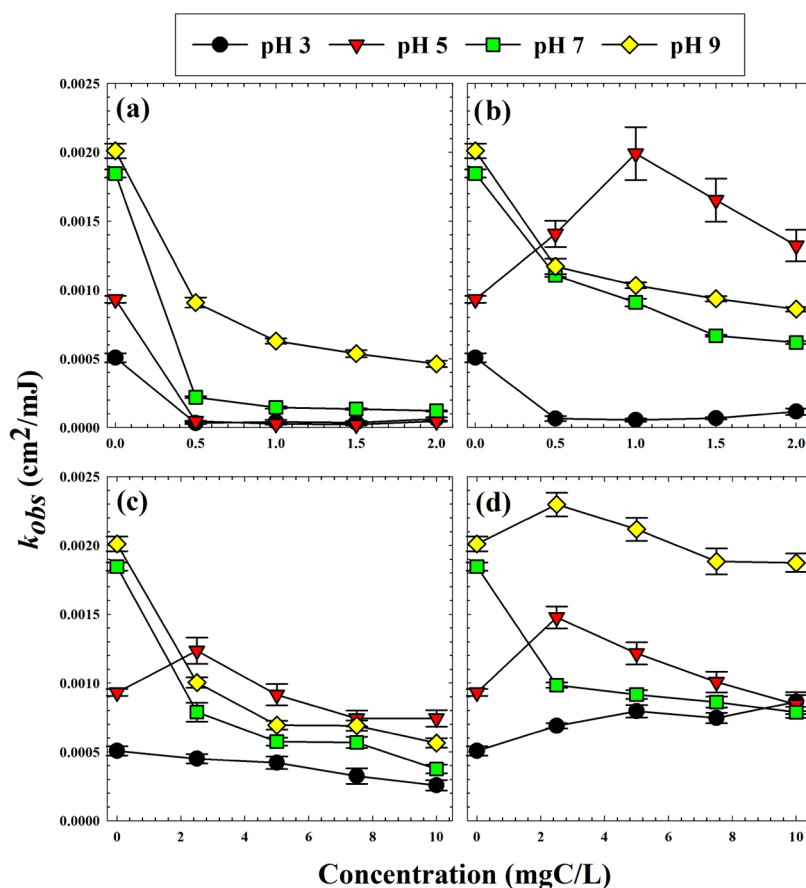


**Figure 2.**  $\text{TiO}_2$  particle  $\zeta$ -potential values as a function of DOM concentration at pH 3, 5, 7, and 9 for (a) HA, (b) colloids, (c) TPI, and (d) HPO. Error bars indicate standard error.

Several trends within individual experimental factors revealed notable trends in particle  $\zeta$ -potentials. The addition of HA at pH 5 changed the  $\zeta$ -potential from a low absolute value of  $+11.6 \pm 2.5$  mV to a higher, negative value of  $-27.8 \pm 3$  mV (average value calculated within the DOM concentration range, Table S5). The HA also decreased  $\text{TiO}_2$  particle size from  $276 \pm 31$  to  $211 \pm 26$  nm. With positively charged  $\text{TiO}_2$  at pH 5, the HA likely forms a water-rich, highly polar adlayer, as documented for a system of HA adsorbing onto positively charged, amine-terminated self-assembling monolayers by Armanious et al.;<sup>14</sup> this dynamic could stabilize the  $\text{TiO}_2$  by expressing negatively charged moieties out into solution, thereby explaining the reduced particle sizes. This HA-induced particle size reduction occurred at all other pH values tested, where the  $\zeta$ -potential became more negative. Notably, the changes in  $\zeta$ -potential provide good evidence for significant HA adsorption, while decreasing particle sizes show that a stabilization effect precludes conclusions on the basis of particle size only. This observation is in line with a study by Jayalath et al., which noted that HA-adsorbed  $\text{TiO}_2$  had a higher magnitude of  $\zeta$ -potential than bare  $\text{TiO}_2$ , regardless of HA concentration, resulting in greater particle stability in all cases.<sup>10</sup> Modification of surface properties by adsorption of negatively charged DOM also occurs at low pH values; most DOM molecules contain numerous carboxylic functional groups, which provide wide acid/base equilibria ranges at around pH 3–4.<sup>26–28</sup> In our case, the fractionated bulk supernatant samples yielded successively less negative particles

at pH 3—where  $\text{TiO}_2$  is normally positively charged—in order of  $\text{C} < \text{TPI} < \text{HPO}$ , and this trend matched well with reported carboxylic acid functionality in naturally derived DOM isolates, where an HPO fraction had significantly less carboxylic content than a TPI counterpart (colloidal matter were not studied in that report).<sup>28</sup> With fewer carboxylic groups, HPO is consequently less prone to electrostatic adsorption to the hydrophilic, positively charged  $\text{TiO}_2$ . At pH 5, the addition of colloids increased the magnitude of  $\zeta$ -potential to a lesser extent than HA, but particle sizes still decreased, showing that the colloidal fraction exerted a stabilization effect similar to HA at pH 5. Here again we note that colloids likely form a highly polar, stabilizing adlayer on the surface of  $\text{TiO}_2$ .<sup>14</sup> A decrease in particle sizes was also observed for the TPI case at pH 5, but no change was observed in the size in the case of HPO. The TPI fraction did not follow the expected  $\zeta$ -potential-size stabilization trend at basic pHs.  $\text{TiO}_2$  particle sizes increased from  $211 (\pm 24)$  at pH 5 to averages of  $281 (\pm 36)$  or  $282 (\pm 47)$  nm at pHs 7 or 9, respectively, despite higher  $\zeta$ -potential magnitudes. Overall, particle  $\zeta$ -potentials were generally inversely correlated to sizes, as expected. Notably, the different DOM samples had varying charge neutralization capacities; at pH 3, DOM imparted negative charge in order of  $\text{HA} > \text{colloids} > \text{TPI} > \text{HPO}$  in terms of magnitude of charge per concentration.

**Impacts of pH-Driven DOM– $\text{TiO}_2$  Interactions on Photocatalysis.** A series of *p*CBA photodegradation experiments were conducted in the presence or absence of DOM



**Figure 3.** Observed *p*CBA degradation rate constants (first order with respect to fluence) in the presence of 5 mg/L TiO<sub>2</sub> and various DOM concentrations at pH 3, 5, 7, and 9 for (a) HA, (b) colloids, (c) TPI, and (d) HPO. Error bars indicate standard error of linear trendlines with 95% confidence.

samples, as shown in Figure 3. Data used to generate these rate constants are shown in Figure S1. The lowest photodegradation rate constant (highest inhibition by DOM when present) occurred at pH 3 for nearly all cases. This observation was somewhat surprising since  $\bullet\text{OH}$  radical production is known to increase at low pHs via reactions between electron holes in the TiO<sub>2</sub> surface with protons in solution.<sup>29</sup> The reason for the reduced rate is likely related to limited adsorption of *p*CBA compound onto the TiO<sub>2</sub> surface. At pH 3, deprotonation occurs for only about 10% of *p*CBA molecules since its  $\text{pK}_a$  is 3.98.<sup>30</sup> Consequently, *p*CBA is less likely to adsorb onto the TiO<sub>2</sub> surface, since most of the molecules are not electrostatically attracted to the positively charged TiO<sub>2</sub>.<sup>31</sup> Because  $\bullet\text{OH}$  are generated at the TiO<sub>2</sub> surface and only a small portion of them diffuse away,<sup>11</sup> the adsorption of *p*CBA on the surface is a critical factor in its degradation rate. The low photodegradation constants at pH 3 could also be explained by competitive adsorption of Cl<sup>−</sup> (added via HCl) for adsorption sites on TiO<sub>2</sub> surface. Piscopo et al. noted that chloride competes with target compounds (*p*CBA in this case) for adsorption sites, resulting in reduced in photodegradation rates.<sup>32</sup> The decrease in the rate constants for *p*CBA degradation here indicates the adsorption of target contaminant onto the photocatalyst surface is a rate-controlling factor regardless of the concentration of generated  $\bullet\text{OH}$ . The addition of HA completely inhibited the photodegradation. Colloids also showed strong inhibition of photocatalysis at pH 3, and their nonlinear inhibitory profile of colloids indicated

strong affinity toward TiO<sub>2</sub>. HA exhibits a net negative charge in all ranges of pH used here,<sup>33</sup> and similar to the colloids, HA adsorbs strongly onto positively charged TiO<sub>2</sub> surfaces under acidic conditions. For both HA and the colloidal DOM, surface-phase inhibition was the dominant quenching mechanism at pH 3, whereas TPI presented a profile representative of bulk quenching.<sup>3</sup>

Given the different pathways of DOM interference, the inhibitory effect of TPI at pH 3 was milder compared to HA or colloids where the degradation rate in the presence of TPI was  $4.5 \times 10^{-4} \text{ cm}^2 \cdot \text{mJ}^{-1}$  at a concentration of 2.5 mgC/L, much higher than the same in the presence of HA or colloids at a concentration of 2.0 mgC/L ( $\sim 1.10^{-4} \text{ cm}^2 \cdot \text{mJ}^{-1}$ ). A comparatively low affinity between TiO<sub>2</sub> and TPI could be inferred by this observation, but the opposite was expected according to the analysis of the average hydrodynamic diameters of TiO<sub>2</sub>, which indicated increased aggregation under these conditions. Although the average hydrodynamic diameter of TiO<sub>2</sub> increased with the addition of TPI (390  $\pm$  47 nm) compared to HA (224  $\pm$  32 nm) or colloids (285  $\pm$  37 nm), the resultant inhibition profiles were strikingly different, suggesting that the aggregation of TiO<sub>2</sub> nanoparticles is not the key factor in the inhibition of photocatalysis. Destabilization of TiO<sub>2</sub> particles via DOM addition does not imply direct competition for photocatalytic surface sites.

HPO added to the TiO<sub>2</sub> system improved its photocatalytic performance at pH 3, counter to the expectation that nearly any DOM will compete with *p*CBA to scavenge  $\bullet\text{OH}$  in the

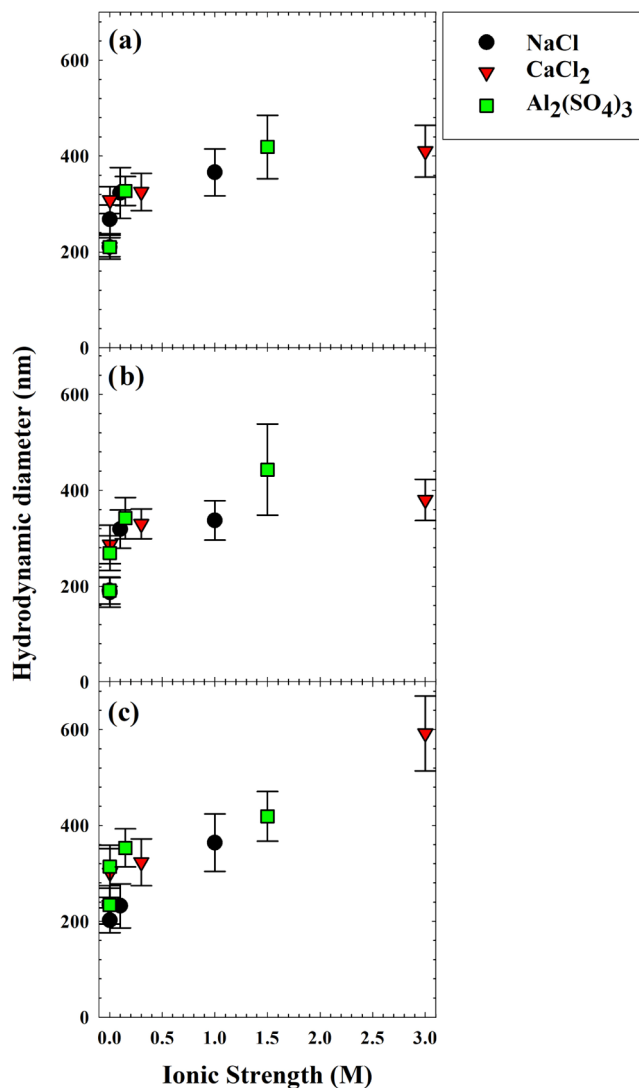
bulk phase if not on the  $\text{TiO}_2$  surface. This surprising enhancement to degradation kinetics suggests that HPO at pH 3 acts to accelerate *p*CBA oxidation in some way. The role of photochemically produced reactive intermediates is important in some DOM systems, especially where singlet oxygen is sensitized by DOM.<sup>34</sup> However, in our study, the kinetic enhancement effect is pH-dependent, which suggests that surface phenomena (e.g., favorable co-adsorption processes) are a more likely explanation. HPO also improved *p*CBA photodegradation kinetics at pH values 3, 5, and 9 but appeared to cause mild inhibition via surface and bulk competition at pH 7.<sup>3</sup> A kinetic enhancement was also observed for the colloidal DOM at pH 5, in stark contrast to its strong, surface competition inhibition at other pHs. With the exception of HA, this enhancement effect was also observed for all DOM samples at pH 5; the increases occurred at low DOM/ $\text{TiO}_2$  ratios, typically followed by up decreases with higher DOM concentrations. For example, the addition of up to 1 mgC/L colloids at pH 5 improved the photodegradation kinetics, at which point the trend reversed, suggesting that inhibitory mechanisms began to counterbalance the improvements. For TPI and HPO, increasing the concentration beyond 2.5 mgC/L caused inhibition. Drosos et al. noted a similar improvement at low ratios of DOM/ $\text{TiO}_2$  and concluded that the enhancement is caused by a reduction in diffusion limitations when DOM attracts targeted contaminants to the surface of  $\text{TiO}_2$ .<sup>35</sup> They also showed that at high DOM: $\text{TiO}_2$  ratios, DOM blocks the target contaminants from reaching the surface of  $\text{TiO}_2$ .<sup>35</sup>

At pH 7, all DOM types inhibited *p*CBA photodegradation, with HA exerting the strongest effect. The inhibition by HA and colloids was likely caused by strong adsorption interactions, given the exponential decrease of rates with increasing DOM, while HPO and TPI appeared to inhibit partly via surface and partly by bulk quenching. Photodegradation experiments at pH 9 showed largely the same results, with the notable exception of HPO, which demonstrated an enhancement effect similar to HPO at pH 3 or pH 5. At neutral and basic conditions,  $\text{TiO}_2$  and *p*CBA both have negative charges and experience some electrostatic repulsion, and poor *p*CBA- $\text{TiO}_2$  adsorption at high pHs was reported by Jayalath and co-workers in 2018, where they showed a decreasing trend in the adsorption of HA with the increase in pH.<sup>10</sup> On the other hand, large DOM molecules with varied functional groups and charge distributions are more likely to sorb partly, if not wholly, onto  $\text{TiO}_2$ . Indeed, partial adsorption of DOM molecules is precisely the mechanism we suppose is responsible for the photodegradation enhancement observed in some cases, and the HPO fraction is the least likely to have fully favorable interactions with hydrophilic  $\text{TiO}_2$ . Note that  $\text{TiO}_2$  is least charged, therefore most hydrophobic, at pH 7, which is where HPO competes most strongly with *p*CBA for adsorption sites.

**Impacts of IS-Driven DOM- $\text{TiO}_2$  Interactions on Photocatalysis.** The effect of ionic strength on the aggregation of  $\text{TiO}_2$  was investigated by adding different salts to solutions in the presence or absence of different fractions, and the pH of solutions was adjusted to 6.4. The TPI and HPO fractions were selected for comparison because linear DOM-inhibitory profiles were observed for these fractions under dilute conditions in our previous work,<sup>4</sup> implying that the inhibition was primarily bulk-phase competition.<sup>3</sup> These fractions, then, provide an excellent test case to determine

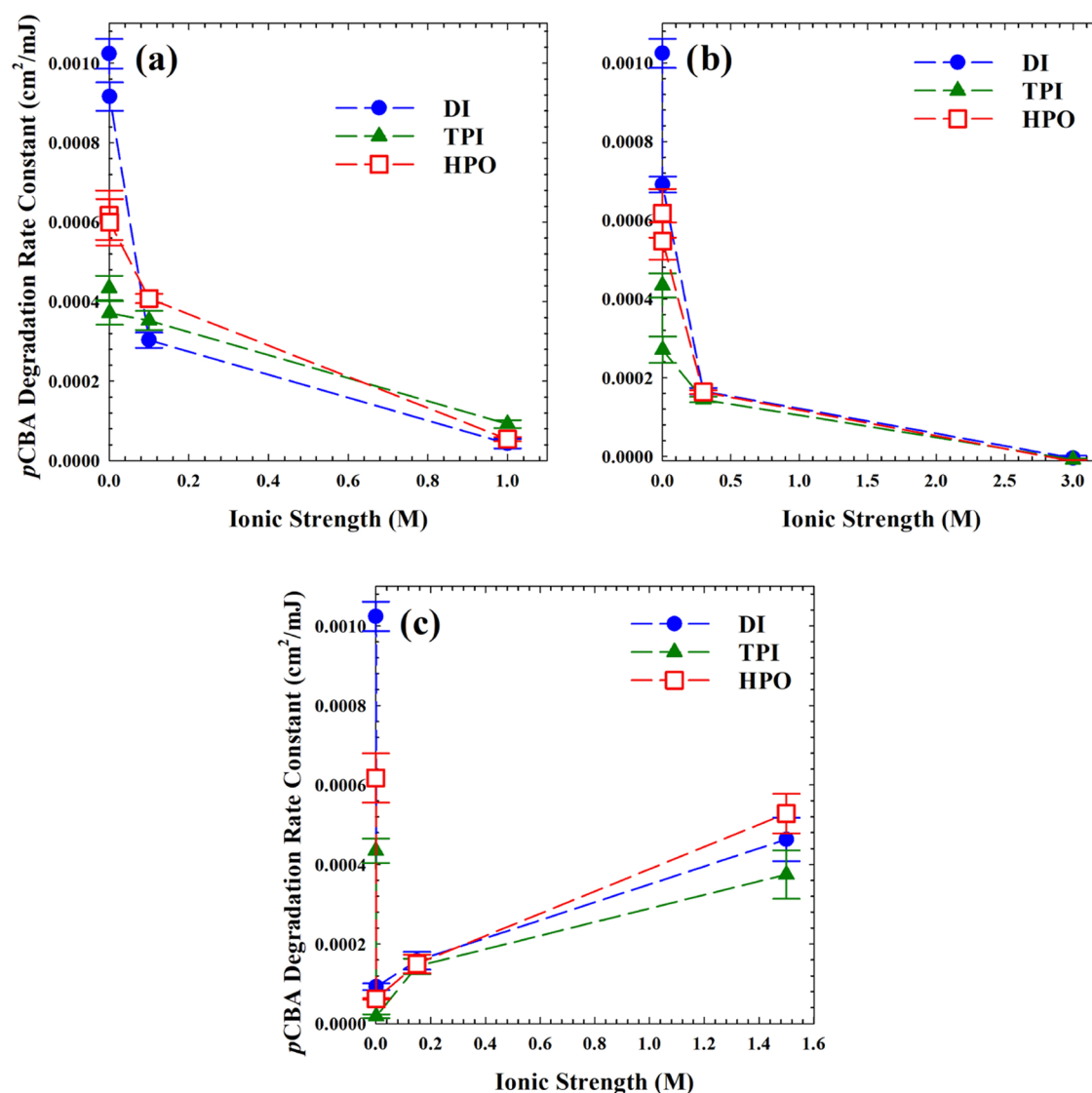
whether particle destabilization will induce surface-phase inhibition or if the presence of multivalent ions will otherwise affect the system.

DLS measurements were collected for  $\text{TiO}_2$  particles under a variety of cosolute conditions; these data are plotted in Figure 4 and shown in Table S9. Corresponding ANOVA results are



**Figure 4.** Hydrodynamic diameters of  $\text{TiO}_2$  particles as a function of IS (pH 6.4) added via NaCl,  $\text{CaCl}_2$ , or  $\text{Al}_2(\text{SO}_4)_3$  for cases (a) without DOM, (b) with 10 mgC/L TPI, and (c) with 10 mgC/L HPO. Error bars indicate standard error.

shown in Tables S10–S12. With a fixed pH value, surface charge characteristics should not change meaningfully in direction or magnitude with changing IS; the electrical double-layer distance should compress with increasing IS,<sup>36,37</sup> but the PALS platform was not suitable to discern these changes in terms of  $\zeta$ -potential. The three-factor ANOVA found that only IS impacted particle diameters significantly across the conditions tested (Table S10); the  $\text{TiO}_2$  particles aggregated more with increasing ionic strength (Table S12). The particle diameters in these solutions were largely unaffected upon further addition of HPO or TPI, with two exceptions. When  $\text{CaCl}_2$  was added to yield an ionic strength of 3 M, larger aggregates formed with HPO compared



**Figure 5.** *p*CBA degradation rate constants in the presence of 5 mg/L TiO<sub>2</sub>, TPI, and HPO in various ionic strengths of (a) NaCl, (b) CaCl<sub>2</sub>, and (c) Al<sub>2</sub>(SO<sub>4</sub>)<sub>3</sub>. Error bars indicate standard error of linear trendlines with 95% confidence.

to the TPI and no-DOM cases. Likewise, when Al<sub>2</sub>(SO<sub>4</sub>)<sub>3</sub> was added to an ionic strength of 1.5 M, a small particle size increase was observed with HPO. These observations suggest first that the TPI and HPO fractions generally do not stabilize or destabilize TiO<sub>2</sub> particles with increasing ionic strength and second that the HPO molecules may be affected differently by multivalent cations, enhancing TiO<sub>2</sub> aggregation in the presence of Ca<sup>2+</sup> with Cl<sup>−</sup> (3 M ionic strength) or of Al<sup>3+</sup> with SO<sub>4</sub><sup>2−</sup> (1.5 M ionic strength). Note that this amount of aluminum sulfate is expected to induce sweep coagulation with the formation of Al(OH)<sub>3</sub> solids, kept in suspension via stirring. The extreme condition here provides a case of maximum particle destabilization. While 3 M ionic strength may not be relevant in real systems since typical seawater has an ionic strength of about 0.7 M,<sup>38</sup> the results suggest that conformational changes in DOM do not necessitate changes in TiO<sub>2</sub> agglomeration. For instance, millimolar concentrations of CaCl<sub>2</sub> were found to cause significant changes in size and conformational to humic substances by Baalousha and co-workers.<sup>18</sup>

**Effects of IS and DOM on Photocatalysis.** Photodegradation experiments were also performed across ranges of IS and DOM concentrations to determine if destabilization of DOM–TiO<sub>2</sub> systems induces inhibition via surface competition. Results from these experiments are arrayed in Figure 5. Data used to generate these rate constants are shown in Figure S2. The increase in IS decreased the photodegradation rates in all cases, even with only a small increase in IS (0.01–0.03 M). Generally, decreases in photoactivity may be expected with increasing IS due to compression of the electrical double layer and consequent agglomeration of TiO<sub>2</sub> with itself or DOM. However, our scrutiny of changes in agglomeration due to pH demonstrated that particle size may not be a reliable predictor of photoactivity. Although reaction rates do depend on the physical properties of TiO<sub>2</sub> nanoparticles, including size,<sup>39</sup> Lin et al. noted that there are disagreements in this area; some researchers claim that the efficiency of photocatalytic processes did not increase monotonically with a decrease in particle size.<sup>40</sup> According to Dionysiou et al., increasing IS decreased *p*CBA adsorption onto TiO<sub>2</sub> surfaces, reducing the photodegradation rate in their system.<sup>41</sup> Reductions in *p*CBA



adsorption onto  $\text{TiO}_2$  is a more likely explanation of the decrease in *p*CBA degradation rates observed here. Researchers have shown that chloride in particular competes with some aromatic compounds for surface adsorption sites.<sup>32</sup>

The addition of different salts in small amounts (0.01–0.03 M as IS) caused a range of inhibition according to the following order:  $\text{Al}_2(\text{SO}_4)_3 > \text{CaCl}_2 > \text{NaCl}$ . Recent work by Raza et al. (2016) demonstrated that electrolyte type plays an important role in the morphology of  $\text{TiO}_2$ –DOM aggregation:  $\text{Cl}^-$  induced tight aggregates, while  $\text{NO}_3^-$  promoted loose aggregation with larger particle sizes.<sup>42</sup> The observed ordering of inhibition here suggests that morphological changes associated with the valency of ions in solution may explain the differential photocatalytic performances. Moreover, Wang et al. also confirmed that  $\text{Ca}^{2+}$  and other divalent cations caused more inhibition than  $\text{Na}^+$ .<sup>43</sup> Increasing the IS lowered the observed reaction rate constants in both NaCl and  $\text{CaCl}_2$  cases, but  $\text{Al}_2(\text{SO}_4)_3$  accelerated the reaction after the initial decline. This effect was also observed by Wang et al.; they postulated that the mechanism was a hydrolytic effect by  $\text{Al}^{3+}$ , where poly-aluminum hydroxides form by reactions of  $\text{Al}^{3+}$  with  $\text{H}_2\text{O}$ , releasing some  $\text{H}^+$ .<sup>43</sup> Notably, this effect did not occur in a low concentration of  $\text{Al}_2(\text{SO}_4)_3$  as the results showed almost complete inhibition by increasing the IS with  $\text{Al}_2(\text{SO}_4)_3$  to 0.015 M.

The effects of ionic species on DOM– $\text{TiO}_2$  interactions were examined using TPI and HPO. Since both TPI and HPO exhibit bulk phase inhibition in dilute solutions,<sup>4</sup> ion-induced adsorption of the DOM onto  $\text{TiO}_2$  should be apparent. The decreases in photodegradation rate constants caused by increasing IS were smaller for TPI and HPO compared to DI. The addition of 0.015 M  $\text{Al}_2(\text{SO}_4)_3$  caused a decrease of  $0.0009 \text{ cm}^{-2}\cdot\text{mJ}$  for the rate constant in DI, which was about twice the change observed with TPI ( $0.0004 \text{ cm}^{-2}\cdot\text{mJ}$ ) or HPO ( $0.0005 \text{ cm}^{-2}\cdot\text{mJ}$ ) in solution. This observation was surprising because DOM is a major source of quenching for  $\cdot\text{OH}$  and adsorption of the DOM onto  $\text{TiO}_2$  was expected to increase with IS due to particle destabilization. The resulting rate constants, however, were clustered near a floor value of  $\sim 0.0001 \text{ cm}^{-2}\cdot\text{mJ}$  with minimal photoactivity retained. In the NaCl and  $\text{CaCl}_2$  cases, the added salts appeared to cause the same suppression of photoactivity at all IS values, regardless of the presence of TPI or HPO, since the DI case tracked closely with the TPI and HPO trends. Manipulation of IS did not meaningfully alter the surface competition dynamics (or lack thereof) of TPI or HPO onto  $\text{TiO}_2$  particles.

## CONCLUSIONS

In the search for applied, practical photocatalytic water treatment, recent work revealed an important parameter for designing effective systems; adsorption interactions of target and interfering molecules onto catalyst surfaces can be used as a tool to optimize photocatalysis in complex waters.<sup>3,4</sup> Here, the DOM– $\text{TiO}_2$  surface adsorption mechanism was scrutinized by inducing adsorption and particle aggregation via changes in pH and IS across conditions expected to completely destabilize the DOM– $\text{TiO}_2$  colloidal particles. Resulting photocatalytic performances highlighted several critical points for controlling the unwanted DOM quenching reactions in photocatalytic systems:

- Adjustments to pH induced aggregation near  $\text{TiO}_2$ 's isoelectric point, but observed increases in aggregate size

did not correlate to increased inhibition, contrary to prior reports.<sup>44,45</sup>

- Particle  $\zeta$ -potential predicted aggregate sizes but did not directly correlate with inhibition. The DOM types which imparted more negative surface charge at pH 3 were the stronger inhibitors, in order of HA, colloids, TPI, and HPO.
- Increases to IS reduced photocatalytic performance in most cases, and aggregate sizes increased with increasing IS. But aggregate sizes were not predictive of inhibition for the ionic conditions tested.
- The effect of IS on photocatalytic performance was unchanged by the presence of TPI or HPO regardless of salt type used.

The simple fact that DOM– $\text{TiO}_2$  agglomerate size cannot predict photocatalytic inhibition has profound implications on prospective photocatalytic applications and quenching mitigation strategies. Neither the compression of the electrical double layer nor DOM conformational changes by multivalent cations were important in the DOM– $\text{TiO}_2$  system. Site-specific interactions between DOM molecules and the photocatalyst surface are fundamental to the inhibitory process, as demonstrated by the stark differences between inhibition profiles of the DOM isolates studied here at different pHs. In fact, DOM can promote photodegradation of target molecules at rates beyond pure water cases under certain circumstances; this phenomenon appears to be directly related to the dynamics of molecular interactions between the target (*p*CBA here), DOM molecules, and the  $\text{TiO}_2$  surface. Further investigations on the fundamental interactions of moieties of target and inhibitory compounds will be fruitful for efforts aimed at predicting and preventing inhibition of photocatalysts by non-target organics.

## ASSOCIATED CONTENT

### Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/acs.langmuir.2c03487>.

Tables of DLS and PALS data and three-way ANOVA statistics for  $\text{TiO}_2$  particle sizes and  $\zeta$ -potential with variable solution conditions, and figures of *p*CBA photodegradation profiles according to pH, DOM type, IS, and ion species (PDF)

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## Notes

The authors declare no competing financial interest.

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