Supplementary Materials: Removal of Algae, and Taste and Odor Compounds by Combination of Plant-Mineral Composite (PMC) Coagulant with UV-AOPs: Laboratory and Pilot Scale Studies

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Section S1. Measurement of OH radical scavenging factor

Application of the $R_{OH,UV}$ concept for measuring the OH radical scavenging rate has been recognized as a more reliable method for •OH water demand measurement than other methods based on theoretical calculations [1]. The degradation of a well-known chemical probe compound via direct photolysis and •OH reaction parameters were used to determine the $R_{OH,UV}$ value (Eq. S1). In this study, para-chlorobenzoic acid (*p*CBA) was used as a probe compound for measuring the scavenging factor and this method was applied in this work to determine the scavenging factor.

$$R_{OH,UV} = \frac{\int_0^t [\cdot OH] dt}{E_0 \times t} = \frac{k_T^{\prime app} - k_d^{\prime app}}{k_{OH,pCBA}^{app}}$$
(S1)

where $k_d^{'app}$ is the fluence-based apparent rate constant for direct UV photolysis, $k_T^{'app}$ is the fluencebased apparent rate constant when H₂O₂ is added, and $k_{OH,pCBA}^{app}$ is the apparent rate constant of *p*CBA solution decolorization (5 × 10⁹ M⁻¹ s⁻¹) [1].

The •OH water background demand can be calculated using the measured ROH, UV values with Eq. (S2).

$$\sum k_{s,OH}[S]_i = \left(k_{H_2O_2,OH} \times \frac{m}{b}\right) - k_{OH,pCBA}^{app}[\text{pCBA}]$$
(S2)

where $\sum k_{s,OH}[S]_i$ is the •OH water demand (s⁻¹);, *k*_{H2O2,OH} is the second-order rate constant for the •OH reaction with H₂O₂ (2.7 × 10⁷ M⁻¹ s⁻¹) [2, 3]; m and b are the slope and the intercept of the plot of 1/*R*OH,UV vs. 1/[H₂O₂]₀ (Fig. S1), respectively; and [*p*CBA] is the initial concentration of *p*CBA.



Figure S1. Determination of scavenging factor for the water sample before and after PMC treatment used in collimated beam work: *R*OH,UV variation with the initial H₂O₂ concentration;

References

- 1. Rosenfeldt, E.J.; Linden K.G. The R-OH,R-UV concept to characterize and the model UV/H₂O₂ process in natural waters. Environ. Sci. Technol. **2007**, *41*, 2548–2553.
- 2. Buxton, G.V.; Greenstock, C.L.; Helman, W.P.; Ross, A.B. Critical review of rate constants for reactions of hydrated electrons, hydrogen atoms and hydroxyl radicals (OH/O⁻ in aqueous solution. J. Physic. Chem. Reference Data, **1988**, *17*, 513–886.
- 3. Kwon, M.; Kim, S.; Yoon, Y.; Jung, Y.; Hwang, T.M.; Kang, J.W. Prediction of the removal efficiency of pharmaceuticals by a rapid spectrophotometric method using Rhodamine B in the UV/H₂O₂ process. Chem. Eng. J. **2014**, *236*, 438–447.