

Article

Research about Organic Matter Removal and Biofilms Development of Pilot-Scale UV/H₂O₂-BAC Process

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Abstract: As a green advanced process for drinking water treatment, the UV/hydrogen peroxide (UV/H₂O₂) process has been gradually applied in China. To study the effect and mechanism of organic matter removal and the development of microbial communities in the UV/H₂O₂-biological activated carbon (UV/H₂O₂-BAC) process, a pilot-scale UV/H₂O₂-BAC system was built and operated over one year. Low water temperature affects the UV/H₂O₂ process efficiency, the biofilms in the BAC system were mature and stable after 240 days, and the contribution rate of BAC adsorption to dissolved organic carbon (DOC) removal was approximately 14.2% after one year of operation. The liquid chromatography-organic carbon detection (LC-OCD) analysis shows that UV/H₂O₂ process can increase the amounts of Low Molecular Weight (LMW) neutrals, and the specific UV absorbance (SUVA₂₅₄) value is not suitable for predicting Trihalomethanes (THMs) precursor contents in water after UV/H₂O₂ treatment. High-throughput sequencing results prove that microbial species in the middle section are the most abundant compared to those in the influent and effluent sections, hydrogen peroxide has lower inhibition on the development of microbial community than ozone and the low concentration of hydrogen peroxide (<0.25 mg/L) promotes the development of the microbial communities, hydrogen peroxide can reduce *Proteobacteria* abundance by inhibiting the growth of anaerobes. *Acidobacteria* may have a certain contribution to the degradation of soil organic matter (SOM), and the effluent section of BAC with low DOC concentration cannot form the dominant species of *Rhodobacter*.

Keywords: drinking water treatment; granular activated carbon; UV-AOPs; biofilm microbial community



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1. Introduction

With the development of the economy and the improvement of productivity, developing countries are facing more serious problems of surface water pollution, such as eutrophication of water body and toxic organic matter pollution, and source water is gradually micro-polluted, which also increases the difficulty of ensuring the safety of drinking water. Relevant investigation and research showed that the average antibiotic concentration in the middle and lower reaches of the Yangtze River Basin was about 92.95–156 ng/L [1], and the highest concentration of ciprofloxacin in the Yamuna River in India even reached 1440 ng/L [2]. During the eutrophication period of surface water bodies, the content of 2-MIB, a typical odor compound in the source water of eastern China, reached 2000 ng/L [3,4].

Conventional treatment processes are inadequate at controlling the levels of organic pollutants in the source water; hence, water quality safety has become an urgent problem to solve. At present, the O₃-BAC process is the mainstream advanced treatment process in China [5], the O₃-BAC process can effectively remove organic pollutants, but the removal effect of high-concentration odor substances of this process water is not good since the characteristics of selective oxidation of ozone, and it easily causes bromate problems [6]. Therefore, it is necessary to find a more efficient and safe alternative technology.

The UV/H₂O₂ process can effectively remove odor substances in drinking water [7]. Our previous study showed that the UV/H₂O₂-biological activated carbon (UV/H₂O₂-BAC) combined process had higher organic matter removal efficiency and effluent safety than the O₃-BAC process [8]. Granular activated carbon (GAC) with large specific surface areas and suitable pore structures demonstrates high adsorption capacities, which can effectively remove the dissolved organic compounds, including biodegradable organic compounds, halogenated hydrocarbons, odor compounds and other pollutants [8,9]. With the operation of the GAC system, biofilms are formed on the surface of activated carbon, and GAC gradually changes into BAC, and biodegradation becomes the main reason for organic matter removal [10,11]. Identifying the dominant bacteria population composition can increase the understanding of activated carbon filter operations, which will improve the BAC filter designs and maximize the BAC filter application realization. At present, the effect of the influent water quality on the development of microbial populations in BAC filter systems has been reported, such as that DOC and ammonia nitrogen levels could affect the bacterial diversity and community composition in BAC filters [12], and High residual ozone decreased the attached bacterial density and removal of DOC [13]. However, the effect of the effluent water with H₂O₂ derived from the UV/H₂O₂ process on the BAC filter systems requires additional investigation, and most studies mainly focused on the effect of the water quality derived from the ozone-oxidized effluent on the population structure of the BAC filter [14–16], while the effect of residual H₂O₂ from the UV/H₂O₂ process on the BAC filter microorganism was neglected. Furthermore, most experimental studies were limited to laboratory-scale and short-term research, and few studies regarding the removal efficiency of organic matter and microbial communities with different carbon layer depths in the pilot-scale UV/H₂O₂-BAC combined process have been published.

To solve these problems, a pilot-scale UV/H₂O₂-BAC process was built. The organic matter removal efficiency was studied, and liquid chromatography-organic carbon detection (LC-OCD) was used to analyze the changes of organic matters during the UV/H₂O₂-BAC process. Meanwhile, the development of biofilms in the activated carbon was studied. The structure and abundance of microbial communities at different carbon depths were compared through high-throughput sequencing. The effect of different doses of H₂O₂ on the development of microbial communities was obtained, which contributes to the better application of the UV/H₂O₂-BAC process in actual water treatment.

2. Materials and Methods

2.1. Experimental Setup and Operating Conditions

The pilot plants were located in the Quehua Water Plant, which takes water directly from the Yellow river reservoir. The process flowrate of the UV/H₂O₂ process was 5 m³/h. The UV/H₂O₂ system was provided by Trojan, and the UV system consists of four 500 W low-voltage UV-C lamps, and the theoretical UV dose is 350 mJ/cm². The process flow diagram is shown in Figure 1. The operation mode of the activated carbon column was upward flow, the diameter of the activated carbon column was 20 cm, the process flowrate was 80 L/h, the empty bed contact time (EBCT) was 10.6 min, the filling height was 180 cm, and the height of the supporting layer was 20 cm. The used carbon was coconut shell activated carbon, the parameters are shown in Table S1, and the influent water quality indices of BAC are shown in Table 1.

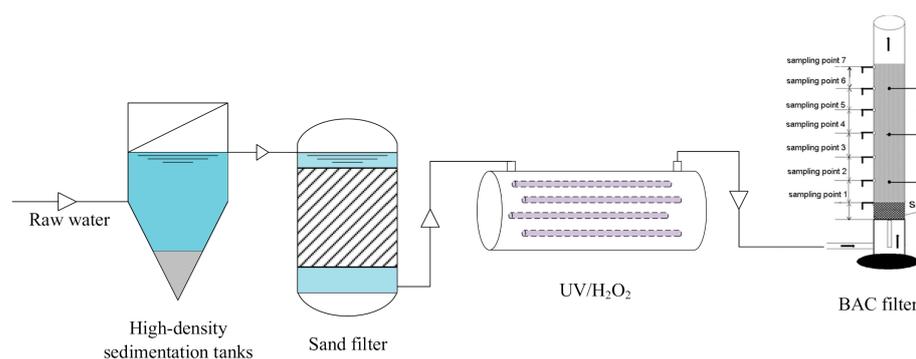


Figure 1. Process flow and structural of UV/H₂O₂-biological activated carbon (UV/H₂O₂-BAC).

Table 1. Characteristics of BAC influent.

Temp (°C)	Turbidity (NTU)	pH	UV ₂₅₄ (cm ⁻¹)	DOC (mg/L)	NH ₄ ⁺ -N (mg/L)	H ₂ O ₂ (mg/L)	DO (mg/L)	Br ⁻ (ug/L)	I ⁻ (ug/L)
10–22	0.80–2.15	8.0–8.3	0.21–0.28	2.69–3.35	0.41–0.53	5–6.5	4.0–5.0	243–262	14–17

Note: DO is the abbreviation of dissolved oxygen.

2.2. Experimental Analysis Methods

2.2.1. Conventional Index

The conventional indices for detection are the DOC, UV₂₅₄, H₂O₂ content, specific surface area (SBET), molecular weight distribution and phospholipid biomass. The detection methods are listed in Table 2.

Table 2. Conventional indexes detecting methods or instruments.

Number	Detection Index	Determination Method
1	DOC	Shimazu TOC analyzer
2	DO	Thermo Orion 3star DO analyzer
3	UV ₂₅₄	Ultraviolet spectrophotometry
4	H ₂ O ₂ content	Spectrophotometry
5	SBET	AUTOSORB 6b analyzer
6	Molecular weight distribution	Ultrafiltration-TOC analyzer
7	Phospholipid biomass	Phosphatide method [17]

2.2.2. LC-OCD Scanning

The principle of LC-OCD is to separate organics according to molecular weight using size exclusion chromatography and subsequently analyze them by OCD, ultraviolet detector (UVD) and OND detector. The organics in water can be divided into biopolymers, humic acid, building blocks, acid and LMW humics, LMW neutral [18,19].

The detection conditions of LC-OCD were as follows: The instrument model is LC-OCD-Model 9. The chromatographic column is a polymethacrylate-based weak cation exchange column, the sample volume was 1000 uL, the total running time of each sample was 70 min, and the experimental results were obtained by peak integration with specific software.

2.2.3. High-Throughput Sequencing and Parameter Analysis

The carbon samples were taken at depths of 30 cm (GAC 1), 90 cm (GAC 2), and 150 cm (GAC 3) from the water influent end of the activated carbon layer, each position was sampled three times in parallel to reduce the errors. According to the Illumina Miseq high-throughput sequencing requirements, bidirectional sequencing was performed to design the target region with a “5” Miseq linker. Using a two-step PCR amplification method, first, a specific primer (hereafter described as an inner primer) was used to amplify the

target fragment, which was subjected to gel recovery with the recovered product used as a template for secondary PCR amplification. The specific detection steps and the calculation of Chao 1, Shannon and Simpson indexes can refer to the research of Zhang [20].

3. Results and Discussion

3.1. Organic Matter Removal by UV/H₂O₂-BAC Process

Figure 2 shows the variations and removal rates of the DOC concentration at the influent and effluent areas of the UV/H₂O₂-BAC process. After the UV/H₂O₂ process oxidation, the average DOC value was reduced from 4 mg/L (influent) to 3 mg/L (effluent), and the DOC value of the activated carbon filter effluent water was maintained at an average low level of 1.2 mg/L. By analyzing the change trend of the DOC removal rate in the UV/H₂O₂ process, we observe that the DOC removal rate in winter (December–February) has a significant downward trend. This trend was interesting because the temperature coefficient of most photochemical reactions was close to 1, the removal efficiency of organic matter by the UV/H₂O₂ process generally does not change with the change in temperature [21,22], and DOC is not a gas-phase organic matter and does not involve heterogeneous mass transfer process. We speculate that water temperature affects the UV output efficiency of mercury lamps, which reduces photon production [23]. The result of the H₂O₂ ratios also supports this speculation (Figure S1). According to Einstein's law of photochemical equivalence, since the absorption of photons by molecules is a single-photon process, the reduction of photon production will reduce the generation of hydroxyl radicals, so the removal rate of DOC will be reduced.

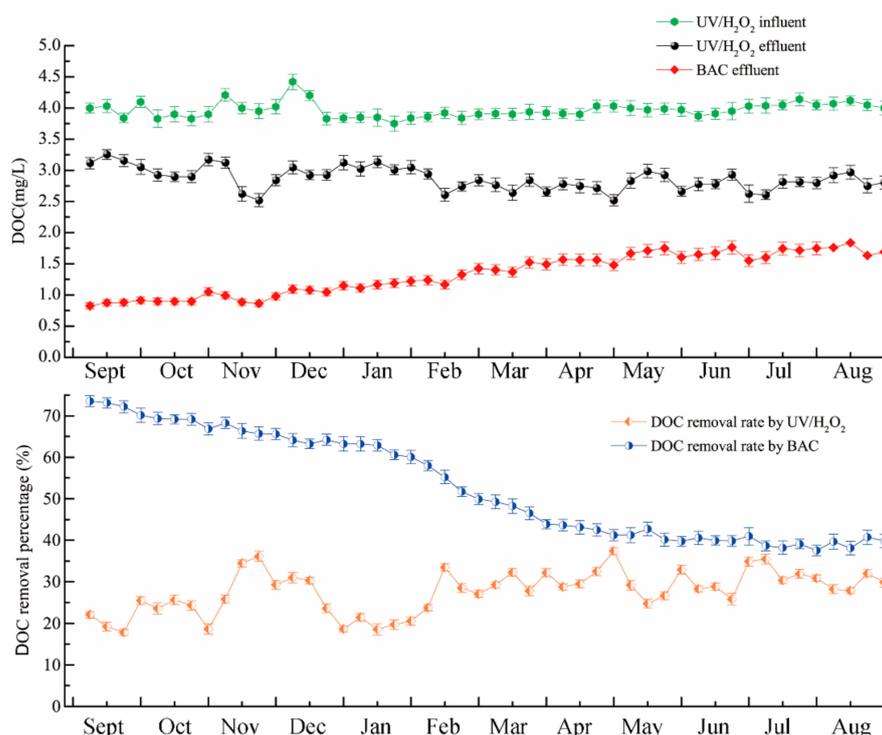


Figure 2. Dissolved organic carbon (DOC) removal of UV/H₂O₂-BAC process.

In the initial stage of the process operation, the removal of DOC by the activated carbon filter maintained a high efficiency with an average removal rate of approximately 70%. The high DOC removal rate was directly attributed to the adsorption of the activated carbons. Afterwards, when the adsorption capacity of activated carbon tends to be saturated, the biofilms on the surface of activated carbon gradually forms and BAC gradually transits to GAC; the DOC value removal rate gradually decreased and finally maintained at approximately 38.1%. Compared with pilot-scale O₃-BAC, UV/H₂O₂-BAC has a higher

organic matter removal efficiency. Ross et al. showed that the DOC removal rate by BAC in the stable operation stage of the O₃-BAC process was approximately 24% [24]. Our research group also conducted a pilot test on the O₃-BAC process using the same pretreatment equipment, and the results showed that the removal rate of BAC to DOC in the stable stage was approximately 34 ± 2% [25].

Figure 3 shows the specific surface area change of the activated carbons as a function of the carbon depths. The specific surface area can reflect the adsorption capacity of the activated carbon filter and subsequently explain the high DOC removal efficiency of the activated carbon filter from the physical viewpoint. S_{BET} decreases from 974 m²/g initially to 704 m²/g (GAC 1), 717 m²/g (GAC 2), and 730 m²/g (GAC 3) during operation, respectively. The decrease in SBET value at the influent section (GAC 1) occurred earlier than that at the middle section (GAC 2) and effluent section (GAC 3). After 240 days of operation, the SBET value tended to stabilize, which was similar to the change trend of the BAC removal rate of DOC in Figure 2, indicating that the high efficiency of the BAC removal of DOC in the early stage of system operation may be attributed to adsorption. When the adsorption capacity of activated carbon tends to be saturated, the removal rate of DOC also tends to be stable. At this stage, biodegradation was the main reason for the removal of organic matter by activated carbon columns. Liang's mathematical model showed that the efficiency of biodegradation to DOC removal in BAC at this stage accounted for approximately 88% [26]. Our previous study showed that the average DOC removal rate of the same activated carbon column was approximately 38.1% and 32.7% after continuous operation for one year and five years, respectively [8]. Since the biofilms on the activated carbons had been stabled after one year of operation, it was assumed that the decrease in DOC removal rate was due to the decrease in adsorption capacity, and we assumed that the adsorption capacity of BAC has reached the limit after 5 years of operation; then, the calculated contribution rate of adsorption to the DOC removal in the activated carbon column after one year of operation was approximately $\eta = \frac{(38.1\% - 32.7\%)}{38.1\%} = 14.2\%$, and the biodegradation efficiency was approximately 85.8%, which was in agreement with the conclusion of Liang et al.

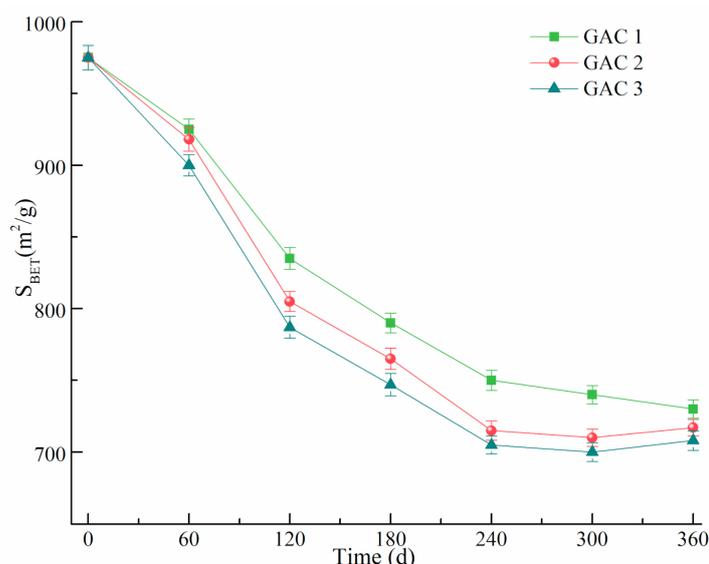


Figure 3. Specific surface area of granular activated carbon (GAC) in different carbon layers.

3.2. Organics Transform Analysis of UV/H₂O₂-BAC Process

3.2.1. Change of Molecular Weights of Organic Matters

Figure 4 shows the change in molecular weight distribution in the UV/H₂O₂-BAC process. Compared with the BAC process, the UV/H₂O₂ process had lower DOC value removal efficiency, which was determined by the characteristics of the UV/H₂O₂ process.

Previous studies had also shown that the simple advanced oxidation process was difficult to have a good mineralization effect on organics [27,28], but the UV/H₂O₂ process can decompose macromolecular organics in raw water into micromolecular organics, which promotes the subsequent removal of organics by the BAC process [8,29]. In Figure 4, the removal rate of organics with a molecular weight greater than 30 kDa by the UV/H₂O₂ process was approximately 50%, and the DOC values of small-molecule organics did not decrease significantly, which indicated that hydroxyl radical can oxidize macromolecular organics into small molecules or double-bond organics. The activated carbon filter had a better removal effect on small molecular organics (<3 kDa) with a removal rate of over 70%.

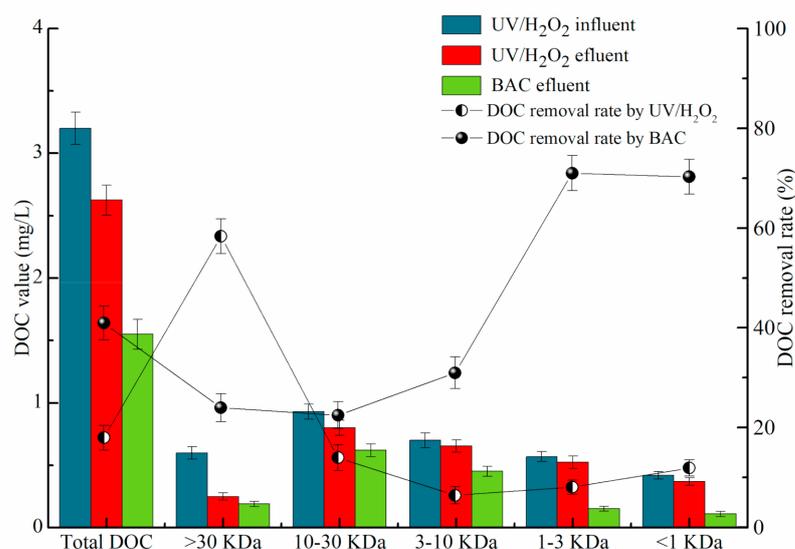


Figure 4. Change of molecular weights distribution of UV/H₂O₂-BAC process.

3.2.2. LC-OCD-OND Fractionation Analysis

LC-OCD-OND was used to scan the influent and effluent water of the process, and the results are shown in Figure 5. The peaks A, B, C, D and E of the black curves represent biopolymers, humic, building blocks, LMW acids and LMW neutrals respectively. The molecular weights decrease in turn, and the blue curves were the UV₂₅₄ response value of each component. It can be seen from Figure 5 that the peak areas of biopolymers and humic decreased significantly after the UV/H₂O₂ process, the peak areas of building blocks, LMW acids and LMW neutrals with smaller molecular weights increased, which also confirmed the fragmentation effect of advanced oxidation on macromolecular components. The response values of UV₂₅₄ decreased after advanced oxidation, which indicated that advanced oxidation had good degradation effect on aromatic organic compounds; the results of quantitative analysis (Table 3) also showed that SUVA₂₅₄ value decreased from 2.19 to 1.02 after UV/H₂O₂ process. Although Marais proved that there is a positive correlation between the number of Trihalomethanes (THMs) precursors and SUVA₂₅₄ values in natural water [30], our detection results showed that the content of THMs and haloacetic acid (HAA) precursor increases after UV/H₂O₂ process (Table S2). This indicates that the SUVA₂₅₄ value is not suitable for characterizing DBP precursors in water after advanced oxidation treatment; another possible reason was that the SUVA₂₅₄ value of natural water involved in this study is low, and some studies have shown that there is no good linear relationship between SUVA₂₅₄ value and the content of THMs precursors in low SUVA₂₅₄ water [30,31].

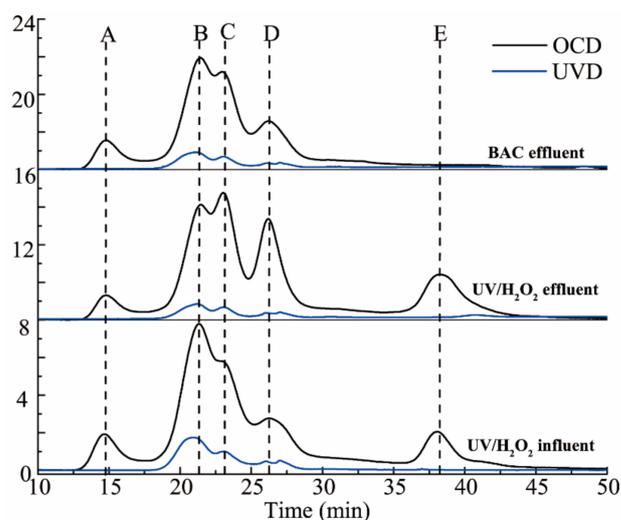


Figure 5. Variation tendency of organic carbon detection (OCD) and UVD signals.

Table 3. Results of liquid chromatography-organic carbon detection (LC-OCD) fractionation.

	Humic		SUVA ₂₅₄ (L/mg·m)	SOM(mg/L)	Biopolymers
	DON(mg/L)	Mol.-Weight(Mn)			%
UV/H ₂ O ₂ influent	0.1	445	2.19	0.328	100%
UV/H ₂ O ₂ effluent	0.069	355	1.02	0.259	90%
BAC effluent	0.072	403	1.53	<1	100%

The quantitative analysis of LC-OCD scanning results is shown in Tables 3 and 4. After UV/H₂O₂ oxidation, the biopolymer content reduced 33.9%, and the proportion of protein in biopolymers reduced from 100% to 90%, which also confirms the degradation effect of hydroxyl radicals on protein organics with larger molecular weight [28]. After advanced oxidation, the content of LMW neutrals increased by 16%, but decreased by 56.3% by the BAC system, which also confirmed the conclusion in Section 3.2.1. Table 4 shows that BAC has a good removal effect on DON, and it was speculated that BAC also has a good control effect on the nitrogenous disinfection by-product (N-DBP) precursor, which was confirmed by Zheng et al. [32].

Table 4. DOC value of each component.

	Biopolymers	Humic	Building Blocks	LMW-Acids	LMW-Neutrals
UV/H ₂ O ₂ influent	0.59 (16)	1.82 (50)	0.51 (14)	0 (0)	0.75 (20)
UV/H ₂ O ₂ effluent	0.39 (11.5)	1.56 (46.2)	0.53 (15.7)	0.03 (1)	0.87 (25.7)
BAC effluent	0.31 (17)	1.32 (54)	0.40 (15)	0 (0)	0.38 (14)

Note: the italics in brackets indicate the proportion (%) of each component to total DOC value, and the units of DOC value is mg/L.

3.3. Analysis of Biofilms in Activated Carbons

3.3.1. Changes of Biomass Concentration and Inner DOC Removal Rate of Activated Carbon

The biomass on the surfaces of activated carbons at different times and depths of carbon layers was measured by the phospholipid method. The results are shown in Figure 6a. The growth rate of biomass at the influent section of the activated carbon column was the fastest, since the influent section can absorb more organic matters, and the growth of microorganisms is basically not limited by nutrients and shows an exponential growth trend, since the decomposition of H₂O₂ will produce O₂, which increases the DO concentration (10.5 mg/L) in the influent section, the growth of microorganisms is also not limited by DO concentration. The adsorption of activated carbon on organic matters was

shown as a decrease in DOC value. In Figure 6b, at the initial stage of operation, when the depth of the carbon layer increased, the DOC value significantly decreased. The lack of nutrients caused the biomass growth rate of the middle section and effluent section to be lower than that of the influent section, which results in vertical stratification.

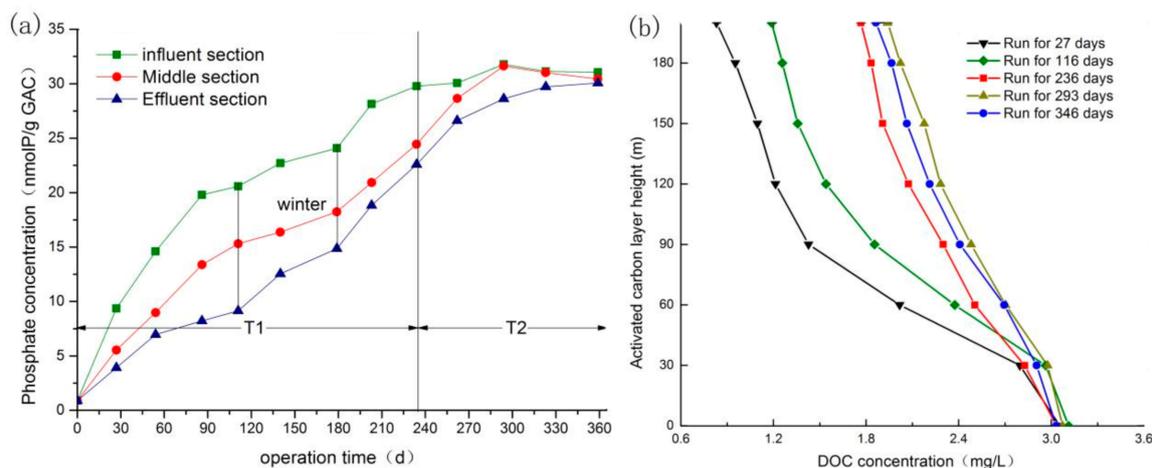


Figure 6. (a) Biomass change of carbon surfaces with different carbon depths (GAC 1 = 30 cm, GAC 2 = 90 cm, GAC 3 = 150 cm); (b) DOC removal rates in the BAC (27 days, 116 days, 236 days and 346 days).

In addition, Figure 6a shows that the growth rate of biomass in the influent and middle sections in winter decreased, while the increase in biomass in the effluent section is faster than that in the influent and middle sections. The reason was that the activity of microorganisms was reduced under low-temperature conditions, which reduced the degradation of microorganisms on organic matters [25]. In addition, the organic matter in water easily combines into macromolecular organic matter under low-temperature conditions, which increased the difficulty of adsorption and removal [33]. Therefore, the content of organic matters in the effluent section increased and the growth rate of biomass accelerated.

From the perspective of time change, the biofilms rapidly accumulated in the T1 stage (the first 240 days) and subsequently gradually stabilized, and the vertical stratification in the activated carbon column gradually disappeared after 293 days. Compared with other research results, the biofilms on the surface of the activated carbon column had longer mature and stable time. Oriol's study on BAC showed that the stable biofilms were formed after 180 days of operation, and the vertical stratification disappeared after 280 days [34]. This difference may be attributed to the temperature difference of water. The experiment was run in the local winter from 120 to 180 days, and the water temperature was 2.5–5 °C. The temperature reduced the metabolism of microorganisms, led to the decrease in the utilization rate of microorganisms and nutrients in the biofilm, and inhibited the growth of biofilms [35].

3.3.2. Microbial Richness and Diversity

To further identify and analyze the growth characteristics of activated carbon bed biofilms, a Roche 454 high-flux sequencing system was applied to analyze the microbial community of the activated carbon biofilm, and the results are shown in Table 5. The optimized sequence numbers of the three samples were 30,515 (GAC 1), 31,935 (GAC 2), and 29,917 (GAC 3), and the coverage rate of all samples was >99%. Three samples were divided into 1160 (GAC 1), 1215 (GAC 2) and 1181 (GAC 3) OTUs with a difference of 3%, and the theoretical maximum OTU values calculated through Chao 1 theory were 1570 (GAC 1), 1612 (GAC 2) and 1594 (GAC 3), the above data showed that in the up-flow UV/H₂O₂-BAC process, the microbial species were more abundant in the middle section than in the influent and effluent sections. This finding may be related to the up-flow

operation mode. The larger flow shear force in the influent section was not conducive to the formation of thicker biofilms. Silvana's study also showed that the highest biomass accumulation on the surfaces of activated carbons in the activated carbon pool was in the middle section [36]. In addition, the low concentration of H₂O₂ (0.20–0.25 mg/L) in the middle section can promote the development of microbial communities [37], the analysis of the Simpson index and Shannon index showed that the biodiversity in the middle section and effluent section was significantly higher than that in the influent section. Hence, we speculate that the environment with a high concentration of H₂O₂ (5.0–6.0 mg/L) in the influent section has a "screening" effect, which reduces the biodiversity.

Table 5. Diversity estimators of microbial communities in GAC 1, GAC 2 and GAC 3 ($\alpha = 0.03$).

Sample	OTUs	Chao 1	ACE	Shannon	Simpson	Coverage
GAC 3	1181	1594	1265.32	5.79	0.77	99.1%
GAC 2	1215	1612	1295.44	5.80	0.73	99.2%
GAC 1	1160	1570	1210.12	5.90	0.61	99.3%
O ₃ -BAC-1	291	-	-	4.39	0.757	-
O ₃ -BAC-2	282	-	-	4.57	0.821	-

Note: O₃-BAC-1 was middle section, O₃-BAC-2 was effluent section.

Our previous collaborator Han had used the identical retreatment process to study the microbial community on the surface of activated carbon in the O₃-BAC system, by comparing with his results (Table 5), we found that the Shannon index and OTUs of the microbial community affected by H₂O₂ were higher than that affected by O₃ in the identical-particle-size activated carbons [5], which indicated that H₂O₂ had lower inhibition on the development of the microbial community than ozone.

3.3.3. Comparison of Biological Community Structures

Figure 7 shows the microbial composition and clustering relationship of the three samples at the phylum level. There was a correlation between the microbial populations in the middle section and effluent section. The dominant species in the three samples were *Proteobacteria*, *Planctonella*, *Acidobacteria*, *Chloroflexi*, *Actinobacteria* and *Bacteroidetes*. Previous studies have confirmed the correlation between *Proteobacteria* and organic matter removal, and *Proteobacteria* also had an important contribution to denitrification [38]. Its content sequence was GAC 3 > GAC 2 > GAC 1; since the H₂O₂ content decreases with the increase in carbon layer depth, the effluent section has basically no H₂O₂, so the existence of H₂O₂ may have an inhibitory effect on *Proteobacteria*. Wang et al. showed that anaerobes were sensitive to H₂O₂, and the presence of H₂O₂ limited their growth [37]. Therefore, H₂O₂ can reduce their abundance by inhibiting the growth of anaerobes in *Proteobacteria*. *Chloroflexi* was detected as the dominant bacteria on the activated carbon biofilm and principally responsible for the degradation of carbohydrates and cellular substances [39]. As the concentration of nutrients in the activated carbon filter gradually decreased, the richness of *Chloroflexi* along the profile exhibited a gradual decrease. *Acidobacteria* is a type of acidophilic bacteria with high content in soil [40]. Combined with the LC-OCD analysis, we can speculate that it contributes to the degradation of SOM by BAC. In addition, *Acidobacteria*, which can degrade H₂O₂, has a relatively high content in the influent section.

Figure 8 mainly depicts the relative abundancies of the main genus level of GAC 1, GAC 2, and GAC 3. Compared with the O₃-BAC process in our previous study [25], there was a significant difference in genus level. The main genus of the O₃-BAC filter was *Nevskia*, *Flavobacterium*, *Acidovorax*, *Methylomicrobium* and *Pseudomonas*. In the UV/H₂O₂-BAC process, the main genera of the activated carbon filters were *Gemmata*, *Gaiella*, *Leptospirillum*, *Pirellula*, and *Planctomyces*. *Nitrospira* is a nitrifying bacteria, which can oxidize nitrite to nitrate [41]. In Figure 8, the *Nitrospira* content in GAC 1 was significantly higher than those in GAC 2 and GAC 3, and we speculated that nitrification mainly occurs in the influent section of BAC. *Rhodoplanes*, a genus of Proteus, was an anaerobe [42]. Its

content sequence was GAC 3 > GAC 2 > GAC 1, which also proved the inhibition of H₂O₂ on *Proteobacteria*. *Rhodobacter* is an important participant in the degradation of organic pollutants in water [43], and its abundance order was GAC 1 > GAC 2 > GAC 3, which implies that the abundance of *Rhodobacter* was positively related to the concentration of organic matters in the water environment. The effluent section with low DOC concentration cannot form the dominant species of *Rhodobacter*. Similar rules can be observed from the analysis of microbial communities of activated carbon with different carbon ages. *Rhodobacter* dominant species can be formed in the aged carbon (6 years) with a relatively high organic matter content in the effluent but not in new carbon [8].

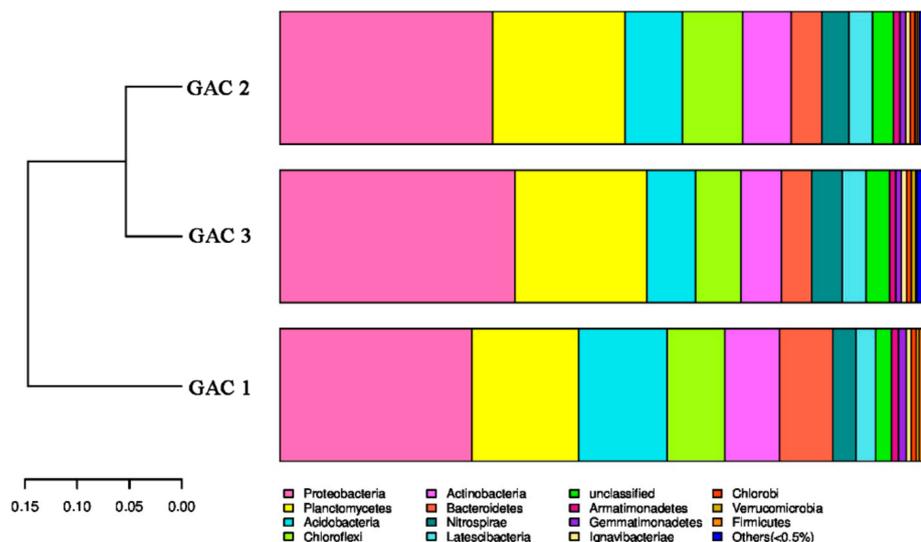


Figure 7. Microbial population classification of activated carbons from different carbon depths (phylum-level).

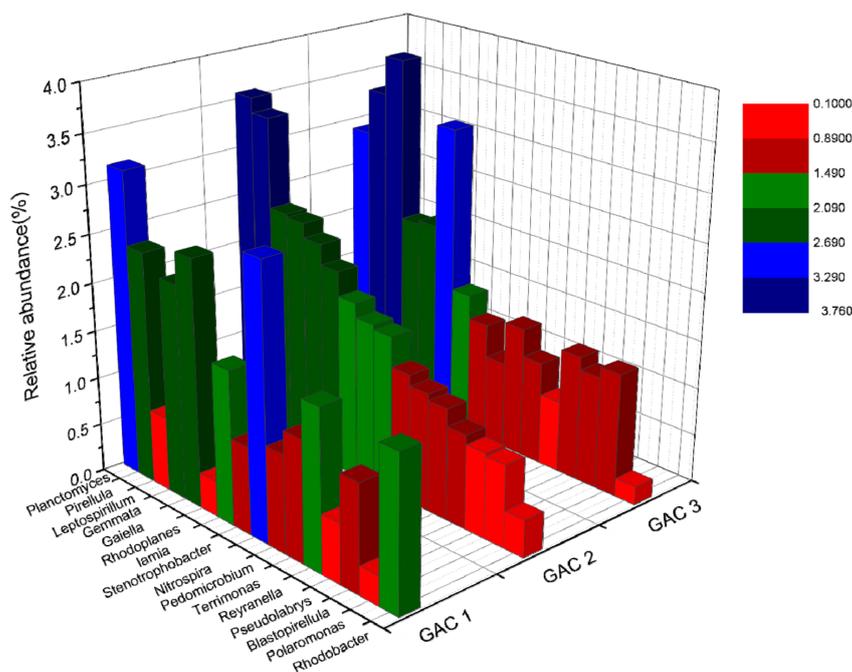


Figure 8. Microbial population classification of activated carbons from different carbon depths (class-level).

4. Conclusions

In this work, the organic matter removal and microbial development of the UV/H₂O₂-BAC process were studied in detail. The results showed that water temperature reduces the efficiency of the UV/H₂O₂ process by affecting the photon production, and the calculated contribution rate of adsorption to the DOC removal in BAC after one year of operation was approximately 14.2%; UV/H₂O₂ process can increase the amounts LMW-neutrals, and SUVA₂₅₄ value is not suitable for characterizing DBP precursors in water after advanced oxidation treatment, and BAC has a good removal effect on SOM. The growth rate of biomass at the influent section of the activated carbon column was the fastest. Microbial species were more abundant in the middle section than in the influent section and effluent section. The high concentration of H₂O₂ (5–6 mg/L) in the influent section has a “screening” effect, which reduces the biodiversity, and a low concentration of H₂O₂ (<0.25 mg/L) promotes the development of the microbial communities. Shannon index shows that H₂O₂ had lower inhibition on the development of microbial community than ozone. H₂O₂ can reduce the *Proteobacteria* abundance by inhibiting the growth of anaerobes, *Acidobacteria* may contribute to the degradation of SOM, *Nitrification* mainly occurs in the influent section of BAC. The effluent section of BAC with low DOC concentration cannot form the dominant species of *Rhodobacter*.

Supplementary Materials: The following are available online at <https://www.mdpi.com/2073-4441/13/4/565/s1>, Figure S1: H₂O₂ utilization rate by UV/H₂O₂ process, Table S1: characteristics of the activated carbons, Table S2: the contents of THMs and HAAs precursor during UV/H₂O₂ process.

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Conflicts of Interest: We declare that we have no financial and personal relationships with other people or organizations that can inappropriately influence our work, there is no professional or other personal interest of any nature or kind in any product, service and/or company that could be construed as influencing the position presented in, or the review of, the manuscript entitled “Research About Organic Matter Removal and Biofilms Development of Pilot-Scale UV/H₂O₂-BAC Process”.

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