Article

Performance and Microbial Diversity in a Low-Energy ANF-WDSRBC System for the Post-Treatment of Decentralized Domestic Wastewater

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Abstract: Recently, more decentralized wastewater treatments are of great interest for rural regions. In this work, a novel ANF-WDSRBC system combined with an anoxic filter (ANF) and a four-stage water-dropping-self-rotating biological contactor (WDSRBC) was designed as a post-treatment option. With a total hydraulic retention time (HRT) of 8.8 h and reflux ratio of 1:1, the ANF-WDSRBC system was operated 160 days. The results showed the ANF-WDSRBC system had better performance without mechanical aeration devices, the removal efficiencies of chemical oxygen demand (COD), ammonia (NH$_4^+$–N) and total nitrogen (TN) were 61.4% ± 4.3%, 86.1% ± 3.7%, and 54.5% ± 3.9%, respectively. By means of high-throughput MiSeq sequencing, the results suggested that Proteobacteria, Bacteroidetes, Firmicutes, and Chloroflexi were the predominant phyla in the system. In the WDSRBC units, Nitrosomonas, Nitrospira, Bacillus, and Nitrospira were the main genera to take part in nitrification. Longilinea, Bellilinea, Thiobacillus, and Thauera in the ANF unit were the main genera to participate in denitrification and organic matters degradation. The novel ANF-WDSRBC system had great potential in the post-treatment of decentralized domestic wastewater.

Keywords: decentralized domestic wastewater; ANF-WDSRBC; nitrification; denitrification; high-throughput MiSeq sequencing

1. Introduction

In recent years, more and more attention has been focused on the treatment of decentralized domestic wastewater by anaerobic techniques in developing countries, particularly in the rural areas of China [1–4]. That may be attributed to the energy recovery and low sludge production in anaerobic bioreactors [1–3]. Despite of these advantages, anaerobic techniques are not widely adopted due to the lack of nitrogen removal capability [4–7]. Even though nitrogen is one of the important nutrients for agriculture and landscaping irrigation reuse [8], nitrogen also plays a major role in oxygen depletion and eutrophication in the receiving water bodies [9]. Therefore, it is necessary to reduce nitrogen concentration of the decentralized domestic wastewater and polish up the effluent of anaerobic reactors using a post-treatment system.

It was reported that aerated filters [10], water-dropping aeration submerged biofilm reactor (WDABR) [4], aerobic membrane bioreactor (MBR) [6,7], stabilization ponds [11], and constructed wetlands (CW) [12,13] had been applied in decentralized domestic wastewater treatment. Nevertheless, there still have many challenges, such as high energy consumption in aerobic systems, low organic load rate, flow clogging of constructed wetlands and high land footprint of stabilization ponds existed during the whole process. Rotating biological contactors (RBCs) are cost-effective and flexible attached growth bioreactors, which are widely used to treat various wastewater including domestic wastewater [14–17], dye wastewater [18], and landfill leachate [19]. The RBCs are fascinating options
to decentralized wastewater treatment due to high specific surface area, high biomass content, low footprint, and easy operation [20]. Even though RBCs are mostly reliable, they still have some shortcomings in terms of costs and stable nitrogen removal efficiency. Compared to constructed wetlands and water-dropping aeration submerged biofilm reactors, RBCs had higher ammonium removal efficiency but higher energy costs [4,21].

In order to further save the energy costs and improve the performance of RBCs, great effort has been spent on modifying RBCs configuration, such as self-rotating discs [22,23], net-like rotating biological contactor [24], and RBC-MFC [25]. Self-rotating discs (SRDs) are cost-effective devices to improve the self-purification efficiency of water stream via increasing dissolved oxygen in the water stream with natural mechanical energy of rivers [22]. Although SRDs has been proven efficient in river self-purification, there is a little information about the application of SRDs on treating decentralized domestic wastewater.

Microbial communities are the main drivers of biological wastewater treatment. Knowledge regarding the microbial consortia in biological systems is of great importance for comprehensive understanding the performance and stability of biological reactors [26,27]. In recent years, molecular biology tools have been widely developed to determine the microbial communities in the wastewater treatment [15,25,26]. It is necessary to identify the microbial compositions and structures of novel biological reactors by this way. Currently, high-throughput sequencing technology is becoming one of the efficient molecular tools for analyzing the microbial diversity. However, there is no study focused on the relationship between microbial communities and nitrogen removal in the SRD system.

To overcome these aforementioned challenges, based on the SRD device configuration, we explored a modified Anoxic/Oxic (A/O) process of an anoxic filter (ANF) and four-stage water-dropping-self-rotating biological contactor (WDSRBC) as an alternative post-treatment. In this system, the ANF unit packed with wool-felt was mainly used for COD removal and denitrification with WDSRBC effluent reflux. The WDSRBC units were used for nitrification via natural mechanical energy of wastewater. Unlike the traditional A/O activated sludge process with complex mechanical mixing devices in the anoxic unit, this system enhanced denitrification for nitrogen removal in the ANF unit with packing wool-felt carrier. The wool-felt carriers were suitable for deploying high biomass and reducing washout of biomass. Without mechanical aeration devices in the oxic unit for improving oxygen transfer capacity and nitrification, the WDSRBC units are more energy-efficient than traditional RBC, MBR [7], and aeration filter [10] due to water-dropping gravity rotation.

The aim of this study was to evaluate the performance of the ANF-WDSRBC system in treating decentralized domestic wastewater. Moreover, high-throughput MiSeq sequencing was used to determine the compositions and structures of microbial communities in each unit of the system.

2. Materials and Methods

2.1. Experimental Setup

Figure 1 shows the schematic diagram of the experimental set-up of the ANF-WDSRBC system. The system consisted of an anoxic filter and a four-stage WDSRBC made of PVC sheets. The ANF (ø150 mm × 1500 mm) packed with vertical wool-felt carriers had a 25 L effective volume. Each WDSRBC unit (350 mm × 200 mm × 150 mm) had a 7.5 L effective volume and 15 discs with 150 mm diameter. The discs of WDSRBC were 40% submerged. The water-dropping height between each WDSRBC was 0.6 m. The wastewater first pumped through the ANF and then drained to the WDSRBC. In order to utilize the carbon source from wastewater to enhance TN removal, the effluent of WDSRBC was partially recycled to the ANF, serving as a nitrate source. The remaining effluent of the WDSRBC was drained to the wetland for irrigation.
was performed under the following conditions: initial denaturing for 30 s at 98 °C, followed by 30 cycles of 10 s at 98 °C, 30 s at 56 °C, 45 s at 72 °C, and a final extension step of 10 min elongation at 72 °C. The PCR products were confirmed by 2% agarose gel electrophoresis and purified using the PCR products were confirmed by 2% agarose gel electrophoresis and purified using the PCR products were confirmed by 2% agarose gel electrophoresis and purified using the PCR products were confirmed by 2% agarose gel electrophoresis and purified using

### 2.2. Wastewater Characteristics and Seeding Sludge

The ANF-WDSRBC system wastewater was the effluent of the anaerobic filter reactor that treated domestic wastewater obtained from Southeast University located in Wuxi, China. The main characteristics of the influent of the ANF-WDSRBC system are presented in Table 1. The seeding sludge used in the study was received from an aeration unit of a WWTP located at Wuxi city in China. The initial MLSS and MLVSS concentrations of seeding sludge were 3000 mg/L and 4000 mg/L, respectively.

<table>
<thead>
<tr>
<th>pH</th>
<th>COD (mg/L)</th>
<th>NH$_4^+$–N (mg/L)</th>
<th>NO$_3^-$–N (mg/L)</th>
<th>NO$_2^-$–N (mg/L)</th>
<th>TN (mg/L)</th>
<th>DO (mg/L)</th>
</tr>
</thead>
<tbody>
<tr>
<td>7.1 ± 0.2</td>
<td>91.4 ± 21.0</td>
<td>26.7 ± 5.5</td>
<td>0.37 ± 0.2</td>
<td>ND *</td>
<td>37.6 ± 7.1</td>
<td>0.20 ± 0.03</td>
</tr>
</tbody>
</table>

* where, ND: not detect.

### 2.3. Analytical Methods

#### 2.3.1. Chemical Analysis

Concentrations of COD, NH$_4^+$–N, NO$_3^-$–N and TN were measured according to the standard method [28]. Dissolved oxygen (DO) and pH were measured with a DO meter (YSI-DO200, YSI, Yellow Springs, OH, USA) and a pH meter (YSI-pH100, YSI, Yellow Springs, OH, USA).

#### 2.3.2. Microbial Community Analysis by MiSeq Sequencing

In order to study the complete microbial community structures, biofilm samples were collected from each unit (namely ANF, WDSRBC1, WDSRBC2, WDSRBC3, and WDSRBC4) of the system on day 160. The total DNA was extracted using OMEGA Soil DNA Kit D5625-01(Omega Bio-Tek, Norcross, GA, USA) according to the manufacturer’s protocol. The quality and quantity of the extracted DNA was measured by a Nanodrop 1000 spectrophotometer (NanoDrop Technologies, Wilmington, DE, USA).

The V3–V4 region of bacterial 16S rRNA genes was amplified using primers 319F/806R (5'-ACTCCTACGGGAGGCAGCAG-3'/5'-GGACTACHVGGGTWTCTAAT-3'). PCR reactions were carried out in 25 µL mixture contained 25 ng template DNA, 2.5 µL forward and reverse primer (1 µM), and 12.5 µL Premix Ex TaqTM Hot Start Version (Takara Bio Inc., Shiga, Japan). Amplification was performed under the following conditions: initial denaturing for 30 s at 98 °C, followed by 30 cycles of 10 s at 98 °C, 30 s at 56 °C, 45 s at 72 °C, and a final extension step of 10 min elongation at 72 °C. The PCR products were confirmed by 2% agarose gel electrophoresis and purified using

![Figure 1. Schematic diagram and photos of the ANF-WDSRBC system.](image-url)
the AxyPrepDNA Gel Extraction Kit (Axygen, Union City, CA, USA) following the manufacturer’s protocol. At last, the purified amplicons were pooled in equimolar and paired-end sequenced using the Illumina-Miseq platform (Illumina Inc., San Diego, CA, USA). After removing the short and low-quality reads, Illumina-Miseq sequencing analysis obtained 90 782 effective sequences for five samples. The resulting high-quality sequences were clustered into operation taxonomic units (OTU) at 97% similarity level by Mothur. Representative sequences selected for each OTU were assigned a taxonomy using a Ribosomal Database Project (RDP) classifier with a confidence threshold of 80%.

3. Results and Discussion

3.1. Nitrogen Removal of the ANF-WDSRBC System at Different Reflux Ratios

Previous studies have reported the impact of reflux ratio on TN removal in various reactors [4,29,30]. Hiras et al. [14] showed that TN removal efficiency increased up to a reflux ratio of 300% in a two-stage rotating biological contactor. Chiou et al. [29] found that the maximum TN removal efficiency in a pre-denitrification/nitrification biofilter occurred at a reflux ratio of 250%. Our previous study [4] demonstrated the ABR-WDASB system achieved high nitrogen removal efficiency at a reflux ratio of 100%. Based on these previous studies, considering the influent concentrations of COD and TN, the ANF-WDSRBC system started its operation at three different reflux ratios (50%, 100%, and 200%) to treat decentralized domestic wastewater.

As shown in Figure 2, the average nitrogen removal efficiencies of the reflux ratios 50%, 100%, and 200% were 31.4% ± 5.7%, 52.9% ± 2.8%, and 54.5% ± 2.8%, respectively. It indicated that TN removal efficiency increased when reflux ratio increased. This result was consistent with our previous study using the ABR-WDASB system to treat rural domestic wastewater [4]. It was also shown that the reflux ratio ranging from 50% to 100% had a significant promote effect on TN removal efficiency than that of from 100% to 200%. Although denitrification in the ANF unit depended on the amount of returned nitrifying solution, a high reflux ratio might increase the returned nitrifying solution and enhance TN removal [29], but it is not cost-effective due to high energy consumption. Hence, a cost-effective reflux ratio of 100% was chosen for the system operation.

![Figure 2. Effects of reflux ratios (R) on the TN removal efficiencies.](image)

3.2. Performance of the ANF-WDSRBC System at Steady State

In order to evaluate the performance of the system at steady state condition, the ANF-WDSRBC system was operated as a continuous flow with a stable flow of 150 L/day (The reflux ratio 1:1). The performance of the system in terms of COD, ammonia, nitrate, nitrite, and TN are shown in Figure 3.
(over time) and Table 2 (in average). Throughout the whole operation, the system was operated with a total HRT 8.8 h (ANF 4 h, WDSRBC 4.8 h) for 160 days. Even though the influent concentration of COD was fluctuated in range of 54.2–147.4 mg/L, the effluent concentration of COD remained stable, resulting in the total average COD removal efficiency of 61.4% ± 4.3%, which was higher than previous reports for aerated fixed bed (47.6% of COD removal) treating the effluent of UASB [31]. For the ANF unit, the average COD removal efficiency was 43.0% ± 4.0%, while the effluent COD concentration was 51.6 ± 10.4 mg/L. In contrast, the COD removal efficiency was 18.3% ± 3.3% for the four-stage WDSRBC, and the final effluent COD concentration was 35.0 ± 7.7 mg/L. Among these reactors of the system, it was remarkably observed that ANF was the predominant contributor in COD removal with the WDSRBC effluent recirculation.

Figure 3 shows the variation of ammonia concentration in the influent and effluent of each unit during 160 days’ operation. When the influent concentration of ammonia varied from 13.9 to 40.6 mg/L with an average value of 26.7 mg/L, the effluent ammonia concentration of the system basically stabilized at 3.7 ± 1.1 mg/L, corresponding to the average ammonia removal efficiency of 86.1% ± 3.7%. Compared with the COD removal in the ANF-WDSRBC system, the total removal efficiency of ammonia was much higher, indicating that the activity of ammonium oxidation bacteria in the system was high during the whole operation period. Along the flowpaths of the system, the effluent concentration of ammonia decreased from 22.3 ± 4.4 mg/L in ANF unit to 16.9 ± 4.0 mg/L (WDSRBC1), 8.3 ± 1.7 mg/L (WDSRBC2), 6.8 ± 1.2 mg/L (WDSRBC3), and 3.7 ± 1.1 mg/L (WDSRBC4), respectively. WDSRBC1 played a pivotal role in the ammonia removal. It might be due to the fact that the influent concentration of COD in the WDSRBC1 was much lower, favoring the nitrifying biomass growth [32]. Previous studies [4,33] reported that ammonia was always removal via nitrification under aerobic condition, which was highly dependent on dissolve oxygen (DO). Compared to other traditional mechanical aeration methods such as brush aeration and blast aeration, the water-dropping rotating aeration supplied DO at a level of 5.15 ± 0.05 mg/L in the WDSRBC units (Table 2), achieving higher oxygen transfer capacity and lower ammonia effluent with low energy consumption.

Figure 3. Influent, effluent, and removal efficiency of COD throughout the whole operation.

Table 2. Summary of the ANF-WDSRBC process performances.

<table>
<thead>
<tr>
<th>Unit (mg/L)</th>
<th>COD</th>
<th>NH$_4^+$–N</th>
<th>NO$_3^-$–N</th>
<th>NO$_2^-$–N</th>
<th>TN</th>
<th>DO</th>
</tr>
</thead>
<tbody>
<tr>
<td>Influent</td>
<td>91.4 ± 21.0</td>
<td>26.7 ± 5.5</td>
<td>0.37 ± 0.2</td>
<td>ND#</td>
<td>37.6 ± 7.1</td>
<td>0.20 ± 0.03</td>
</tr>
<tr>
<td>ANF</td>
<td>51.6 ± 10.4</td>
<td>22.3 ± 4.4</td>
<td>1.01 ± 0.5</td>
<td>ND#</td>
<td>23.8 ± 5.5</td>
<td>0.35 ± 0.02</td>
</tr>
<tr>
<td>Four-stages</td>
<td>35.0 ± 7.7</td>
<td>3.7 ± 1.1</td>
<td>12.6 ± 3.4</td>
<td>0.23 ± 0.18</td>
<td>17.1 ± 4.2</td>
<td>5.15 ± 0.05</td>
</tr>
<tr>
<td>WDSRBC</td>
<td>8.3 ± 1.7</td>
<td>6.8 ± 1.2</td>
<td>3.7 ± 1.1</td>
<td>0.23 ± 0.18</td>
<td>17.1 ± 4.2</td>
<td>5.15 ± 0.05</td>
</tr>
<tr>
<td>Re * (%)</td>
<td>61.4 ± 4.3</td>
<td>86.1 ± 3.7</td>
<td>-</td>
<td>-</td>
<td>54.5 ± 3.9</td>
<td>-</td>
</tr>
</tbody>
</table>

* where, ND: not detect; * where, Re: Removal efficiency.
The distributions of TN, ammonia, nitrate, and nitrite in each unit of the system are shown in Figure 5. It was observed that ammonia was nitrified in the WDSRBC units and approximately 62.3% of ammonia was found to be converted into nitrate. The nitrogen compound in the effluent mainly existed in the form of nitrate (12.6 ± 3.4 mg/L) and the effluent concentration of nitrite was 0.23 ± 0.18 mg/L. The nitrite concentration was not accumulated, indicating that autotrophic denitrification might occur in the inner biofilm of the WDSRBC system [34,35]. The average TN removal efficiency in the ANF-WDSRBC system was 54.5% ± 3.9% (Table 2), which was similar to previous reports for AT-MBR system treating digested domestic wastewater [7]. The reduction of TN might be attributed to the denitrifier in the ANF. In addition, denitrification might partially occur in the WDSRBC. These results were subject to confirmation by Illumina-MiSeq sequencing analysis.

In order to better evaluate the performance of the ANF-WDSRBC system, the treatment efficiency of this system was compared with other post-treatment systems. The summary of comparison is shown in Table 3. Remarkably, different system configurations entailed variations of system performance. Gao et al. [7] used the AT-MBR system to treat digested domestic wastewater with a total HRT of 8 h, reaching 87% of COD removal, 91% of NH$_4^+$–N removal, and 58% of TN removal. Even though the AT-MBR system found to have the best performance among these systems, the energy consumption of the AT-MBR system was significantly higher than others. On the other hand, ANF-WDSRBC,
which requires lower energy consumption, showed similar \( \text{NH}_4^+\)–N and TN removal efficiencies to the AT-MBR system. Although the COD removal efficiency of the ANF-WDSRBC system was lower than the AT-MBR system, the effluent COD concentration was at a low level of 35.0 ± 7.7 mg/L. In addition, the \( \text{NH}_4^+\)–N and TN removal efficiencies of the ANF-WDSRBC system were higher than those in CW, ABR-WDASB, and AFB system. That might be explained that the ANF-WDSRBC system combined the advantages of WDASB and RBC to enhance oxygen transfer capability and nitrification processing via water-dropping gravity rotation. Consequently, based on a balance between the treatment efficiency and energy consumption, the ANF-WDSRBC system was a relatively better alternative option of post-treatment.

**Table 3.** Comparison of the performance of representative post-treatment systems in terms of COD removal, ammonia removal, TN removal, and energy consumption.

<table>
<thead>
<tr>
<th>Post-Treatment</th>
<th>COD_{inf} (mg/L)</th>
<th>Removal (%)</th>
<th>( \text{NH}<em>4^+)-N</em>{inf} (mg/L)</th>
<th>Removal (%)</th>
<th>TN_{inf} (mg/L)</th>
<th>Removal (%)</th>
<th>Energy Consumption</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>AT-MBR</td>
<td>186.8 ± 10.4</td>
<td>87</td>
<td>44.1 ± 4.8</td>
<td>91</td>
<td>58.8</td>
<td>58</td>
<td>Higher</td>
<td>[7]</td>
</tr>
<tr>
<td>AFB</td>
<td>103 ± 25</td>
<td>47.6</td>
<td>-</td>
<td>-</td>
<td>38 ± 6</td>
<td>21</td>
<td>High</td>
<td>[31]</td>
</tr>
<tr>
<td>ABR-WDASB</td>
<td>167</td>
<td>53.3</td>
<td>22.3</td>
<td>50</td>
<td>22.3</td>
<td>33</td>
<td>Low</td>
<td>[4]</td>
</tr>
<tr>
<td>CW</td>
<td>175 ± 89</td>
<td>72</td>
<td>27.9 ± 14.0</td>
<td>38</td>
<td>55 ± 9.5</td>
<td>32</td>
<td>Lower</td>
<td>[30]</td>
</tr>
<tr>
<td>ANF-WDSRBC</td>
<td>91.4 ± 21.0</td>
<td>61.4 ± 4.3</td>
<td>26.7 ± 5.5</td>
<td>86.1 ± 3.7</td>
<td>37.6 ± 7.1</td>
<td>54.5 ± 3.9</td>
<td>Low</td>
<td>This study</td>
</tr>
</tbody>
</table>

Where, AT anoxic tank; MBR aerobic membrane bioreactor; AFB aerated fixed bed; ABR anoxic biofilm reactor; WDASB water-dropping aeration submerged biofilm reactor; CW: constructed wetland.

### 3.3. Diversity of Microbial Community in the ANF-WDSRBC System

As the core of biological processing, the functional bacteria in each reactor of the ANF-WDSRBC system mainly determined the performance of the system including the removal of COD and nitrogen. In order to get a full understanding of the difference of bacterial communities in each unit of the ANF-WDSRBC system, Illumina-Miseq sequencing was applied to investigate the microbial consortium in ANF, WDSRBC1, WDSRBC2, WDSRBC3, and WDSRBC4, respectively. Figure 6 illustrates the relative abundance of the bacterial community at the phylum level. The bacterial community in the ANF-WDSRBC system demonstrated high diversity, 31 taxonomic categories at phyla level were summarized. The top four bacterial phyla (Proteobacteria, Bacteroidetes, Firmicutes, and Chloroflexi) accounted for 83.1%, 87.3%, 85.3%, 77.8%, and 85.4% of total bacteria in ANF, WDSRBC1, WDSRBC2, WDSRBC3, and WDSRBC4, respectively. Despite the same major phyla in these units, the relative abundances of bacteria were significantly different. The phylum Proteobacteria predominated in the WDSRBC units (52.7%–63.3%) of the system, while it showed a lower relative abundance in the ANF unit (39.2%). These results were similar to the previous study of bacterial communities in activated sludge, in which the phylum Proteobacteria accounted for 36%–65% of the total effective bacterial sequences [26]. Different with Proteobacteria, the dominant phylum Chloroflexi was much higher in ANF (16.7%) than in WDSRBC units (4.9%–5.3%). Interestingly, the relative abundance of the Gram-negative Gemmatimonadetes was higher in WDSRBC units (1.9%–6.4%) than in ANF (0.92%), this is quite different from a previous study which reported that the phylum Gemmatimonadetes accounted for 0.49%–0.51% of the total bacteria in activated sludge [36]. These differences were likely due to the environment factors including dissolved oxygen concentration, the concentrations of organic matter, and nutrients. Different relative abundance of bacterial community in the ANF and WDSRBC units might explain the performance of these units in decentralized domestic wastewater treatment. Meanwhile, other phyla in the system might also provide the potential pathways for COD and nitrogen removal, including Acidobacteria, Actinobacteria, Chlorobi, Planctomycetes, Verrucomicrobia, Nitrospirae, and so on [26,37,38].
In order to further get a deeper insight into the bacterial community in the ANF-WDSRBC system, the phylogenetic classification of bacterial sequences from the samples at class level was presented in Figure 7. At the class level, the difference in the bacterial community of ANF and WDSRBC units was significantly distinct. The top two classes in ANF were Betaproteobacteria (16.1%) and Anaerolineae (14.5%), whereas the predominant classes in WDSRBC units were Betaproteobacteria (22.5%–33.5%) and Gammaproteobacteria (13.3%–16.2%). In addition, the relative abundance of the class Bacilli remarkably increased from 1.83% in WDSRBC1 to 8.32% in WDSRBC4. On the contrary, the relative abundances of the classes Sphingobacteria, Bacteroidia, and Flavobacteria gradually decreased along the flowpath of WDSRBC units. Just like at phylum level, the distribution of some classes might also depend on the composition of wastewater, organic loading, and DO concentration in each unit of the system [26].

At a genus level, 18 representative genera identified in ANF and WDSRBC units were shown in Table 4. Among these genera, the detected nitrifying bacteria in WDSRBC units mainly included those from genera Nitrosomonas (0.23%–1.59%), Nitrospira (2.59%–4.01%), Bacillus (0.73%–4.29%), and Nitrobacter (0.96%–2.39%). In this study, high ammonia removal efficiency in the WDSRBC unit might be due to the leading role of these main nitrifying bacteria. Unlike the traditional method which misses the minor functional bacterial community, the minor genera Nitrobacter (0.03%–0.10%), Pseudomonas (0.05%–0.20%), and Luteimonas (0.03%–0.14%) were also detected by high-throughput Illumina-Miseq sequencing. These minor genera might coexist and contribute considerably to the nitrification of the system process. Even though previous study reported that most heterotrophic bacteria were denitrifiers [39–41], the denitrifiers in ANF mainly included the bacteria from genera

Figure 6. Bacterial community structures in units at phylum level.

Figure 7. Bacterial community structures in units at class level.
Thiobacillus (2.01%), Thauera (1.02%), and Bacillus (2.43%). The genus Thauera belonging to the class Betaproteobacteria was also detected in the activated sludge system [33]. Windey et al. reported that Bacillus played important roles in denitrification [42]. In our study, TN removal efficiency in the ANF was much higher than in WDSRBC (Figure 5), indicating that these genera might be key contributors to converting nitrate to nitrogen gas in ANF unit. In addition, based on the Illumina-Miseq sequencing platform, the minor genus Nitratifactor (0.02%), which likely participated in denitrification processes, was identified only in the ANF unit. Interestingly, the denitrification genus Flavobacterium (0.18%–0.58%) was detected in WDSRBC units, which was also identified in the aerobic tanker [43]. It indicated that some bacteria of the genus Flavobacterium probably had the denitrification capability under aerobic conditions to participate in TN removal.

Table 4. The relative abundance of dominant bacterial populations at genus level.

<table>
<thead>
<tr>
<th>Phylum</th>
<th>Class</th>
<th>Genus</th>
<th>ANF</th>
<th>WDSRBC1</th>
<th>WDSRBC2</th>
<th>WDSRBC3</th>
<th>WDSRBC4</th>
</tr>
</thead>
<tbody>
<tr>
<td>Proteobacteria</td>
<td>Alphaproteobacteria</td>
<td>Nitrobacter</td>
<td>0.00</td>
<td>0.05</td>
<td>0.10</td>
<td>0.03</td>
<td>0.03</td>
</tr>
<tr>
<td></td>
<td>Betaproteobacteria</td>
<td>Nitrosomonas</td>
<td>0.12</td>
<td>0.26</td>
<td>1.59</td>
<td>0.47</td>
<td>0.23</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Nitrosospira</td>
<td>1.17</td>
<td>2.59</td>
<td>3.25</td>
<td>3.95</td>
<td>4.01</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Thiobacillus</td>
<td>2.01</td>
<td>3.70</td>
<td>4.12</td>
<td>3.78</td>
<td>2.96</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Thauera</td>
<td>1.02</td>
<td>0.04</td>
<td>0.04</td>
<td>0.07</td>
<td>0.03</td>
</tr>
<tr>
<td></td>
<td>Epsilonproteobacteria</td>
<td>Nitratifactor</td>
<td>0.02</td>
<td>0.02</td>
<td>0.04</td>
<td>0.06</td>
<td>0.03</td>
</tr>
<tr>
<td></td>
<td>Gammaproteobacteria</td>
<td>Latemones</td>
<td>0.03</td>
<td>0.14</td>
<td>0.06</td>
<td>0.08</td>
<td>0.03</td>
</tr>
<tr>
<td>Bacteroidetes</td>
<td>Flavobacteria</td>
<td>Flavobacterium</td>
<td>0.25</td>
<td>0.18</td>
<td>0.30</td>
<td>0.58</td>
<td>0.32</td>
</tr>
<tr>
<td>Sphingobacteria</td>
<td>Meniscus</td>
<td>4.69</td>
<td>0.04</td>
<td>0.07</td>
<td>0.10</td>
<td>0.10</td>
<td>0.04</td>
</tr>
<tr>
<td>Firmicutes</td>
<td>Bacillus</td>
<td>Bacillus</td>
<td>2.43</td>
<td>0.73</td>
<td>4.29</td>
<td>1.12</td>
<td>2.65</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Falsibacillus</td>
<td>0.52</td>
<td>1.07</td>
<td>2.41</td>
<td>2.39</td>
<td>3.81</td>
</tr>
<tr>
<td>Clostridia</td>
<td>Anaerovorax</td>
<td>1.07</td>
<td>0.01</td>
<td>0.00</td>
<td>0</td>
<td>0</td>
<td>0.01</td>
</tr>
<tr>
<td>Chloroflexi</td>
<td>Anaerolineae</td>
<td>Bellilinea</td>
<td>4.55</td>
<td>0.43</td>
<td>0.36</td>
<td>0.35</td>
<td>0.31</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Longilinea</td>
<td>7.50</td>
<td>1.82</td>
<td>2.02</td>
<td>2.11</td>
<td>1.80</td>
</tr>
<tr>
<td>Caldilineae</td>
<td>Caldilinea</td>
<td>1.91</td>
<td>1.57</td>
<td>2.10</td>
<td>2.47</td>
<td>3.12</td>
<td></td>
</tr>
<tr>
<td>Dehalococcoides</td>
<td>Dehalogenimonas</td>
<td>0.29</td>
<td>1.69</td>
<td>0.53</td>
<td>1.11</td>
<td>0.60</td>
<td></td>
</tr>
<tr>
<td>Nitrospira</td>
<td>Nitrospira</td>
<td>0.53</td>
<td>1.37</td>
<td>0.96</td>
<td>2.39</td>
<td>0.47</td>
<td></td>
</tr>
</tbody>
</table>

The other representative genera Meniscus (4.69%), Longilinea (7.5%), Bellilinea (4.55%) were typical identified in the ANF unit. Members of these genera may be correlating to both carbohydrate and amino acid degradation [44,45]. This might partially explain the reason for COD removal in ANF units. The genus Caldilinea (1.57%–3.12%) was detected in the WDSRBC unit, which may affect the formation of sludge flocs [46]. Taken together, the microbial communities coexisted in each unit ensured the desired performance of ANF-WDSRBC system.

4. Conclusions

The combined ANF-WDSRBC system was used to treat decentralized domestic wastewater under continuous-flow operation with an HRT of 8.8 h and reflux ratio of 1:1. The average removal efficiencies of 61.4% ± 4.3%, 86.1% ± 3.7%, and 54.5% ± 3.9% for COD, NH₄⁺-N, and TN were achieved by the system. Compared to some representative post-treatment systems, the ANF-WDSRBC showed a relatively better performance with low energy consumption. Deeper sequencing results indicated that Proteobacteria, Bacteroidetes, Firmicutes, and Chloroflexi were the predominant phyla in the system. The genera Nitrosomonas, Nitrosospira, Bacillus, and Nitrospira in the WDSRBC units played vital roles in nitrification, whereas Longilinea, Bellilinea, Thiobacillus, and Thauera in the ANF unit were the main genera to participate in denitrification and organic matter degradation. Therefore, the ANF-WDSRBC system is a viable option for decentralized domestic wastewater post-treatment.

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Author Contributions: Juanhong Li and Xiwu Lu conceived and designed the experiments; analyzed the data; and wrote the paper.

Conflicts of Interest: The authors declare no conflicts of interest.

References


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