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Dynamics of naphthenic acids and microbial community structures in a membrane bioreactor treating oil sands process-affected water: impacts of supplemented inorganic nitrogen and hydraulic retention time†

 Jinkai Xue,^a Yanyan Zhang,^{ab} Yang Liu^{*a} and Mohamed Gamal El-Din^{*a}

This study was focused on how different operating conditions affected the biodegradation of naphthenic acids (NAs) and the microbial community architectures in an anoxic–aerobic membrane bioreactor (MBR) for oil sands process-affected water (OSPW) treatment. After 442 days of continuous optimization, a supplemented $\text{NH}_4\text{-N}$ concentration of 25 mg L^{-1} and a hydraulic retention time (HRT) of 12 h demonstrated the best removal rates of total classical NAs (37.6%) and total oxidized NAs (23.9%). Neither higher HRTs nor higher supplemented $\text{NH}_4\text{-N}$ concentrations resulted in a better overall removal of NAs. In addition, NAs with larger carbon numbers were generally better degraded, whereas higher cyclicity tended to lessen the biodegradability of NAs. MiSeq sequencing analysis disclosed that orders under Proteobacteria (*i.e.*, Rhodocyclales, Burkholderiales and Nitrosomonadales), Bacteroidetes (*i.e.*, Cytophagales, [Saprospirales] and Flavobacteriales), and Nitrospirae (*i.e.*, Nitrospirales) were the major microbes over the whole study though their relative abundances varied. The results of this study provide insightful information for future studies and application of biological processes for OSPW treatment on a large scale.

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1. Introduction

Due to the huge water demand (3.1 barrels of water per barrel of bitumen produced) in the modified Clarke caustic hot water extraction process in the oil sands industry, the total area of existing tailings ponds (storing oil sands process-affected water, OSPW) in Canada reached 182 km^2 (including associated structures) by the year 2013.¹ Among all the organic acids in OSPW, naphthenic acids (NAs) have been reported as the major contributors to OSPW toxicity and persistency. They are known to be acutely toxic to a range of organisms, including microorganisms, algae, vertebrates and invertebrates.^{2–5} Therefore, NAs are regarded as the target pollutants that need to be treated urgently.^{6–9}

NAs are a wide group of aliphatic and alicyclic, alkyl-substituted carboxylic acids with a generalized formula $\text{C}_n\text{-H}_{2n+z}\text{O}_x$, in which n refers to molecular carbon number, z is zero or a negative even integer indicating hydrogen deficiency

caused by the formation of rings or double bond equivalents,^{4,10} and x refers to the number of oxygen atoms. Species with $x = 2$ are referred to as classical NAs, while those with $x \geq 3$ are termed oxidized NAs (oxy-NAs).¹¹ It has been estimated that more than 20 000 individual NAs are potentially associated with OSPW.¹² Hanging on the ore composition, extraction processes, the tailings age and quantification approaches, the NAs concentrations reportedly varied greatly within the range of $20\text{--}120 \text{ mg L}^{-1}$.^{3,13–15} It has been suggested that NAs with a carbon number $n \leq 21$ and/or linearly grouped carbon rings may be more toxic in OSPW;^{4,16–18} whereas NAs with ring structures tend to be more persistent in environment.² It is, therefore, desirable to remove NAs from OSPW to mitigate its environment impact.

It is suggested that biodegradation is responsible for the natural, but slow dissipation of NAs in tailings ponds.¹⁹ However, the *in situ* biodegradation half-lives of NAs are within a range of 12.8–13.6 years.²⁰ To accelerate the biodegradation of OSPW NAs, it is crucial to develop engineered biological processes that are efficient in mitigating OSPW organic compounds. Choi and Liu performed a study on the treatment of OSPW by using two sequencing batch reactors (SBRs) and achieved an acid extractable fraction (AEF) removal rate of 8.7% and 16.6% at a hydraulic retention time (HRT) of 24 h with the inoculation of activated sludge and mature fine tailings (MFT), respectively.⁶ Hwang *et al.* reported that 13.8% of OSPW AEF and 18.5% of the OSPW parent NAs were eliminated by OSPW

^aDepartment of Civil and Environmental Engineering, University of Alberta, Edmonton, Alberta, T6G 1H9, Canada. E-mail: yang.liu@ualberta.ca; mgamalel-din@ualberta.ca; Fax: +1-780-492-0249; +1-780-492-0249; Tel: +1-780-492-5115; +1-780-492-5124

^bDepartment of Civil Engineering, New Mexico State University, Las Cruces, New Mexico 88003, USA

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indigenous microorganisms in a biofilm reactor with an HRT of 19 h.⁷ Hence, it is suggestive that disparate reactor configurations have different capabilities in degrading OSPW organic substances.^{6,21,22}

Membrane bioreactor (MBR) is finding wider application for municipal and industrial wastewater treatment because of its compactness, high treatment capacity, low sludge production and excellent effluent quality.^{23,24} To investigate the feasibility of MBR for OSPW treatment, an anoxic-aerobic MBR system was performed at an HRT of 48 h, and attained an OSPW NA removal of 25% after the startup period (day 1–425).²⁵ To further improve its performance for OSPW treatment, more effort is needed to optimize the operating conditions.

HRT is one of the essential parameters in MBR operation. It not only affects the extent to which the target contaminants are degraded,²⁶ but also profoundly impacts membrane fouling development.^{27,28} Longer HRTs are usually required for industrial wastewater, especially those containing complex organic contaminants.²⁹ However, the effect of HRT on the degradation of organic pollutants was still unclear as indicated by some studies.^{28,30–32} On the other hand, a shorter HRT is favorable as it reduces the required reactor volume (less investment) for the same treatment capacity. In addition, HRT has an indirect yet profound impact on membrane fouling through altering of sludge properties (*i.e.*, the concentration of mixed liquor suspended solids (MLSS), the production of extracellular polymeric substances (EPS) and the viscosity of sludge).^{33–35} Due to increased membrane flux, shortened HRTs usually result in severer membrane fouling,^{27,28,33} which could increase the operating cost substantially. Therefore, a balance between contaminant removal and operation cost must be considered for HRT selection.²⁹

Further, ammonia monooxygenase (amo) produced by ammonia oxidizing bacteria (AOB) is able to degrade a wide spectrum of refractory organic substrates (including hydrocarbons) through co-metabolism.^{36–40} Nitrifiers are not necessarily stimulated by higher ammonium nitrogen (NH₄-N) concentrations as free ammonia (FA) is inhibitory to many microbial activities including nitrification.⁴¹ Given the alkaline nature of OSPW, the effect of FA should not be ignored. Therefore, it is needed to find out an optimal supplemented NH₄-N concentration to maximize the MBR's performance of NA degradation.

The objectives of this study were: (1) to explore the optimal operation conditions to improve the MBR's NA degradation efficiency; and (2) to characterize the microbial community structure in the MBR with the aim of uncovering the most effective bacterial groups associated with NA removal. According to our literature search, this is the very first study evaluating the performance (particularly the biodegradation of classical and oxy-NAs) and characterizing the microbial communities of an MBR for OSPW treatment under various operating conditions.

2. Methodology

2.1 Source waters, external carbon source and supplemented nutrients

The OSPW was collected from the same tailings pond in Northern Alberta region. The OSPW used for the optimization

operation had a pH of 8.8 ± 0.3 , chemical oxygen demand (COD) of $224.1 \pm 46.3 \text{ mg L}^{-1}$, 5 day biochemical oxygen demand (BOD₅) of $5.4 \pm 0.8 \text{ mg L}^{-1}$, classical NAs of $49.9 \pm 4.0 \text{ mg L}^{-1}$, and oxy-NAs of $39.8 \pm 4.2 \text{ mg L}^{-1}$. Raw OSPW was received in 200 liter barrels and preserved in dark at 4 °C prior to treatment. To facilitate the growth of microorganisms in the system, sodium acetate was supplied as an external carbon (C) source at a constant organic loading rate (OLR) of 125 g COD per (m³ d) throughout the whole operation. To ensure the sufficiency of nitrogen (N), phosphorus (P), and other trace nutrients, supplemented KNO₃, NH₄Cl, NaH₂PO₄, and modified Bushnell-Haas medium (BHM)¹⁷ without KH₂PO₄, Na₂HPO₄, NH₄NO₃, and (NH₄)₂SO₄ were added in the feed water. To avoid undesirable microbial growth in the feed tank, the stock solution of external C source, and the stock solution of external N (to achieve equivalently 75 mg N per L in the feed) and P (to achieve equivalently 10 mg P per L in the feed) source were continuously injected in the MBR at a pre-set rate by using a syringe pump (KDS Infusion Pump, Model 220, KDS Scientific Inc., USA).

2.2 MBR configuration and operating conditions

The configuration of the MBR is illustrated in Fig. S1.† Detailed description on the system configuration, biomass inoculation and acclimatization is available elsewhere.^{25,42} Dissolved oxygen (DO) concentrations in the anoxic tank and the aerobic tank were maintained at a level of $<0.3 \text{ mg L}^{-1}$ and $>4.0 \text{ mg L}^{-1}$, respectively. The retentate in the aerobic tank was recirculated to the anoxic tank by overflow at a recycling ratio of 2.

The different operating conditions examined in this study are tabulated in Table 1. Despite the changes made to NO₃-N and NH₄-N concentrations, the total inorganic nitrogen (TIN, *i.e.*, NH₄-N, NO₂-N and NO₃-N) concentration supplemented to the system was maintained at 75 mg N per L throughout the study. The system automatically performed a 30 second back-washing after every 9.5 minutes of membrane filtration, comprising a 10 minute working phase. The relaxation duration between two working phases was adjusted to achieve a desired equivalent HRT. According to the manufacturer's suggestion, chemical cleaning (soaking the membrane module in 0.1% HClO solution for 1 hour followed by rinsing with running tap water) was applied once the transmembrane pressure (TMP)

Table 1 Operating conditions examined

Stage	Period (days)	Supplemented nitrogen		HRT (h)
		NH ₄ -N (mg L ⁻¹)	NO ₃ -N (mg L ⁻¹)	
A25H48	300–449	25	50	48
A75H48	449–546	75	0	48
A50H48	546–630	50	25	48
A25H72	630–680	25	50	72
A25H12	680–712	25	50	12
A25H24	712–742	25	50	24



exceeded -35 kPa (severe fouling event). Sludge in the MBR was wasted through weekly sampling (30 mL from each tank for solids content measurements) to achieve a calculated solid retention time (SRT) of ~ 187 days.

2.3 Analyses

Methods of conventional water chemistry analyses and acute aquatic toxicity bioassay are described in the (ESI†).

2.3.1 NA measurement. To assess the MBR performance on OSPW NA degradation, ultra-performance liquid chromatography coupled with high-resolution mass spectrometry (UPLC/HRMS)¹¹ was performed to quantify classical and oxy-NAs in the feed and permeate samples at the end of each operating condition stage. With respect to permeate sampling, a volume of 10 mL was collected from totally 2.4 L of permeate water produced over three consecutive days for UPLC/HRMS analysis. The second of the three days was used as the nominal sampling date. To evaluate the MBR's NA degradation performance, percentile removal rates were calculated using the initial and remaining concentrations.

2.3.2 MiSeq sequencing. A PowerSoil® DNA Isolation Kit from Mo-Bio Laboratories, Inc. (CA, USA) was used to extract DNA from collected sludge samples according to the manufacturer's manual. In addition, the fouling layer on the membrane surface was carefully scraped off by using a spatula and collected for DNA extraction. Extracted DNA samples were diluted to a concentration of ~ 20 ng μL^{-1} before sending to a commercial laboratory (Research and Testing Laboratory, Texas, US). Samples were sequenced using Illumina MiSeq sequencing instruments and reagents according to the manufacturer's guidelines. Primers used for DNA amplification prior to MiSeq sequencing were 28F (GAGTTTGATCCTGGCTCAG) and 388R (TGCTGCCTCCCGTAGGAGT). Raw sequence data were processed with the Quantitative Insights Into Microbial Ecology (QIIME, <http://qiime.org>) software package with default settings. Operational taxonomic units (OTUs) were picked through using open-reference clustering algorithm. More details are given in our published reports.^{25,42}

2.3.3 Statistical method. Paired two sample *t*-test was used in this study to compare the MBR performance under different operating conditions with the significance level of 0.05. Pearson correlation coefficient (r_p) was used to estimate linear correlations. The coefficient r_p is a numerical value within the range of -1 and 1 that expresses the strength of the linear relationship between two parameters. An r_p closer to 1 indicates a strong positive correlation. An r_p of 0 signalize that there is no relationship. Values closer to -1 indicate a strong negative relationship between the two variables.

3. Results and discussion

3.1 Classical and oxidized NA removal performance

Table 2 presents total classical NAs concentrations of the feeds and permeates under different conditions and the estimated half-life of classical NAs under each condition. By degrading 37.6% of classical NAs within 12 h, A25H12 stage demonstrated

the shortest half-life (*i.e.*, 17.6 h) of the contaminants among the six stages. In contrast, A25H72 achieved the longest half-life (*i.e.*, 159.7 h) of classical NAs by removing 26.8% within 72 h. The half-lives of classical NAs in all the six operating conditions were substantially shorter than the estimated half-life of 44–240 days in Mahdavi *et al.*⁴³ and Han *et al.*'s¹⁴ suspended biological systems, and even negligible compared with the estimated ~ 13 year *in situ* half-life in tailings ponds.²⁰ The shortest half-life exhibited by our previous batch suspended reactors was 20 days.⁴⁴ In Hwang *et al.*'s continuous biofilm reactor, 18.5% of classical NAs was removed within 19 h (half-life = 64.4 h).⁷ More recently Huang *et al.*'s, integrated fixed-film activated sludge system (IFAS) demonstrated a half-life of 59.0 h of classical NAs.⁴⁵ Therefore, the performance of MBR for OSPW NA degradation could be advantageous over conventional activated sludge systems and biofilm reactors.

Fig. 1 is prepared to examine the effect of n and Z on biodegradation of NAs. A relative abundance of residual classical NAs ($\text{NA}_{\sum ne}/\text{NA}_{\sum ni}$) is used to indicate the extent of biodegradation of NAs with each n . $\text{NA}_{\sum ne}$ denotes the total concentration of NAs with the same carbon number n in the effluent, while $\text{NA}_{\sum ni}$ is the total concentration of NAs with the same n in the influent. Therefore, a lower $\text{NA}_{\sum ne}/\text{NA}_{\sum ni}$ means a better removal of the group of NAs with n . Similarly, a relative abundance of residual classical NAs ($\text{NA}_{\sum |z|e}/\text{NA}_{\sum |z|i}$) is used to indicate the extent of biodegradation of NAs with each $|Z|$. To better disclose how molecular n and $|Z|$ affected the degradation of NAs, Fig. 2 and 3, and Fig. S3 and S4† are plotted based on the NA concentration profiles.

Although the stages differed in operating conditions, similar patterns could be seen for $\text{NA}_{\sum ne}/\text{NA}_{\sum ni}$ over the n range in Fig. 1A and B. For C9–12 and C19–23, a higher $\text{NA}_{\sum ne}/\text{NA}_{\sum ni}$ ratio is associated with a lower n , whereas a plateau or a slightly climbing trend of the $\text{NA}_{\sum ne}/\text{NA}_{\sum ni}$ ratio is observed when n increases from 13 to 18 in Fig. 1A and B. As demonstrated in Fig. 2 and S3,† species with more carbon numbers were generally better degraded within each Z series, which is in agreement with our previous batch study.⁴⁴ In contrast, previous studies indicated that n has either negatively proportional correlation to or minor influence on biodegradation of NAs.^{14,17,46} This discrepancy evidences that the capabilities of biological systems are considerably influenced by their

Table 2 Total classical NA concentrations of the MBR feed and permeate under different operating conditions

Operating condition	Classical NAs (mg L ⁻¹)		Removal (%)	Estimated half-life (h)
	Feed	Permeate		
A75H48	45.3	34.9	23.0	127.6
A50H48	51.8	35.9	30.7	90.7
A25H48	51.4	34.1	33.7	81.1
A25H72	44.7	32.7	26.8	159.7
A25H24	55.3	35.4	36.0	37.3
A25H12	44.7	27.9	37.6	17.6



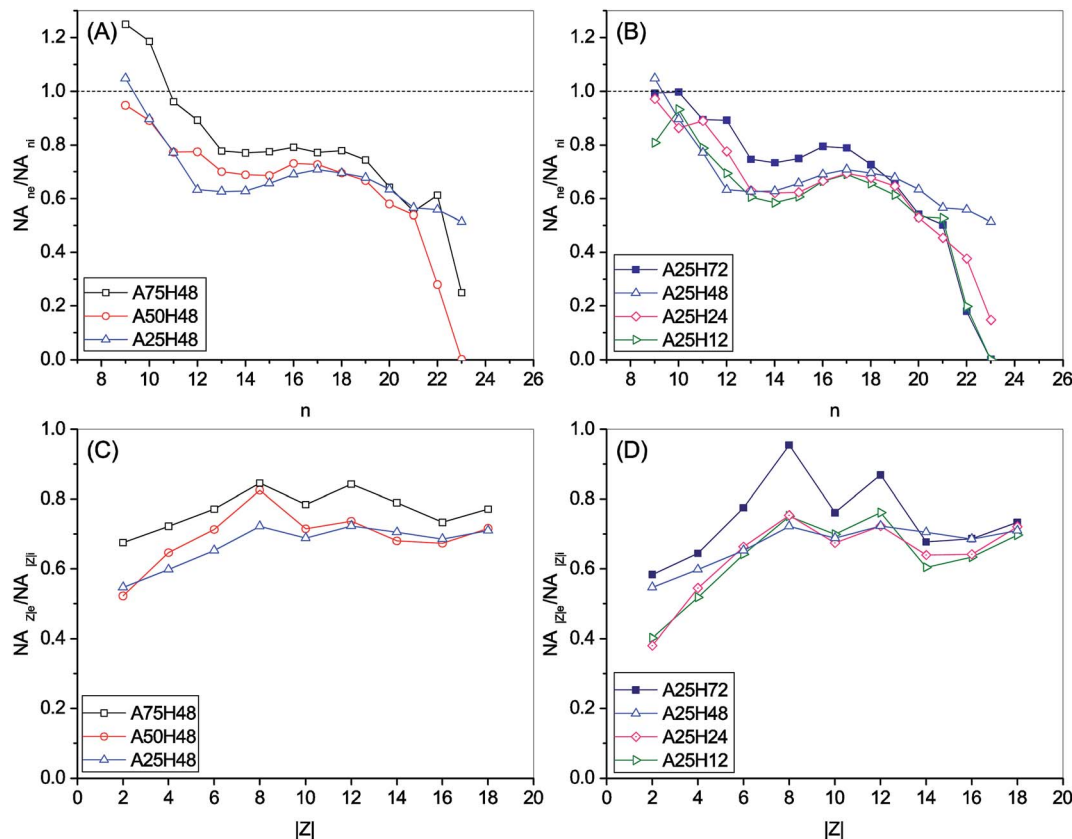


Fig. 1 The relative abundance of residual classical NAs with a carbon number n ($NA_{\Sigma ne}/NA_{\Sigma ni}$) in the MBR permeate under different operating conditions: (A) HRT of 48 h; (B) 25 mg NH_4-N per L and 50 mg NO_3-N per L. And the relative abundance of residual NAs with different cyclicality ($NA_{\Sigma ze}/NA_{\Sigma zi}$) in the MBR permeate under different operating conditions: (C) HRT of 48 h; (D) 25 mg NH_4-N per L and 50 mg NO_3-N per L.

configurations. In addition, it is noticed that, within each Z series, some NAs with smaller carbon numbers (e.g., $n = 9$ and $Z = -6$ in Fig. 2) were even increased after biodegradation. It is thus suggested that some parent NAs with larger carbon numbers were biologically converted into NA species with smaller carbon numbers, which is consistent with our previous observation.^{25,42}

In Fig. 1C and D, for $|Z| = 2$ to 8, the $NA_{\Sigma ze}/NA_{\Sigma zi}$ ratio is positively proportional to $|Z|$, whereas the $|Z|$ range of 9 to 18 witnesses a relatively stable region. It is noticed that within each n series, an increased degree of cyclization (larger $|Z|$) generally resulted in poorer removal efficiency (Fig. 3 and S4†). This trend is not surprising as the biodegradation rate of NAs decreased with increasing cyclicality in both of commercial NAs and OSPW NAs according to previous research.¹⁴ A reduction was observed for all the NAs with specific cyclicality (Fig. 1), which suggested the occurrence of ring cleavage. Some NAs (e.g., $n = 14$ and $|Z| = 10$ in Fig. 3) were even enriched after biodegradation. Hence, it is speculated that microorganisms might convert some parent NAs with higher cyclicality into species with less rings.^{7,25,42}

According to previous studies, the potential biodegradation pathways of aliphatic and alicyclic carboxylic acids include β -oxidation, combined α - and β -oxidation, and aromatization pathways.^{14,47} These researchers suggested that β -oxidation is

the most probable degradation pathways of OSPW NAs in engineered biological systems. Quaternary and tertiary carbons, at β or α position to the carboxylic group, would prevent or impede the proceeding of β -oxidation due to steric hindrance. This explains why the presence of alkyl branches and ring structures result in increased bio-resistance of NAs. In addition, it is claimed that a tertiary carbon in a ring is less favorable than tertiary carbons in an acyclic NAs as the ring creates further steric hindrance. Moreover, the previous researchers suggested that the intermediate product species generated during biodegradation of NAs could be even more recalcitrant to microbes depending on the occurrence and position of tertiary carbons.¹⁴ Furthermore, NAs with larger n are generally more hydrophobic.⁴⁸ Within the same n series, a higher cyclicality tends to decrease the NA hydrophobicity.⁴⁸ Organic compounds of higher hydrophobicity (or higher lipophilicity) could easily attach onto or penetrate the bacterial cell envelope, possibly leading to better biodegradation under certain circumstances.^{49–51} For those particular species, stronger hydrophobic interactions of NAs–NAs, NAs–flocs, and NAs–fouling layer were expected, which would result in elevated local concentrations of hydrophobic NAs. And the locally elevated concentrations of NAs would benefit overall biodegradation of them as a threshold level of NAs exists for microorganisms to initiate the biodegradation process.⁵²



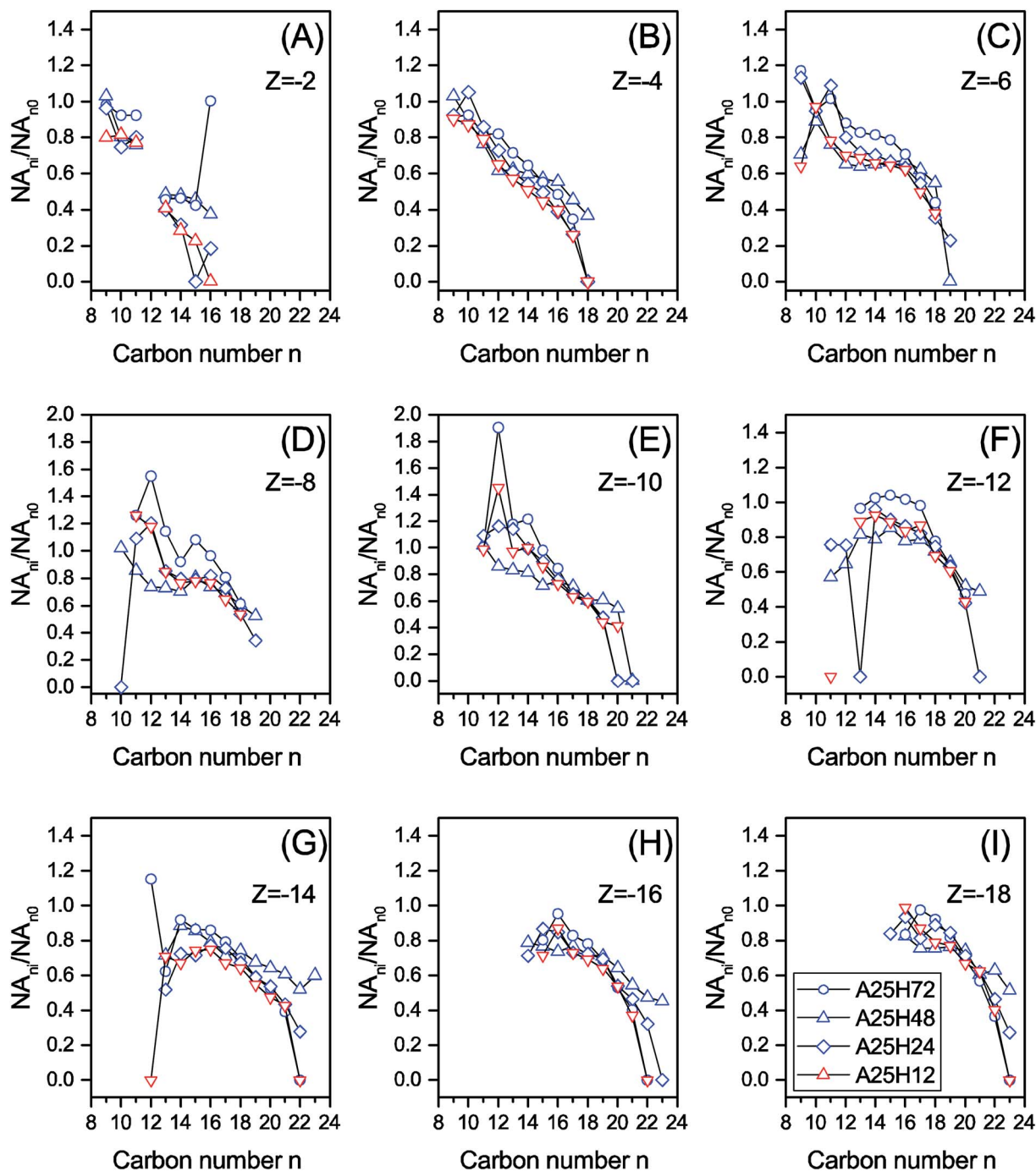


Fig. 2 The relative remaining abundance of NAs with different carbon numbers (NA_{ni}/NA_{n0}) within each Z series of the permeate samples at different HRTs.

Oxy-NAs concentrations of the MBR feeds and permeates were also monitored (Fig. 4). O3 and O4 oxy-NAs are conceivably worth more attention than O5 and O6 oxy-NAs owing to their dominating abundance. O4, O5, and O6 oxy-NAs were all degraded to a certain extent at each stage. In contrast, more O3 oxy-NAs were generated after the MBR biological process at all stages except A25H48 and A25H12. O3 (probably hydroxylated NAs) and O4 oxy-NAs (probably hydroxylated NAs) are intermediates during

aerobic biodegradation of classical NAs.^{9,14,20,53,54} According to Martin *et al.*, O3 species may be byproducts of more cyclic classical NAs.⁹ The poorer removal of oxy-NAs might be attributed to the bioconversion of classical NAs. Some researchers reported unappreciable changes of O3 and O4 oxy-NAs after 98 days¹⁴ and 84 days⁵⁵ of incubation. Therefore, the MBR in this study is competent for classical NAs and oxy-NA degradation.



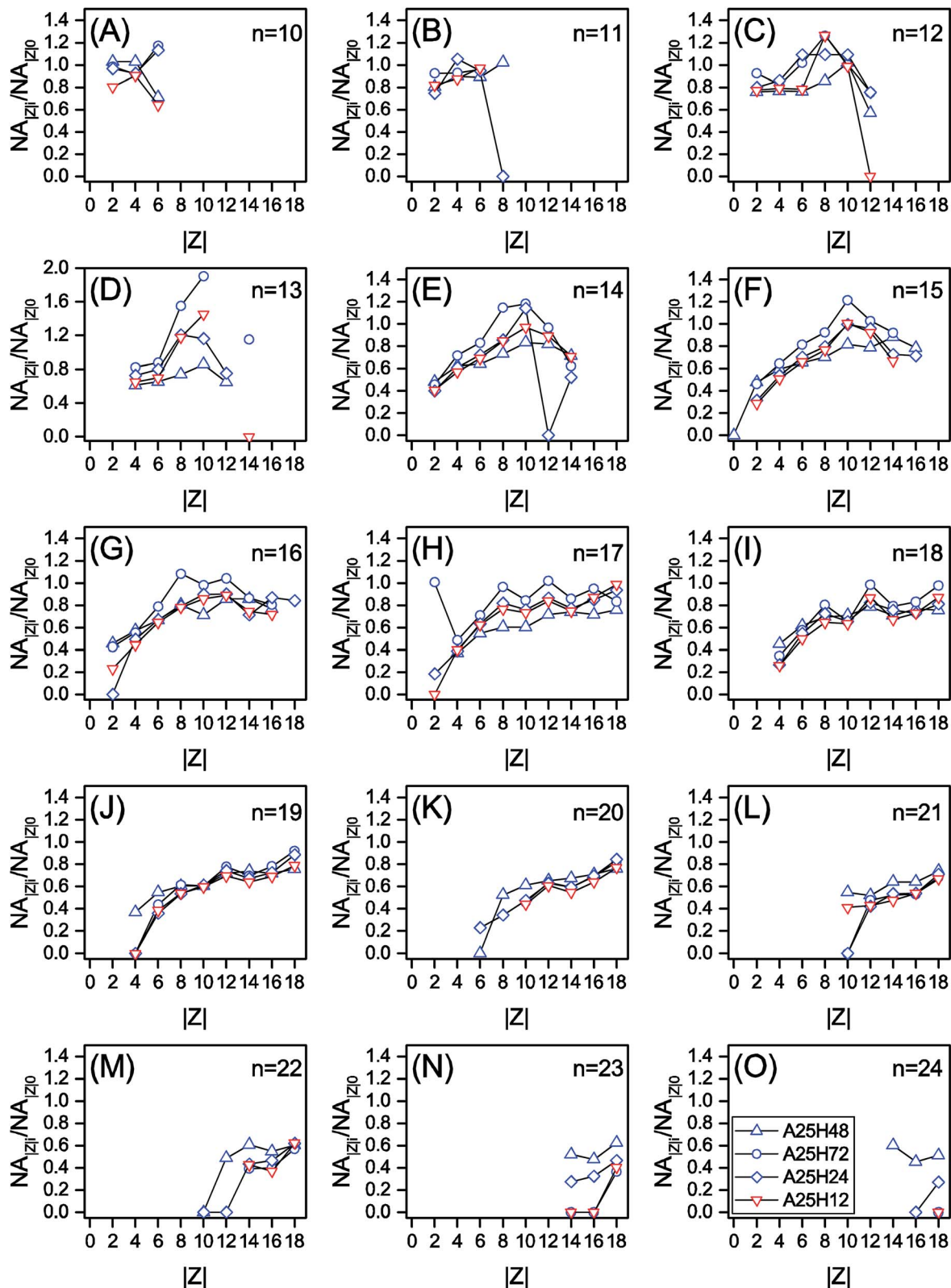


Fig. 3 The relative remaining abundance of NAs with different cyclicity (NA_{zi}/NA_{z0}) within each n series of the permeate samples at different HRTs.

3.1.1 Effect of supplemented nitrogen composition.

Despite that the total concentration of NH_4-N (including OSPW indigenous and supplemented NH_4-N) varied over the whole

operation duration, the MBR constantly produced a permeate NH_4-N concentration of $\sim 0 \text{ mg L}^{-1}$ under all operating conditions, indicating its excellent ammonium oxidation stability



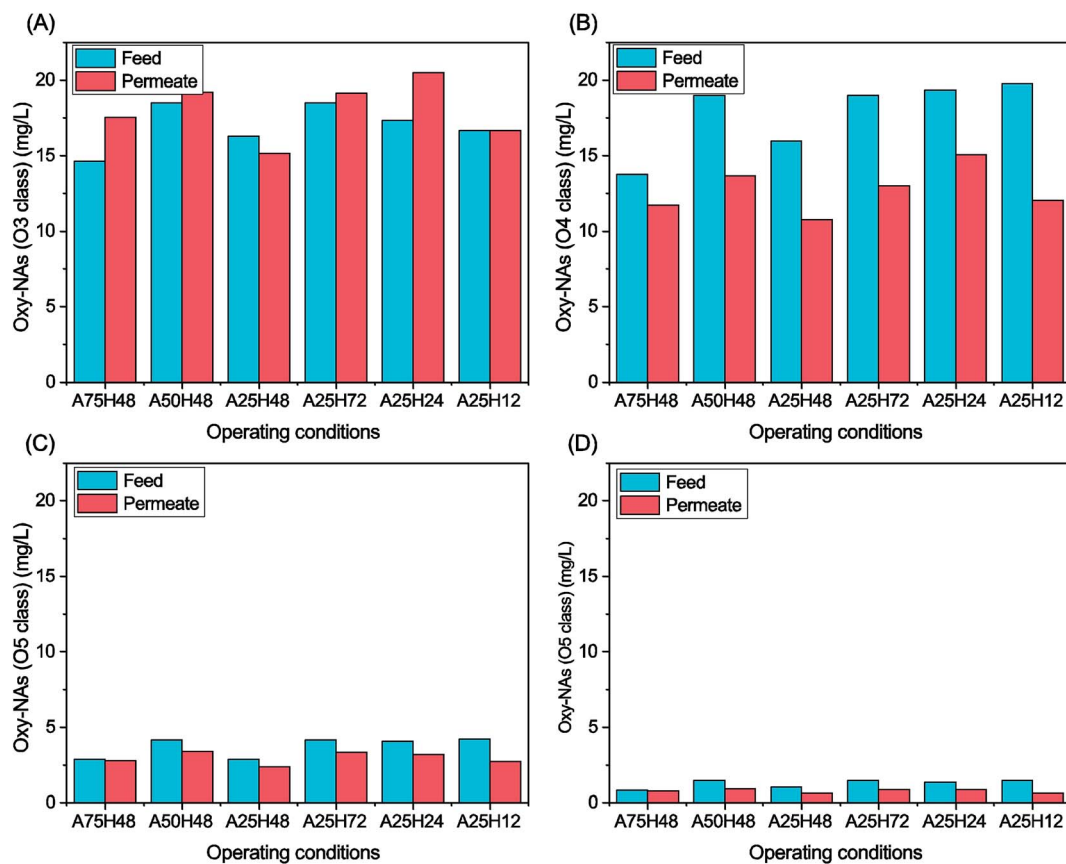


Fig. 4 Oxy-NA concentrations of the MBR feed and permeate under different operating conditions: (A) oxy-NAs with O3; (B) oxy-NAs with O4; (C) oxy-NAs with O5; and (D) oxy-NAs with O6.

(Tables S1 and S2,† and Fig. S2†). The $\text{NO}_2\text{-N}$ concentrations of the MBR permeate were found negligible (≤ 0.25 mg N per L) throughout the study (Fig. S2C†), reflecting the insusceptibility of nitrite oxidizing microorganisms against operating condition variation. With respect to TIN removal performance, the MBR showed more fluctuations as the operating conditions changed, suggesting that denitrifiers might be more susceptible to changes of operating conditions and needed more time to adjust themselves. In addition to volatilization and biomass assimilation, the majority of $\text{NH}_4\text{-N}$ was oxidized into $\text{NO}_3\text{-N}$ by nitrifiers (Tables S1 and S2,† and Fig. S2†). The biomass changes over time are presented in Table S3.†

Discussion within the following paragraphs will therefore be focused on how different influent $\text{NH}_4\text{-N}$ concentrations affected the MBR's efficiency in degrading OSPW NAs when the HRT was 48 h.

Based on the removal efficiency of total classical NAs in Table 2, $\text{NA}_{\sum ne}/\text{NA}_{\sum ni}$ and $\text{NA}_{\sum |z|e}/\text{NA}_{\sum |z|i}$ in Fig. 1, A75H48 showed the lowest effectiveness for OSPW treatment, which may be attributed to the toxicity of free ammonia (FA) caused by high $\text{NH}_4\text{-N}$ concentration. It is known that a higher total ammonia ($\text{NH}_3\text{-N}$ and $\text{NH}_4\text{-N}$) concentration under an alkaline pH environment results in a higher concentration of FA, which has been reported to be inhibitory to nitrifiers and other microbes.^{56,57} Based on Ford *et al.*'s study,⁵⁸ an FA concentration of ~ 13.5 mg L^{-1} is estimated in the MBR at A75H48 stage.

In Fig. 1A, the two lines of A50H48 and A25H48 are weaved over the n range. The difference between the lines for A50H48 and A25H48 is insignificant ($P = 0.3360 > 0.05$). For NAs of C12–17, A25H48 removed more NAs than A50H48 did, whereas the latter achieved higher removal of C9 and C20–23 NA species. Moreover, it is interesting to notice the residual abundance of C23 NAs at A75H48 was lower than that at A25H48. Therefore, it is suggested that higher concentrations of influent $\text{NH}_4\text{-N}$ might be helpful to degradation of NAs with higher carbon numbers (*i.e.*, $n > 21$ in this study). It is noticed that for $|Z|$ of 4, 6, 8 and 10, A50H48 resulted in more ($P = 0.034 < 0.05$) residual NAs than A25H48 (Fig. 1C).

Among the three stages with an HRT of 48 h, A25H48 was the only stage that reduced the O3 oxy-NAs, whereas A75H48 and A50H48 increased the abundance of that group of oxy-NAs by 20% and 4%, respectively (Fig. 4). As for O4 oxy-NAs, A25H48 showed the best degradation (33%). Regarding the removal of O5 and O6 oxy-NAs, A25H48 demonstrated either similar or higher performance than the other two stages.

Therefore, based on the higher removal rates of total classical NAs (33.7%) and total oxy-NAs (20%), the nitrogen composition used at A25H48 (25 mg $\text{NH}_4\text{-N}$ per L and 50 mg $\text{NO}_3\text{-N}$ per L) was selected for further trials on different HRTs.

3.1.2 Effect of HRT. A25H72, A25H48, A25H24 and A25H12 are compared to evaluate the effect of HRT on OSPW NA degradation in the MBR. As indicated in Fig. 1B and D, $\text{NA}_{\sum ne}/$



$NA_{\sum n_i}$ at HRT of 12 h was significantly lower than those at HRT of 48 h ($P = 0.038 < 0.05$) and 24 h ($P = 0.040 < 0.05$). Among the four HRTs examined, the 48 hour HRT showed the poorest removal efficiency of NAs with higher carbon numbers (C19–23), whereas 72 hour HRT and 12 hour HRT demonstrated a similarly good removal rate of C19–23 NAs. With respect to removal of NAs with different Z , the 72 hour HRT returned the worst removal efficiency for NAs with $|Z|$ within the range of 2 to 12. For $|Z| = 2, 4, 14$ and 16 , HRT of 48 h showed higher $NA_{\sum |Z|e}/NA_{\sum |Z|i}$ ratios than the 24 hour and 12 hour HRTs, indicating poorer removal efficiency at 48 hour HRT. For the rest $|Z|$ values, the difference among the HRTs of 48, 24 and 12 h was marginal. Within the whole $|Z|$ range, the difference of $NA_{\sum |Z|e}/NA_{\sum |Z|i}$ ratios between the HRTs of 24 and 12 h was insignificant ($P = 0.671 > 0.05$), indicating that the HRTs of 24 and 12 h made no difference in degrading NAs sharing the same cyclization degree.

No clear correlation between HRT and oxy-NA degradation rate could be elucidated based on the data in this study. Among the four stages, A25H12 exhibited the highest overall removal of oxy-NAs ($O3 + O4 + O5 + O6$) (23.9%). Therefore, an HRT of 12 h is considered as the optimal on the basis of overall classical NA and oxy-NA removal.

Furthermore, it is not unconceivable that NA species with diverse molecular structures may have different bio-persistence, thus requiring different optimal operating conditions. Thousands of different NAs were present in the system, making it intricate to decipher the kinetics underlying. Previous researchers have reported enhancement of contaminant rejection performance by membrane biofouling. It has been suggested that the formation of biofouling may promote the microfiltration (MF) membrane rejection performance up to the nanofiltration (NF) efficiency.⁵⁹ Solutes with higher molecular weights and more complex molecular structures tend to be better rejected by NF membrane.⁶⁰ In addition, it is apparent that the negatively charged deprotonated NA molecules confronted electrostatic repulsion from the membrane fouling layer.⁶⁰ Thus, NAs could be either repelled or adsorbed by the fouling layer, depending on the permeate dragging force, hydrophobic interaction and electrostatic repulsion. In our study, HRT of 12 hours led to the removal of NA with higher hydrophobicity ($n > 21$ and $|Z| < 6$) to a bigger extent (Fig. 1), suggesting the contribution of fouling layer. Microbial community structure analyses indicate that the potential major degraders of NAs were also abundantly present in the membrane's fouling layer (biofilm) (Section 3.2). Thus, the fouling layer formed on membrane surface may enhance the biodegradation of NAs through rejecting NAs thereby elevating the NAs concentration in the MBR sludge and on membrane surface. Organic and biological fouling in the MBR was not necessarily unfavorable in light of the improved NA degradation. It is therefore imperative to contain membrane fouling within a certain level to help mitigation of recalcitrant organic pollutants without significantly increasing the demand of chemical cleaning and membrane replacement. Further study could be performed to look into the influences of membrane/fouling-layer filtration on the removal of NAs in MBR.

Regarding the aquatic toxicity reduction, all six operating conditions effectively reduced the OSPW's inhibition effect to *Vibrio fischeri* (Fig. S5†). Although the best detoxification percentage (37%) was achieved at the A25H12 stage based on the Microtox bioassay, the correlation between detoxification efficiency and NA (or oxy-NA) removal was vague. That is not unexpected as the acute toxicity is not only determined by NAs but also associated with other toxic organic compounds, such as aromatic compounds.

3.2 Microbial community characterization

Out of the great similarity caused by the internal recirculation between the anoxic and aerobic tanks, the OTU numbers of the two compartments were averaged for following analyses. As is shown in Fig. 5A–F, Rhodocyclales, Burkholderiales and Nitrosomonadales of the phyla Proteobacteria, Cytophagales, [Saprosirales] and Flavobacteriales of the phyla Bacteroidetes, Nitrospirales of the phyla Nitrospirae, and RB41 of the phyla Acidobacteria were arguably the most abundant microorganisms in the MBR throughout the whole operation.

It is found that the variations in relative abundances of some microbial species corresponded to the changes in MBR performance over the different operating conditions. The relative abundance of Rhodocyclales of sludge samples over the stages was positively correlated to the MBR's denitrification performance with an r_p of 0.790. More interestingly, Rhodocyclales demonstrated strong positive correlations ($r_p > 0.700$) to removal rates of C12–17 NAs and 3–4 ringed NAs (Tables S4 and S5†). In comparison, Cytophagales showed positive correlations to removal rates of C19–23 NAs ($r_p \geq 0.755$) and 7–8 ringed ($r_p \geq 0.680$) NAs. In addition, relative abundances of Burkholderiales, RB41, and Deinococcales showed moderate positive correlation with removal rate of C23 classical NAs. It is, therefore, suggested that degradation of NAs with different molecular structures were completed by different microbial species either synergistically or independently. To confirm this, further effort should be made by using isolated bacterial cultures and model compounds.

Fig. 5G and H delineate the 10 most abundant bacterial orders of the two samples collected from membrane cake layers. All the major orders were commonly found in both of sludge samples and their corresponding cake layer samples. Some bacteria, such as Rhodocyclales and Cytophagales, were found more in the sludge samples, whereas some others, such as Nitrospirales, were more abundant in the cake layer samples. It is also found that the microbial community of membrane cake layer was affected by the suspended growth community. For instance, stage A25H24 witnessed more Rhodocyclales in both of sludge and cake layer microbial communities than stage A50H48 did. Furthermore, the similar dominant microorganisms in the sludge samples and their corresponding cake layer samples imply that the microorganisms inhabiting membrane cake layer might contribute to degradation of NAs given that biofilm systems have proven their effectiveness on OSPW NA degradation.^{21,45}



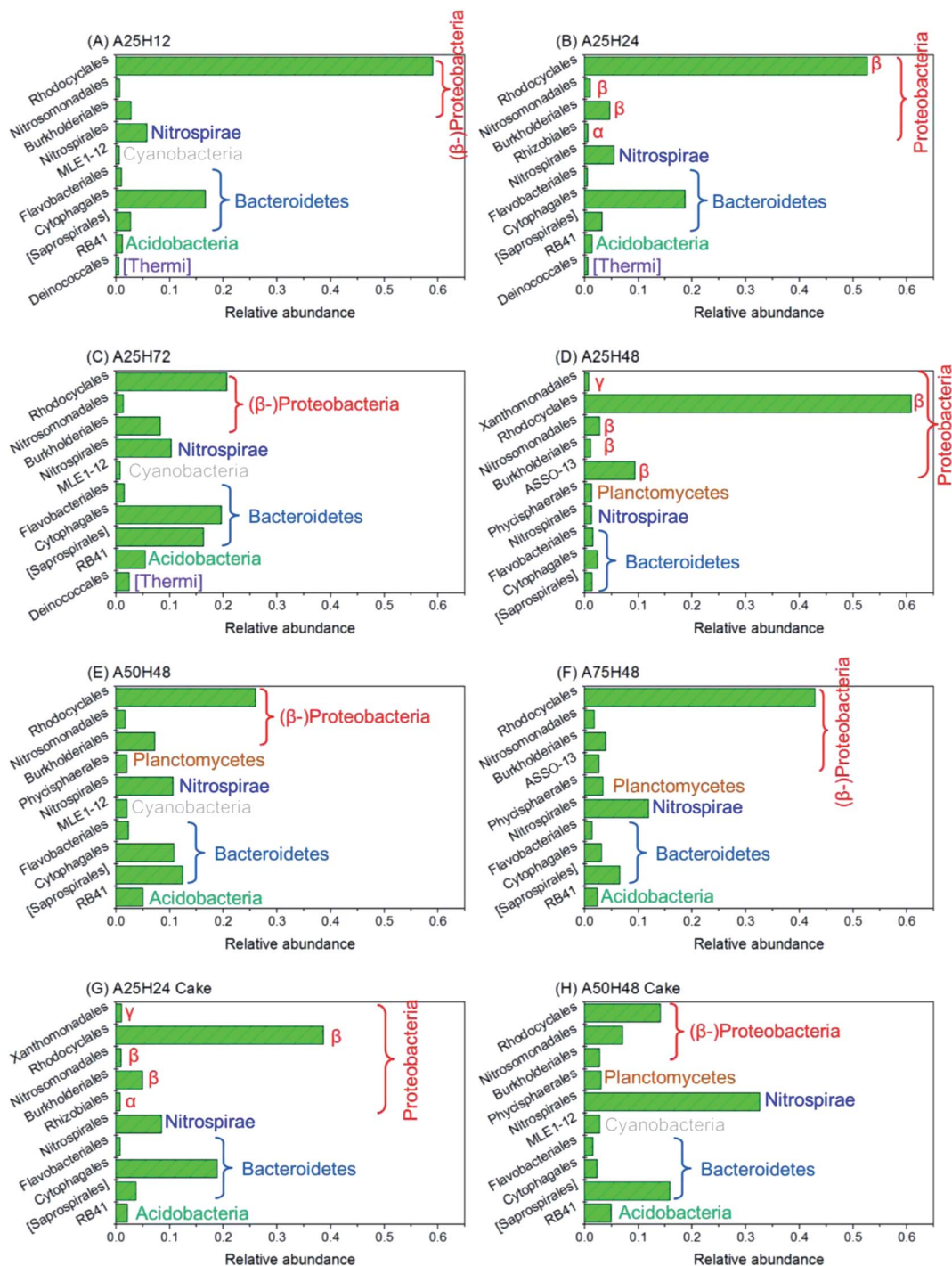


Fig. 5 The 10 most abundant orders detected in sludge samples from all operating stages (A–F) and membrane cake layer samples at stage A25H24 and A50H48 (G and H) (note: square brackets indicate taxonomic changes recommended by Greengenes).

Researchers have reported dominating β -Proteobacteria in oil sands tailings ponds as degrader of NAs and aromatic hydrocarbons,⁶¹ which explains the high abundance of β -Proteobacteria detected in this study. Rhodocyclales are responsible for denitrification and degradation of aromatic hydrocarbons.⁶² Burkholderiales, Cytophagales, Nitrosomonadales and Flavobacteriales have been linked to degradation of bio-recalcitrant hydrocarbons including NAs.^{63–69} The

high population of Nitrosomonadales (ammonium oxidizers) and Nitrospirales (nitrite oxidizers) justifies the excellent nitrification performance of the MBR over the whole study.

Alpha diversity indices, Chao1, observed species, and Shannon were computed through rarefaction (Table S6†). A25H12 had the lowest bacterial richness among the six stages. It is interesting to notice that 5% of the count of genera constituted over 70% of the total bacterial abundance (Table S7†) for every



operating condition, suggesting that the majority of the species richness was originated from numerically minor microbial species. Given that the A25H12 stage demonstrated the best classical and oxidized NA removal rates, it is suggested that numerically minor microbial species might not contribute to the MBR's NA degradation performance though their existence largely affected the species richness of the community. This is in accordance to previous research.⁷⁰

The MBR also demonstrated excellent anti-fouling performance. The first severe fouling was not observed until after 433 days of continuous operation. Within the whole 742 day operation, only six severe fouling events occurred. Detailed results on membrane fouling behavior, major foulants, and fouling layer microbial community structure are available elsewhere.⁷¹

4. Conclusion

After more than one year of continuous operation, the MBR's performance on OSPW NA degradation was successfully improved through changing the supplemented inorganic nitrogen composition and HRT. The study showed that neither higher supplemented $\text{NH}_4\text{-N}$ concentrations nor longer HRTs guaranteed a better NA removal. Instead, an $\text{NH}_4\text{-N}$ concentration of 25 mg L^{-1} and an HRT of 12 h demonstrated the best removal rates of total classical NAs (37.6%) and total oxy-NAs (23.9%). Classical NAs with larger carbon numbers were generally better mitigated in the MBR; and species with higher cyclicality were more persistent to microorganisms. Moreover, MiSeq sequencing analyses revealed that orders of Proteobacteria (*e.g.*, Rhodocyclales and Burkholderiales), Bacteroidetes (*e.g.*, Cytophagales and Flavobacteriales), and Nitrospirae (*i.e.*, Nitrospirales) were the dominating microbial groups over the whole study duration though their relative abundances varied as the operating conditions changed. In the next stage, biofilm process would be added into the reactor to further enhance the removal efficiency. For instance, biodegradation of commercial NA model compounds has been studied based on biochar and granular activated carbon (GAC), which demonstrated quite encouraging performance. It is interesting to investigate whether the integration of bio-adsorption into the MBR would enhance its OSPW NA degradation.^{72,73} Further efforts may also be made using model compounds and isolated bacteria cultures to verify the contribution and specificity of those microbes to degradation of NAs with certain molecular structural features. The findings from this study could be illuminating to future studies and projects on MBRs and other biological processes for not only OSPW treatment but also other impaired waters containing recalcitrant organic contaminants, such as shale gas produced water.

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