Ammonium and organic carbon co-removal under feammox-coupled-with-heterotrophy condition as an efficient approach for nitrogen treatment

Chung Phuong Le1,4, Hai Thi Nguyen2, Toi Duy Nguyen2, Quyen Huynh Minh Nguyen2, Hai The Pham3,4 & Hang Thuy Dinh2*

Nitrification is the rate limiting step in the nitrogen removal processes since nitrifiers have high oxygen demand, but poorly compete with aerobic heterotrophs. In a laboratory-scaled system, we investigated a process of ammonium oxidation under ferric-iron reducing condition (feammox) in the presence of organic carbon using influents with high NH4+ and COD contents, and ferrihydrite as the only electron acceptor. Batch incubations testing influents with different NH4+ and COD concentrations revealed that the [COD]/[NH4+] ratio of 1.4 and the influent redox potential ranging from −20 to +20 mV led to the highest removal efficiencies, i.e. 98.3% for NH4+ and 58.8% for COD. N2 was detected as the only product of NH4+ conversion, whereas NO2− and NO3− were not detected. While operating continuously with influent having a [COD]/[NH4+] ratio of 1.4, the system efficiently removed NH4+ (> 91%) and COD (> 54%) within 6 day retention time. Fluorescence in situ hybridization analyses using Cy3-labeled 16S rRNA oligonucleotide probes revealed that gamma-proteobacteria dominated in the microbial community attaching to the matrix bed of the system. The iron-reduction dependent NH4+ and COD co-removal with a thorough conversion of NH4+ to N2 demonstrated in this study would be a novel approach for nitrogen treatment.

Wastewaters with high ammonia content if discharged inappropriately can cause adverse environmental effects to aquatic systems, i.e. toxicity to living organisms and eutrophication in water bodies1. Ammonium removal from wastewater has been for long time a controversial research topic for the scientific community. Traditionally, NH4+ is removed by the two-stage process of nitrification (aerobic, lithotrophic) and denitrification (anaerobic, organotrophic)3. An alternative technology for NH4+ removal is the partial nitritation/anammox process in which half NH4+ is first oxidized to nitrite by nitrifying bacteria, afterward the other half is oxidized with nitrite to N2 by Planctomycetes species3. Both technologies employ nitrification step which is highly demanding in oxygen and chiefly responsible for the elevated cost of the treatment process. Although the partial nitritation/anammox reduces up to 60% of that cost, technologies with no or lower oxygen demand are still desired.

In the last decade, a novel pathway of NH4+ oxidation coupled to ferric iron (Fe3+) reduction in the absence of oxygen and mediated by microorganisms, termed feammox was proposed4–7. Clement et al.8 and Shrestha et al.9 reported the feammox process in wetland soils, where NH4+ was oxidized to NO2− and considered as a significant mechanism of nitrogen loss in wetland environment8,9. In a laboratory model, Sawayama4 demonstrated the feammox process with NH4+ conversion to NO2−, and thus proposed feammox as an alternative mode of NH4+ removal from wastewater. Using the 16S rRNA gene clone library approach, the major microbial groups in the model were identified as Exiguobacterium spp., Pseudomonas spp. and Carnobacterium maltaromaticum4. None of the delta-proteobacterial iron reducers was detected.

1Nha Trang University, Nguyen Dinh Chieu 2, Nhatrang, Khanhhoa, Vietnam. 2VNU Institute of Microbiology and Biotechnology, E2 Building, Xuan Thuy 144, Caugiai, Hanoi, Vietnam. 3GREENLAB-Center for Life Science Research (CELIKE), VNU University of Science, Nguyen Trai 334, Thanh Xuan, Hanoi, Vietnam. 4Department of Microbiology, Faculty of Biology, VNU University of Science, Nguyen Trai 334, Thanh Xuan, Hanoi, Vietnam. *email: dthangimbt@gmail.com
In other studies, \( \text{NH}_4^+ \) oxidation directly to \( \text{N}_2 \) was also proven for microbial communities in tropical rainforest soil. The first pure culture of feamnox bacterium was strain A6 of the Acidimicrobiaceae group (supposed to be a new taxon) isolated from a low pH ferricyanide-containing enrichment culture. This strain grew under autotrophic and acidic condition (at pH 5) and used ferric iron to oxidize \( \text{NH}_4^+ \) to nitrite. With the ability of oxidizing \( \text{NH}_4^+ \) via the feamnox pathway, strain A6 was proposed for application in a new feamnox-based process of \( \text{NH}_4^+ \) removal from wastewater.

In the feamnox process, \( \text{NH}_4^+ \) can be oxidized to \( \text{NO}_3^- \), \( \text{NO}_2^- \) or \( \text{N}_2 \), depending on the environmental pH, and thus the amount of energy retrieved by microorganisms can be respectively at different levels:

\[
3\text{Fe(OH)}_3 + 5\text{H}^+ + \text{NH}_4^+ \rightarrow 3\text{Fe}^{2+} + 9\text{H}_2\text{O} + 0.5\text{N}_2 (−245 \text{ kJ mol}^{-1}) \quad (1)
\]
\[
6\text{Fe(OH)}_3 + 10\text{H}^+ + \text{NH}_4^+ \rightarrow 6\text{Fe}^{2+} + 16\text{H}_2\text{O} + \text{NO}_2^- (−164 \text{ kJ mol}^{-1}) \quad (2)
\]
\[
6\text{Fe(OH)}_3 + 8\text{H}^+ + \text{NH}_4^+ \rightarrow 6\text{Fe}^{2+} + 15\text{H}_2\text{O} + \text{NO}_3^- (−207 \text{ kJ mol}^{-1}) \quad (3)
\]

The conversion of \( \text{NH}_4^+ \) directly to \( \text{N}_2 \) (Eq. 1) is energetically most favorable and likely to occur in a wide range of environmental pH conditions, whereas \( \text{NH}_4^+ \) conversion to \( \text{NO}_3^- \) or \( \text{NO}_2^- \) (Eqs. 2 and 3) are more likely to occur in acidic environments (pH < 6.5).

Generally, in the practice of wastewater treatment, COD if presented at high concentrations is needed to be removed prior to \( \text{NH}_4^+ \) removal step (via anaerobic technologies such as biogas reactor, upflow anaerobic sludge blanket—UASB), because nitrifying microorganisms are litho-autotrophic, highly require oxygen for the ammonium oxidation, although they poorly compete with heterotrophic species for oxygen in environments where organic carbon sources are available.

In this paper, we report a process of \( \text{NH}_4^+ \) oxidation in the presence of organic carbon in a laboratory-scaled anaerobic system with ferricyanide as the only electron acceptor (heterotrophic feamnox). The system could achieve a thorough conversion of \( \text{NH}_4^+ \) to \( \text{N}_2 \) while being operated under neutral pH conditions, i.e. close to the conditions of most \( \text{NH}_4^+ \) and COD bearing wastewater sources in daily life. The co-removal of \( \text{NH}_4^+ \) and COD coupled with iron reduction demonstrated in this study provides an alternative nitrogen treatment technology that does not require oxygen, thereby eliminating the most critical disadvantage of the presently applied technologies based on nitrification/denitrification and partial nitritation/anammox principles.

Materials and methods

**Laboratory-scaled treatment system design.** The laboratory-scaled treatment system in this study was designed following the working principle of a wastewater treatment plant (Fig. S1) in Cu Chi (HCM City, Vietnam) that operates in the presence of Fe-containing anaerobic sludge. The laboratory system consisted of three connected acrylic tanks of 20 cm × 12 cm × 40 cm (length × width × height) size. The working volume of each tank was 6 L, and the total working volume of the system was 18 L. Liquid medium as the influent was supplied by a peristaltic pump, and subsequently transferred from one tank to the next by overflow through bottom-oriented connecting tubes to ensure good contact of the influent with the microbial community at the bottom of each tank. A matrix bed consisting of 100 cm³ of HDPE MBBR matrix 12 × 7 mm (specific surface 850 m²/m³, Kaite Chemical, PRC) was loaded onto every tank. Each tank has a gas sampling valve positioned on the lid, respectively, while the full water level was 27 cm (Fig. 1).

The mineral medium that was used as the influent was adapted from Ratering and contained the following components: 1.0 g NaCl L⁻¹, 0.4 g MgCl₂·6H₂O L⁻¹, 0.15 CaCl₂·2H₂O L⁻¹, 0.5 KCl L⁻¹, 0.2 KH₂PO₄ L⁻¹ (pH 7 ± 0.2). Ammonium and COD were loaded at different concentrations depending on the experimental conditions by using solutions of 1.0 mM \( \text{NH}_4\text{Cl} \) and 1.0 M sodium acetate, respectively. The influent was then supplemented with 1 mL L⁻¹ vitamin and microelement solutions. Afterward, the buffering bicarbonate and reducing agent sodium ascorbate solutions were added to the final concentrations of 30 mM and 1 mM, respectively. The slurry of the electron acceptor ferricyanide was freshly prepared according to Ratering and added to the medium for a final concentration of 30 mM. Finally, the synthetic wastewater was flushed with argon to remove dissolved oxygen.

**Enrichment of iron-reducing community in the laboratory-scaled system.** Enrichment of iron-reducing microorganisms in the laboratory-scaled system was carried out by circulating the anoxic influent inside the system (from the first tank to the third tank and back) with a peristaltic pump (Huiyu Weiye (Beijing), PRC) at 2 mL min⁻¹. The inoculum was a Fe-containing sludge from the wastewater treatment plant in Cu Chi, HCM City Vietnam. Accordingly, 1.5 kg seed sludge was resuspended in 3 L anoxic distilled water in a tightly closed glass bottle, shaken at 100 rpm for 5 min, then added to each tank in a ratio of 10% of the working volume. The tanks were flushed with argon to remove oxygen and then sealed with gas-tight lids. The enrichment process was carried out at ambient temperatures (30 ± 2 °C). Gas samples from the head space of each tank were collected every 24 h for \( \text{N}_2 \) analyses. In parallel, water samples from each tank were collected every 24 h for analyses of the relevant parameters, including \( \text{NH}_4^+ \), COD, and \( \text{Fe}^{2+} \). A negative control was established in a separate tank containing mineral medium with compositions similar to those of the medium in the enrichment tanks, except that the organic carbon source (as sodium acetate) was omitted.

**Batch incubation experiment.** After enrichment, the influent of the system was replaced with a freshly prepared anoxic medium containing 200 mg L⁻¹ \( \text{NH}_4^+ \) and COD (as sodium acetate). In the batch incubation
experiment, tanks 1, 2 and 3 were operated separately with different concentrations of COD, which resulted in the \([\text{COD}] / [\text{NH}_4^+]\) ratios of 1.4, 1.1 and 0.7 in the influents respectively. Within 30 days of the experiment, water samples from the tanks were collected every 24 h to monitor changes in the COD and inorganic nitrogen species (\(\text{NH}_4^+, \text{NO}_2^-, \text{NO}_3^-\) and \(\text{N}_2\)), as well as the redox potential (RP). The \([\text{COD}] / [\text{NH}_4^+]\) ratio giving the best \(\text{NH}_4^+\) removal rate was chosen for the experiment conducted under a continuous operating mode.

**Experiment of co-removal of \(\text{NH}_4^+\) and COD in the continuous operating mode.** Co-removal of \(\text{NH}_4^+\) and COD was demonstrated in the laboratory-scaled system operated under continuous mode using an anoxic influent containing 50 mg L\(^{-1}\) \(\text{NH}_4^+\) and COD (as loaded sodium acetate) at the concentration that enabled the best \(\text{NH}_4^+\) removal rate. The influent was pumped into the system at a hydraulic loading rate of 2 mL min\(^{-1}\) (3 L day\(^{-1}\)). Accordingly, the retention times of the influents in the three sequential tanks 1, 2, and 3 were 2, 4 and 6 days, respectively. Water samples were collected every 24 h from each tank to quantify \(\text{NH}_4^+\) and COD concentrations.

**Fluorescence in situ hybridization.** Precipitates scratched from HDPE MBBR matrix surfaces were fixed for 12 h at 4 °C in 4% formaldehyde, washed twice with PBS (sodium phosphate 10 mM, pH 7; NaCl 130 mM), and stored in PBS:ethanol (1:1) solution at –20 °C. Prior to hybridization, the fixed precipitates were collected on polycarbonate filters (0.2 μm pores, Millipore). The filters were then cut into small sections, hybridized with Cy3-labelled GAM42a and Delta385 16S rRNA probes specific for γ- and δ-proteobacteria, respectively at appropriate stringencies (formamide concentrations)\(^{15}\). DAPI (4′, 6-diamidino-2-phenylindole) was used for general cell staining. The hybridized samples were then observed by a fluorescence microscope (Carl Zeiss, Germany) and the fluorescence signals were captured by an AxioCam Icc3 camera. The images were then analyzed by Image J software.

**Analytical methods.** The ammonium concentration was determined photometrically by using sodium nitroprusside reagent\(^{16}\). The calibration curve of the \(\text{NH}_4^+\) concentration was obtained from a serial dilution of the \(\text{NH}_4\text{Cl}\) solution (using previously dried \(\text{NH}_4\text{Cl}\) crystals) in the range of 0.018–1.8 mg L\(^{-1}\).

The ferrous iron concentration was determined photometrically by using O-phenanthrolin reagent\(^{17}\). A calibration curve was obtained from a serial dilution of the \(\text{FeSO}_4\) solution in the range of 0.28–2.8 mg L\(^{-1}\).

The nitrate concentration was determined by using NitraVer® 5 Nitrate Reagent Powder Pillows (Hach Instruments Inc., USA) with a calibration curve obtained from a serial dilution of the \(\text{NaNO}_3\) solution in the range of 0–30 mg L\(^{-1}\).
The nitrate concentration was determined by using NitriVer+3 Nitrite Reagent Powder Pillows (Hach Instruments Inc., USA) with a calibration curve obtained from a serial dilution of the NaNO3 solution in the range of 0–150 mg L−1.

Quantification of N2 was conducted by gas chromatography (GC) on 7890A instrument (Agilent) using the column HT-plot/Q with a TCD detector. The GC operating parameters were as follows: oven temperature 60 °C; detector temperature 250 °C; argon carrier gas (Messer, Vietnam) at a flow rate of 3.0 mL min−1; and 0.2 mL sample injection volume. An N2 calibration curve was constructed by using a mixture of N2 in argon in the range 0–100% volume prepared in serum bottles tightly closed with rubber stoppers. The concentration of N2 was calculated as % in samples based on the width of the peak values.

COD determination was conducted according to USEPA 410.4 method by using COD digestion vials with 1000 mg L−1 COD standard solution (Hach Instruments Inc., USA).

The redox potential (RP) of the medium in the treatment system was measured by using a working graphite electrode in close proximity with an Ag/AgCl reference electrode (BASI, West Lafayette, IN 47906, USA), with a digital multimeter (model 1009, Kyoritsu, Japan). Both electrodes were built on a rubber stopper of a 100 mL Schott flask. Thus, 80 mL liquid sample was taken from the treatment system into the 100 mL Schott flask, and then the flask was immediately sealed with the rubber stopper that carried the working and reference electrodes. The RP was subsequently measured by the digital multimeter using the voltammetry mode (Fig. S3).

Data analysis. All measurements were carried out in triplicate, unless otherwise stated. The data collected during the experiments were processed using Microsoft Excel software (for average and standard deviation functions, the standard deviation was calculated using the ‘n-1’ method) and graphed by using SigmaPlot 14 software.

The C and N balance calculations were carried out with experimental data retrieved from the batch incubation experiment in tank 1 where the best NH4+ removal was observed (details are presented in supplementary material).

Results
Enrichment of iron-reducing community. The enrichment was carried out in the laboratory-scaled treatment system with Fe-containing anaerobic sludge from a wastewater treatment plant as the initial seed. It can be seen from the results Fig. 2 that the enrichment process harbored a microbial community capable of oxidizing NH4+ and reducing Fe3+, as the NH4+ concentration decreased while Fe2+ was released into the medium. In all three tanks NH4+ oxidation started from the 3rd day of incubation and kept occurring at a relatively stable rate toward the end of enrichment. After 30 days of incubation, NH4+ was almost completely removed, from 200 mg L−1 to < 10 mg L−1 in all three tanks (Fig. 2a). In the negative control without organic carbon, NH4+ conversion was much lower, with a remained concentration as high as 155 mg L−1 after 30 days.

In accordance with NH4+ conversion, Fe3+ was detected as the product of Fe3+ reduction (Fig. 2b), the concentration was up to 11.01 mg L−1 in the medium. It could be observed that the dissolved Fe2+ detected in the enrichment medium was not in a cumulative manner throughout the enrichment, at certain time points its concentration was even critically low (Fig. 2b). During the enrichment process, the pH in the system was relatively stable in the range of 6.8–7.2. Distinct NH4+ oxidation in the enrichment experiment compared to that of the control (Fig. 2a) indicated that organic carbon is important for the Fe3+-dependent NH4+ oxidation in this system. Indeed, ~50% of COD was consumed after 30 days of the enrichment (Table 1), implying that the feammox process was coupled with heterotrophy.

Regarding the products of NH4+ oxidation, neither NO2− nor NO3− was detected in the enrichment medium. Instead, N2 accumulated in the head spaces of all three tanks, with the highest concentration reaching 302.75 mg L−1. Nitrate and nitrite were not detectable in all three tanks during the batch incubation experiment. Instead, N2 accumulated in the head spaces of all three tanks, with the only product of NH4+ conversion (Fig. 3c). Gas production in all three tanks could be observed via the formation of gas bubbles, which to different extents depended on the [COD]/[NH4+] ratio (Fig. S3). N2 production in tank 1 was higher than in tank 2 and tank 3, which reflects the higher NH4+ oxidation that occurred in these tanks (Fig. 3a).

The effect of COD on NH4+ removal. After the enrichment process had succeeded in establishing an active “feammox” community in the treatment system (Fig. 2a), batch incubation experiments were carried out independently in the three tanks loaded with influents of different NH4+ and COD concentrations. In detail, the influents contained 200 mg L−1 NH4+ and COD (as sodium acetate) at different concentrations according to the [COD]/[NH4+] ratios of 1.4; 1.1 and 0.7 in tanks 1, 2, and 3, respectively.

The results showed that the NH4+ and COD removal efficiencies to a large extent depended on the ratio of these two chemical species in the influent (Table 2). Thus, after 30 days of incubation the highest NH4+ removal efficiency (98.3%) was achieved in the first tank where the [COD]/[NH4+] ratio was 1.4. That removal was significantly higher than in tanks 2 and 3 (83.3% to 85.5%) where the [COD]/[NH4+] ratios were lower, 1.1 and 0.7, respectively.

Regarding COD removal, approximately 58% of the COD loaded was removed after 30 days in tank 1 and tank 3, where the [COD]/[NH4+] ratios were 1.4 and 0.7 respectively. Slightly higher COD removal efficiency was observed in tank 2, where the [COD]/[NH4+] ratio was 1.1 (Table 1, Fig. 3b). Thus, unlike the NH4+ removal, the COD removal was not clearly affected by the [COD]/[NH4+] ratio in the influent. Surprisingly, the COD removal varied insignificantly from 58.8 to 63.7%, despite significant variation in the [COD]/[NH4+] ratio, from 0.7 to 1.4.

Nitrate and nitrite were not detectable in all three tanks during the batch incubation experiment. Instead, N2 accumulated in the head space of each tank as the only product of NH4+ conversion (Fig. 3c). Gas production in all the three tanks could be observed via the formation of gas bubbles, which to different extents depended on the [COD]/[NH4+] ratio (Fig. S3). N2 production in tank 1 was higher than in tank 2 and tank 3, which reflects the higher NH4+ oxidation that occurred in these tanks (Fig. 3a).
The redox potential (RP) of the medium during the batch incubation experiment was also monitored to determine its correlation to the “feammox” process (Fig. 3d). In tank 1 where [COD]/[NH$_4^+$] ratio was 1.4, the RP was most stable, varying only in the range of −20 to +20 mV throughout the experiment. In contrast, in tanks 2 and 3 as well as in the control without supplemented organic carbon, the RP fluctuated to a large extent.

**Figure 2.** Enrichment of the feammox community in the laboratory-scaled system. The NH$_4^+$ oxidation (a) coupled with Fe$^{2+}$ production (b) leading to N$_2$ formation as product of NH$_4^+$ conversion (c) during the enrichment. The mineral medium used for enrichment contained 200 mg L$^{-1}$ NH$_4^+$ and 250 mg L$^{-1}$ COD (as sodium acetate). The inoculum was anoxic sludge from the wastewater treatment station in Cu Chi (HCM city, Vietnam). The control was a separate tank loaded with the same medium and inoculum, except for the COD content (organic carbon).
Table 1. NH$_4^+$ and COD removal in the enrichment and the control without organic carbon (data were average of measurements in three tanks of the system).

| Time (day) | Enrichment | Control |
|------------|------------|---------|
|            | NH$_4^+$ (mg L$^{-1}$) | COD (mg L$^{-1}$) | NH$_4^+$ (mg L$^{-1}$) | COD (mg L$^{-1}$) |
| 0          | 201.23 ± 2.67 | 250.21 ± 7.23 | 199.56 ± 7.02 | 0.00 ± 0.00 |
| 10         | 146.28 ± 7.87 | 228.98 ± 7.16 | 184.71 ± 3.23 | 0.00 ± 0.00 |
| 20         | 51.03 ± 3.35 | 175.85 ± 7.98 | 170.93 ± 1.8 | 0.00 ± 0.00 |
| 30         | 9.91 ± 0.86 | 128.30 ± 3.75 | 155.54 ± 3.31 | 0.00 ± 0.00 |

Table 2. Effect of COD on the NH$_4^+$ removal in batch incubation experiment after 30 days.

| Initial [COD]/[NH$_4^+$] ratio | NH$_4^+$ removed (%) | COD removed |
|-------------------------------|----------------------|-------------|
| 1.4                           | 98.3                 | 58.8        |
| 1.1                           | 83.3                 | 63.7        |
| 0.7                           | 85.5                 | 57.9        |

Figure 3. Batch incubation experiments. The removal of NH$_4^+$ (a) and COD (b); formation of N$_2$ as the product of NH$_4^+$ conversion (c), and the redox potential (RP) of the medium (d) during the batch incubation with an influent containing 200 mg L$^{-1}$ NH$_4^+$ and organic carbon (as sodium acetate) supplied in different [COD]/[NH$_4^+$] ratios: Tank 1, 1.4; Tank 2, 1.1; Tank 3, 0.7. The control was a separate tank loaded with the same influent and inoculum, except for the COD content (organic carbon).
mostly in the positive ranges of +20 mV and above. Obviously, the RP condition in tank 1 was the most suitable for the reduction of ferric iron.

Thus, via the batch incubation experiment we showed that for an influent with NH$_4^+$ concentration as high as 200 mg L$^{-1}$ and [COD]/[NH$_4^+$] ratio of 1.4, the NH$_4^+$ removal efficiency was impressively high, reaching 98.3%. Simultaneously, 58.8% COD was removed. This process occurred under the conditions of relatively stabilized neutral pH levels (6.8–7.2) and redox potentials (~20 ± 20 mV).

Calculations for the nitrogen balance (Table S4) showed that throughout the incubation experiment, NH$_4^+$ was thoroughly converted to N$_2$ according to the stoichiometric ratio 1:1.5 (reaction 1). Acetate consumptions at days 5, 10 and 15 were in accordance with the theoretical ratio that could be used for NO$_3^-$ or Fe$^{3+}$ reduction (reactions 4 and 5), taking into account that NO$_3^-$ might have been generated as product of feammox reactions 2 and 3.

\[
\text{NO}_3^- + 8/5\text{CH}_3\text{COO}^- + 13/8\text{H}^+ \rightarrow 0.5\text{N}_2 + 5/4\text{CO}_2 + 7/4\text{H}_2\text{O} \quad (-501 \text{ kJ mol}^{-1})
\] (4)

\[
\text{Fe}^{3+} + 5/8\text{CH}_3\text{COO}^- + 1/2\text{H}_2\text{O} \rightarrow \text{Fe}^{2+} + 1/4\text{HCO}_3^- + 9/8\text{H}^+ \quad (-814 \text{ kJ mol}^{-1})
\] (5)

However, this trend was not observed at days 20, 25, and 30 later, as the amounts of oxidized acetate were lower than the theoretical values, i.e. 56% at day 20; 45.8% at day 25, and 42.5% at day 30 (Table S4). The oxidation of acetate to CH$_4$ was not observed (Fig. S2B).

**Co-removal of NH$_4^+$ and COD in a continuous operation mode.** Continuous operation mode in the laboratory system was carried out to assess the efficiencies of NH$_4^+$ and COD removals under the most suitable conditions, as determined in the batch experiment above. In this experiment, an influent with the moderate NH$_4^+$ concentration of 50 mg L$^{-1}$ and [COD]/[NH$_4^+$] ratio of 1.4 was applied at the hydraulic load of 2 mL min$^{-1}$, As a result, NH$_4^+$ was removed constantly at different levels depending on the retention time in each tank of the system (Fig. 4a). The NH$_4^+$ removal required only a short time for adaptation, readily reaching a steady state after 5 days of continuous operation, with the removal efficiency maintained at around 30%, 60% and > 91% in tank 1, tank 2, and tank 3, respectively. The most efficient NH$_4^+$ removal was achieved at 6 day retention time in tank 3, with a stabilized efficiency of > 91% (Tab. S1). As a result, an NH$_4^+$ concentration as low as 0–4 mg L$^{-1}$ was maintained in the effluent at a retention time of 6 days (Fig. 4a, Tab. S1).

Regarding COD removal, a steady removal rate was achieved after 5 days of operation, which was consistent with the NH$_4^+$ removal (Fig. 4b). The highest level of COD removal was observed in the third tank, i.e. with a 6 day retention time and efficiency in the range of 54–66% throughout 30 days of continuous operation (Fig. 4b, Tab. S2). Accordingly, the COD concentration in the effluent was in the range of 23–30 mg L$^{-1}$ (Tab. S2) throughout the experiment.

Fluorescence in situ hybridization using two probes specific for the δ- and γ-proteobacteria showed that the number of DAPI-stained cells hybridized with the δ-proteobacteria specific probe was quite low (Fig. 5a,b). In contrast, a very high number of cells gave signals with the γ-proteobacteria specific probe (Fig. 5c,d), which accounted for more than 90% of the total DAPI cell count (Tab. S3).

**Discussion**

The NH$_4^+$ oxidation coupled with ferric iron reduction called feammox was first proposed as an important step of the nitrogen cycle in saturated sediment$^9$. Since then, the feammox process has been proven to occur in different habitats where NH$_4^+$ can be oxidized to N$_2$, NO$_3^-$ or NO$_2^-$, depending on the pH levels in the environments$^6$. Regarding its application in NH$_4^+$ removal from wastewater, feammox seems to be a promising new treatment technology, that is alternative to conventional nitrification/denitrification and partial nitritation/anammox processes.

The first demonstration of NH$_4^+$ removal by feammox process as a possible wastewater treatment technology was carried out in a pH and oxygen controlled laboratory fixed bed reactor$^4$. The reactor was supplied with NH$_4^+$ and Fe(III)EDTA, operated in bicarbonate buffered environment with neutral to slightly alkaline pH$^4$. Under such conditions, NH$_4^+$ was oxidized to NO$_2^-$, thereby raising the pH from 6.7 to 7.8 (due to intensive consumption of protons, according to reaction 2). No organic carbon was added, instead, feammox microorganisms in the reactor produced organic compounds to some extent. More recently, Huang and Jaffé described the enrichment of feammox microorganisms in a batch incubation experiment carried out under acidic conditions (pH 5.5) without added organic carbon$^{10}$. These authors reported a significant NH$_4^+$ oxidation to NO$_2^-$ when ferrihydrite was used as the only electron acceptor. Both studies showed the feammox process in lithotrophic environments with NH$_4^+$ as the only electron donor for ferric iron reduction, which led to the production of NO$_2^-$. Thus, to thoroughly convert NH$_4^+$ from the wastewater to dinitrogen gas in these studies, two more steps would be needed, i.e. (i) nitrite oxidation to nitrate, which is lithotrophic and requires oxygen and (ii) nitrate reduction to N$_2$, which is heterotrophic and does not require oxygen.

In this study we showed that at a neutral pH and in the presence of organic carbon, feammox was likely to occur with a thorough NH$_4^+$ conversion to N$_2$ (reaction 1). The role of organic carbon in the feammox process was evident via a significantly distinct conversion of NH$_4^+$ to N$_2$ in the experimental tanks in the presence of organic carbon in comparison with that in the control without organic carbon (Fig. 2a,c). This conversion was in good agreement with the stoichiometric ratio of NH$_4^+$ to N$_2$ according to reaction 1. Consumption of organic carbon was evident in all experiments, i.e. enrichment (Table 1), batch incubation (Fig. 3b) and continuous operation (Fig. 4b), indicating that the feammox process was coupled with heterotrophy. Carbon balance calculations (Table S4) also confirmed that acetate was likely to be utilized as the carbon source in the feammox process. On the other hand, Fe$^{3+}$ reduction using acetate as the electron donor was also feasible under the experimental
conditions. It is theorized that Fe³⁺ reduction with acetate might have been dominant in the system throughout the first two weeks, but was soon replaced by the feammox to N₂ reaction, which required less acetate.

In the context of wastewater treatment, anaerobic processes are considered to be more advantageous as energy is saved. In the anaerobic world, ferric iron Fe³⁺ is a special electron acceptor that differs from others by two characteristics, i.e. (i) being water-insoluble and (ii) continuity of the two iron species Fe³⁺ ⇌ Fe²⁺, whereas the products of nitrate reduction N₂, sulfate reduction SH⁻ or methanogenesis CH₄ would escape from the liquid.

Thus, in practice of wastewater treatment, if ferric iron is used as the electron acceptor, it would not need to be added regularly. In daily life, reactive iron-rich sources are relatively abundant, e.g. iron rich sludge in drinking water treatment plants or red mud from bauxite tailing¹⁸. From the viewpoint of achieving an eco-friendly approach, utilizing iron-rich wastes is of special consideration.

In daily life, wastewaters with high NH₄⁺ and COD contents such as discharge from seafood processing plants, poultry farms, or biogas effluents, are very common. Such kinds of wastewater sources are considered to be highly contaminated and must be subjected to treatments to remove NH₄⁺, COD and other contaminants before discharging to the surrounding environment. Thus, in the present study, we attempted to establish an efficient treatment system based on the feammox principle to remove NH₄⁺ in heterotrophic conditions, i.e. in the presence of organic carbon (feammox-coupled-with-heterotrophy condition), which is common to many types of wastewater sources in daily life. The heterotrophic feammox community in this treatment system was enriched from Fe-containing sludge collected from an anaerobic wastewater treatment plant in southern Vietnam. The original treatment plant had been working under a feammox-resembling condition, i.e. with a high NH₄⁺ concentration (~ 300 mg L⁻¹) and excessive Fe³⁺ content in the form of red mud (bauxite residue), which raised the pH in the system to 7.5 ± 0.2. In addition to the high NH₄⁺ concentration, the influent also had a high COD concentration of ~ 2500 mg L⁻¹. The working conditions of our laboratory-scaled system followed the original treatment system (Fig. S1) from where the seed sludge was obtained. It should be noted that the working conditions we applied to our laboratory system were far different from those in Sawayama’s⁴ and Huang’s¹⁰ studies (discussed above) in several aspects, i.e. occurrence of heterotrophy, the high load of NH₄⁺ and neutral pH in the influent. Under such conditions, parallel removals of NH₄⁺ and organic carbon coupled with biological

---

**Figure 4.** Performance of the laboratory-scaled feammox system when operated in continuous mode. The removal of NH₄⁺ (a) and COD (b) under the feammox-coupled-with-heterotrophy condition in the system with the influent containing 50 mg L⁻¹ NH₄⁺ and [COD]/[NH₄⁺] = 1.4. The hydraulic loading rate was 2 mL min⁻¹ and the retention times in the three tanks were 2, 4 and 6 days, respectively.
Fe³⁺ reduction was evident. It was of special interest that N₂ (neither NO₂⁻ nor NO₃⁻) was detected as the only product of NH₄⁺ conversion in the system. Still, it would be early to conclude that the conversion of NH₄⁺ to N₂ was due to the feammox process alone, since in such a complex system, other bacterial groups of nitrogen cycling might also exist.

In this study, based on the Fe²⁺ concentration detected in the medium, it is evident that it did not accumulate in the theoretically expected manner (Fig. 2a,b; Fig. S4). This might be due to the complexity of the system, in which the produced Fe²⁺ could be involved in different chemical transformations. The disappearance of Fe²⁺ when produced in the enrichment environment can be explained by two processes: (i) precipitating with carbonate as white deposition in the enrichment tanks, and/or (ii) reoxidizing to Fe³⁺ by reacting with trace oxygen in the medium or (more likely) via biological oxidation coupled with nitrate reduction⁸. Indeed, the RP in the medium during the batch incubation experiment ranged from −20 mV to +20 mV, which is feasible for both Fe³⁺- and NO₃⁻-reduction processes. Thus, the collected data of the Fe²⁺ time course could not be used as evidence to support the Fe³⁺ reduction process in the system.

It should be noted that the microbial community in the treatment system presented in this study did work with a very high influent NH₄⁺ concentration of 200 mg L⁻¹. Nevertheless, NH₄⁺ was almost completely removed after 30 days of incubation (Fig. 3a). The wastewater treatment practice showed that such a high NH₄⁺ content could not be easily removed by the conventional nitrification/denitrification process, whereas a complex operating condition and energy would be required for the partial nitritation/anammox process¹⁹,²⁰. According to Huang and Jaffé, the NH₄⁺ removal efficiency of 9.8% was achieved in a feammox membrane reactor after 6 days of operation, and efficiency of 64.5% was recorded for longer period of 150 days. Similarly, a low NH₄⁺ removal of 20% after 50 days was observed in the feammox reactor established by Sawayaama⁴. Such a low rate of NH₄⁺ removal was also observed in the control without added organic carbon in this study (Figs. 2a, 3a). It was surprising that the presence of COD significantly enhanced NH₄⁺ removal, as shown by our results. However, the amount of COD needs to be controlled to achieve a certain ratio of [COD] to [NH₄⁺] that can enable the most efficient NH₄⁺ removal. Thus, by setting the [COD]/[NH₄⁺] at the ratio of 1.4, almost all NH₄⁺ was removed from such a high concentration of 200 mg L⁻¹ after 30 days of batch incubation. Similarly, under a continuous operation mode, the same [COD]/[NH₄⁺] ratio was applied to a moderately contaminated influent (containing 50 mg L⁻¹ NH₄⁺), which succeeded in more than 91% NH₄⁺ removal with a retention time of only 6 days. At the same time, COD was efficiently removed, which resulted in an effluent with low contents of NH₄⁺ (Tables S1 and S2).

Studies on species composition and key players involved in the NH₄⁺ removal process under iron reduction conditions in bioreactor systems so far have led to equivocal results. In the study of the feammox process under neutrophilic autotrophic condition using Fe(III)-EDTA as the electron acceptor for the ammonium oxidation, Sawayaama⁴ reported a bacterial community dominated by Exiguobacterium spp., Pseudomonas spp. and Carnobacterium sp. (with 22.5%, 17.5%, and 7.5% of the 16S rRNA gene clone library, respectively)⁴. Being previously

Figure 5. Images of cells scratched from the HDPE MBBR matrix surfaces with signals from the DAPI stain (a,c) and of the same section with signals from hybridized Cy3-labeled Delta385 and GAM42a 16S rRNA oligonucleotide probes specific for δ-proteobacteria (b) and γ-proteobacteria (d), respectively. Bar, 10 μm.
reported as arsenate reducers, bacteria belonging to the Exiguobacterium group were believed to be candidates for the feammox process\(^1\,\,^2\). In the bioreactor operated under acidic autotrophic feammox conditions, a complex microbial community with the key player Acidimicrobiaceae bacterium A6 was identified\(^1\,\,^2\). It could be deduced that under heterotrophic feammox conditions such as in this study, the established microbial community would be quite different from that under the auto-lithotrophic conditions demonstrated by Sawayama\(^4\) or Huang and Jaffé\(^10\). The absence of δ-proteobacteria in the community established in this study is likely to occur intensively if organic carbon was present. The highly efficient NH\(_4^+\) removal process based on the feammox-coupled-with-heterotrophy principle presented in this study would have substantial application potential as a novel technology for wastewater treatment. Evidently, applications using this principle would drive thorough NH\(_4^+\) removal from wastewater while producing \(\text{N}_2\) as the end-product in a single-step treatment. Occurring at a neutral pH, the feammox to \(\text{N}_2\) coupled with heterotrophy as presented in this study would be applicable for a wide range of wastewater sources.

Received: 28 July 2020; Accepted: 11 December 2020

Published online: 12 January 2021

References

1. Abdel-Raouf, N., Al-Homaidan, A. A. & Ibraheem, I. B. M. Microalgae and wastewater treatment. *Saudi J. Biol. Sci.* 19, 257–275 (2012).
2. Bitton, G. *Wastewater Microbiology* (Wiley, New York, 2005).
3. Mulder, A., van-de-Graaf, A. A., Robertson, L. A. & Kuenen, J. G. Anaerobic ammonium oxidation discovery in a denitrifying fluidized-bed reactor. *FEBS Microbiol. Ecol.* 16, 177–183 (1995).
4. Sawayama, S. Possibility of anoxic ferric ammonium oxidation. *J. Bioosci. Bioeng.* 101, 70–72 (2006).
5. Shrestha, J., Rich, J. J., Ehrenfeld, J. G. & Jaffé, P. R. Oxidation of ammonium to nitrite under iron-reducing conditions in wetland soils. *Soil Sci.* 174, 156–164 (2009).
6. Yang, W. H., Weber, K. A. & Silver, W. L. Nitrogen loss from soil through anaerobic ammonium oxidation coupled to iron reduction. *Nat. Geosci.* 5, 538–541 (2012).
7. Ding, L. J., An, X. L., Li, S., Zhang, G. L. & Zhu, Y. G. Nitrogen loss through anaerobic ammonium oxidation coupled to iron reduction from paddy soils in a chronosequence. *Environ. Sci. Technol.* 48, 10641–10647 (2014).
8. Clement, J. C., Shrestha, J., Ehrenfeld, J. G. & Jaffé, P. R. Ammonium oxidation coupled to dissimilatory reduction of iron under anaerobic conditions in wetland soils. *Soil Biol. Biochem.* 37, 2323–2328 (2005).
9. Huang, S. & Jaffé, P. R. Characterization of incubation experiments and development of an enrichment culture capable of ammonium oxidation under iron-reducing conditions. *Biogeosciences* 12, 769–779 (2015).
10. Luther, G. W., Sundby, B., Lewis, B. B., Brendel, P. J. & Silverberg, N. Interactions of manganese with the nitrogen cycle: Alternative pathways to dinitrogen. *Geochim. Cosmochim. Acta* 61, 4043–4052 (1997).
11. Schmidt, I. et al. New concepts of microbial treatment processes for the nitrogen removal in wastewater. *FEBS Microbiol. Rev.* 27, 481–492 (2003).
12. Ratering, S. Iron Cycle in Italian Rice Field Soil: Localization of the Redox Processes and Characterization of the Involved Microorganisms (University of Marburg, Marburg, 1999).
13. Widdel, F. & Bak, F. Gram-negative mesophilic sulfate-reducing bacteria. In *The Prokaryotes* (eds Balows, A. et al.) 3352–3378 (Springer, Berlin, 1992).
14. Perlthaler, J., Glöckner, F. O., Schönhuber, W. & Amann, R. Fluorescence in situ hybridization (FISH) with rRNA-targeted oligonucleotide probes. *Methods Microbiol.* 30, 207–226 (2001).
15. DIN Deutsches Institut für Normung E.V. *DIN 38406-5. German Standard Methods for the Examination of Water, Waste Water and Sludge: Cations (Group E); Determination of Iron (E 1).* (German Institute for Standardization, Berlin, 1983).
16. Din Deutsches Institut für Normung E.V. *DIN 38406-6. German Standard Methods for the Examination of Water, Waste Water and Sludge: Cations (Group F); Determination of Iron (E 2).* (German Institute for Standardization, Berlin, 1983).
17. Davide, M., Silvia, B., Andrea, G. & Carlo, M. Iron recovery from bauxite tailings red mud by thermal reduction with blast furnace sludge. *Appl. Sci.* 9, 4902. https://doi.org/10.3390/app9224902 (2019).
18. Austin, D. & Nivala, J. Energy requirements for nitrification and biological nitrogen removal in engineered wetlands. *Ecol. Eng.* 35, 184–192 (2009).
19. Saxena, N., Nawaz, A. & Lee, M. Comprehensive review of control and operational strategies for partial nitration/ANAMMOX system. *Ind. Eng. Chem. Res.* 58, 10635–10651 (2019).
20. Anderson, C. R. & Cook, G. M. Isolation and characterization of arsenate-reducing bacteria from arsenic-contaminated sites in New Zealand. *Curr. Microbiol.* 48, 341–347 (2004).
21. Lovley, D. R. Dissimilatory Fe(III)- and Mn(IV)-Reducing Prokaryotes. In *The Prokaryotes* (eds Dworkin, M. et al.) 287–308 (Springer, Berlin, 2006).

Acknowledgements

This research was supported by Grant No QG18.42 from Vietnam National University Hanoi (VNUH). We would like to thank Dr. Nam Dong Hoang for providing sludge of the anaerobic treatment system in Cu Chi, Ho Chi Minh city, Vietnam. Special thanks to Prof. Friedrich Widdel for his inspiration to the study. The authors assure that there is no conflict of interest from any other party regarding the research content presented in this paper.
Author contributions
C.P.L. as the first author of the manuscript prepared Figs. 1, 2, 3, 4 and together with H.T.D. and H.T.P. wrote the main manuscript text. H.T.N. contributed in performing FISH analyses and prepared Fig. 5. T.D.N. contributed in designing the laboratory-scaled treatment system and performing measurements in continuous operation experiments. Q.H.M.N. together with H.T.D. contributed as coordinators for whole research.

Competing interests
The authors declare no competing interests.

Additional information
Supplementary Information The online version contains supplementary material available at https://doi.org/10.1038/s41598-020-80057-y.

Correspondence and requests for materials should be addressed to H.T.D.

Reprints and permissions information is available at www.nature.com/reprints.

Publisher's note Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons licence, and indicate if changes were made. The images or other third party material in this article are included in the article's Creative Commons licence, unless indicated otherwise in a credit line to the material. If material is not included in the article's Creative Commons licence and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this licence, visit http://creativecommons.org/licenses/by/4.0/.

© The Author(s) 2021