AEROBIC AND ANAEROBIC AMMONIUM OXIDISING BACTERIA ENRICHMENT FROM MINED MUNICIPAL SOLID WASTE IN SHARON-ANAMMOX PROCESS

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ABSTRACT Ammoniacal nitrogen removal by novel processes like Single reactor system for high activity ammonia removal over nitrite (SHARON) and Anaerobic ammonium oxidation (ANAMMOX) processes are currently considered as advantageous than conventional processes. It requires aerobic ammonium oxidising bacteria (AOB) and anaerobic ammonium oxidising bacteria (AnAOB) to conduct the SHARON and ANAMMOX processes. This paper presents the feasibility of enriching the AOB and AnAOB using mined municipal solid waste (MSW) in batch reactors for a period of 37 days. In AOB reactor with nitrogen loading of 0.5 kg N/d showed Partial Nitrification Efficiency of 82.6% with $3.8 \times 10^8$ MPN/100mL of AOB population obtained. AnAOB reactor enriched with anammox biomass efficiently removed 78% of ammonia with the specific anammox activity reached up to 0.10 mg NH$_3$N/mg MLVSS/d. The nitrogen transformations along with the formation of intermediates (hydrazine and hydroxylamine), biomass development, free ammonia and free nitrous acid concentrations in the batch reactors confirmed the enrichment AOB and AnAOB biomass activity using mined MSW.

(Keywords: Mined municipal solid waste, ammoniacal nitrogen removal, aerobic ammonium oxidising bacteria and anammox)

INTRODUCTION

Landfill is the most common method for the disposal of municipal solid waste (MSW) worldwide. 1 - 4% of MSW is Total Kjeldahl Nitrogen (TKN) composed of proteins. This nitrogen is getting accumulated as ammoniacal nitrogen in the landfill leachate by ammonification and solubilization [10]. Landfill leachate contains ammoniacal nitrogen concentration between 500-3000 mg/L. Ammoniacal nitrogen has to be removed as it leads to algal growth, decreases performance of biological treatment systems, accelerates eutrophication, promotes oxygen depletion and increases aquatic toxicity [4]. Among the several biological processes for the removal of ammoniacal nitrogen, new innovative processes like SHARON (Single reactor system for high activity ammonia removal over nitrite) and ANAMMOX (Anaerobic ammonium oxidation) processes are being explored [1, 8, 12]. It has several advantages than conventional processes like non addition of organic carbon, non production of N$_2$O, lower oxygen, alkalinity consumption, sludge production and investment /operational cost needed [1, 6].SHARON process is a partial nitrification process which oxides half the influent ammonia to nitrite according to Equation 1, by aerobic ammonium oxidizing bacteria (AOB)[8, 12] such as Nitrosomonas europaea, Nitrosomonas eutropha, Nitrosolobus sp, Nitrosopira sp and Nitrosovibrio sp. [1, 18]. ANAMMOX process oxides ammonia anaerobically using the nitrite as electron acceptor, produced in the SHARON process to nitrogen gas (Equation 2) by anaerobic ammonium oxidising bacteria (AnAOB) such as Candidatus Brocadi a anammoxidans and Candidatus Kuenenia stuttgartiensis [5]. As, SHARON-ANAMMOX process has many benefits in terms of energy and resources, this process has been widely applied for removal of ammoniacal nitrogen from different ammonium rich wastewaters [16].

$$\text{NH}_4^+ + 0.75\text{O}_2 + \text{HCO}_3^- \rightarrow 0.5\text{NO}_2^- + 0.5\text{NH}_4^+ + \text{CO}_2 + 1.5\text{H}_2\text{O} \quad (1)$$

$$\text{NH}_4^+ + 1.32\text{NO}_2^- + 0.066\text{HCO}_3^- + 0.13\text{H}^+ \rightarrow 1.02\text{N}_2 + 0.26\text{NO}_3^- + 0.066\text{CH}_2\text{O}_{0.3}\text{N}_{0.15} + 2.03\text{H}_2\text{O} \quad (2)$$
But, this process is limited by the low availability of AOB and AnAOB biomass. Commonly, enrichment of AOB and AnAOB biomass are carried out using seed source such as Activated sludge (for AOB) and Nitrification sludge or Denitrification sludge or Anaerobic digestion sludge or/and Upflow anaerobic sludge blanket (for AnAOB)[18, 6, 9, 22].

Figure 1. Schematic diagram of experimental set up for enrichment of AOB

Usage of mined municipal solid waste as seed for enriching AOB and AnAOB were not studied so far. The aim of this present study is to assess the feasibility for enriching the AOB and AnAOB biomass from mined MSW in batch reactors.

Figure 2. Schematic diagram of experimental set up for enrichment of AnAOBs

MATERIAL AND METHODS

Characterisation of mined MSW

Mined MSW samples (i.e. partially degraded MSW (three years old)) were augured manually from different locations of an MSW dumpsite in Chennai, India. Mined waste was used as seed for enriching AOB and AnAOB biomass. Mined MSW samples were analysed for moisture content (MC), total solids (TS), volatile solids (VS), organic carbon (OC), ammonia nitrogen, nitrite and nitrate nitrogen was determined as per the standard methods [3].

Experimental setup for enrichment of AOB and AnAOB in batch reactors

Enrichment of AOB and AnAOB from mined MSW was carried out in two batch reactors namely, AOB and AnAOB reactor for 37 days as shown in Figure 1 and 2. The reactors were made up of borosilicate glass and run in duplicates. The reactors were
started up using the AOB and anammox enrichment medium as feed which was as specifically described by [6] and [18]. Mined waste and enriched anammox biomass from a laboratory scale anammox reactor, Chennai was used as seed for AOB and AnAOB reactor, respectively. The enriched anammox biomass used was identified as *Candidatus Brocadia anammoxidans* based on the 16s rRNA sequencing analysis (Gene bank database: under accession number-JQ972060). The loading details of the reactors are summarised in the Table 1.

Table 1. Loading details of the batch reactors

<table>
<thead>
<tr>
<th>S.No</th>
<th>Parameters</th>
<th>AOB reactor (Enrichment of AOB)</th>
<th>AnAOB reactor (Enrichment of AnAOB)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1.</td>
<td>Reactor volume</td>
<td>5 L</td>
<td>2.5 L</td>
</tr>
<tr>
<td>2.</td>
<td>Feed/Seed ratio</td>
<td>80/20</td>
<td>60/40</td>
</tr>
<tr>
<td>3.</td>
<td>Feed</td>
<td>4 L</td>
<td>0.86 L</td>
</tr>
<tr>
<td>4.</td>
<td>Seed</td>
<td>831.8 g of TS (Mined waste)</td>
<td>0.58 L (Enriched anammox biomass)</td>
</tr>
<tr>
<td>5.</td>
<td>Nutrients</td>
<td>NH₄HCO₃- 50 mM</td>
<td>NH₄SO₄ – 1mM; NaNO₂–1mM</td>
</tr>
</tbody>
</table>

The experiment was carried out at ambient temperature. Complete mixing was achieved within the reactors by manual methods. In AOB reactor, aerators (5 L/min) are used to maintain the dissolved oxygen level between 0.5 – 1.0 mg/L. AnAOB reactor was covered with black cloth to avoid light interferences and anoxic condition was maintained by suffocation (i.e. cutting the supply of oxygen) method. Gas generation in the AnAOB reactor was monitored by water displacement method. 60% of the total AnAOB reactor volume was used as working volume and 40% as head space.

The reactors were operated in fed batch mode. Batch reactors were fed with enrichment medium for maintaining the working volume in the reactor, usually reduced during sampling and evaporation (especially, in AOB reactors). The quantity of samples (25 mL) withdrawn once in a week was replaced with the addition of enrichment medium without any nitrogen supplement. The samples were analysed every week for enrichment of AOB and AnAOB bacterial population. Nitrogen transformations were studied by analyses the content of ammoniacal nitrogen, nitrite and nitrate nitrogen as per the standard methods [3]. The AOB and anammox development was determined by trace appearance of intermediates such as hydrazine [20] and hydroxylamine [7] and bacterial biomass accumulation by MLVSS and MLSS analysis [3]. AnAOB activity was exhibited by Specific Anammox Activity (SAA).

The population of AOB was enumerated by Most Probable Number (MPN) were carried out for biomass samples from AOB reactors [14, 13]. The medium used for MPN estimations for AOB contained, 0.5 g (NH₄)₂SO₄; 0.2 g KH₂PO₄; 0.2 g MgSO₄.7H₂O; 0.02 g CaCl₂.2H₂O; 0.0075 g phenol red and 10 ml trace elements, dissolved in a litre of distilled water. Trace elements solution contained 10 mg NaMoO₄.2H₂O; 20 mg MnCl₂.4H₂O; 0.2 mg CoCl₂.6H₂O; 2 mg CuSO₄.5H₂O; 10 mg ZnSO₄.7H₂O; 770 mg FeSO₄.7H₂O and 1.03 g Na-EDTA. pH of the medium was adjusted to 8.2 with 0.1 N NaOH and 5 ml of the medium were distributed into test tubes (150 mm x 15 mm) which were then plugged and autoclaved at 121°C for 10 minutes. After sterilisation, pH dropped to 7.7 - 7.8.

Ten fold dilutions of biomass samples from AOB reactors were prepared in distilled water. Aliquots (1 ml) of suitable dilutions were inoculated into test tubes containing medium and incubated at 37°C. The growth of AOB in the medium was estimated at intervals of 1 or 2 weeks. Presence or absence of ammonium oxidising bacteria was monitored visually by observing whether the colour of the test medium remained pink (no growth) or turned
yellow (growth) because of the decrease in pH resulting from the bacterial oxidation of ammonium. MPN was calculated depending upon the positive and negative tubes (Thomas formula). The nitrite accumulation rate in reactors was calculated by Partial Nitritation Efficiency (PNE) according to Liang and Liu [9] as given in the Equation 3.

\[
PNE = \frac{C_{(NO_2-N)_{eff}}}{C_{(NO_2-N)_{eff}} + C_{(NO_3-N)_{eff}}} \times 100% \quad (3)
\]

where, $C_{(NO_2-N)_{eff}}$ - Concentrations of nitrite nitrogen in the effluent (mg/L)

$C_{(NO_3-N)_{eff}}$ - Concentrations of nitrate nitrogen in the effluent (mg/L)

Concentrations of free ammonia and free nitrous acid in the reactors were calculated according to the Equation 4 and 5 suggested by Anthonisen et al. [17].

\[
FA (NH_3, \text{mg/L}) = 17 \times (NH_4) \times 10^{PH} \quad (4)
\]

\[
FNA (HNO_2, \text{mg/L}) = 46 \times (NO_2) \times 10^{PH} \quad (5)
\]

Statistical analyses were carried out using statistical software package SPSS (version 20.0) for Windows (SPSS, USA). The replicability of the batch reactors were validated by Pearson’s correlation analysis. Correlation results are statistically significant when significance coefficient is smaller than 0.05 (P < 0.05).

RESULTS AND DISCUSSION

Seed characteristics

Mined waste was used as seed for enrichment of AOB and AnAOB. The physico-chemical characteristics of the mined waste showed moisture content with 20.8% and had sufficient solids concentration of 79.2% with 13.3% of volatile solids. The total organic carbon was around 5%. Nitrogen load in the waste was ammoniacal nitrogen: 0.5 kg N/d, nitrite nitrogen: 0.01 kg N/d and nitrate nitrogen: 0.03 kg N/d. The enriched anammox biomass from a laboratory scale anammox reactor, Chennai was used as seed for anammox reactor. Inorganic nitrogen load was 0.041 kg N/d (ammoniacal nitrogen: 0.02 kg N/d, nitrite nitrogen: 0.001 kg N/d and nitrate nitrogen: 0.02 kg N/d). The biomass concentration of the anammox seed was 36.4 g/kg of suspended solids.

Enrichment of Aerobic Ammonium Oxidising Bacteria (AOB) in batch reactors

The enrichment of AOBs was conducted in 5L reactors with mined waste as seed. The aim of the study was to enrich AOB using mined MSW in the reactors by inhibiting the growth of Nitrite oxidising bacteria (NOB). NOBs are the main competitor for AOBs in substrate utilisation, if they are inhibited, AOB can build up faster and nitrite accumulation takes place [11]. The important operational parameters for AOBs to grow require optimum pH, temperature and DO. pH of the reactors were in the range of 7.6 - 8.2, DO: 0.5 – 1 mg/L and temp: 30 – 30.6°C, indicated the favourable conditions existed in the reactors for the AOBs to grow [8, 6, 14]. According to Paredes et al. [11], at temperatures greater than 15°C AOBs can grow faster than NOBs and around 25°C, the AOB can outcompete NOB. The conductivity and salinity of the reactors was 9.7 to 12.3 mS/cm and 6.2 to 8.1 psu, respectively.
Nitrogen transformations along with the Partial nitritation efficiency during the enrichment of AOBs in reactors are depicted in the Figure 3. The peak ammonia concentration in the reactor was 543.2 mg/L and 50% removal of ammonia was reached within 30 days achieving to NO₂-N/NO₃-N ratio of 1:1. The nitrite concentration elevated up to 316 mg/L, whereas nitrate level never exceeded 67 mg/L in the reactors. The nitrite accumulation rate is an indication of the activity of AOBs, as the rate increases it shows that the activity of AOBs are more. Partial nitritation efficiency (nitrite accumulation rate) enhanced from 25.6 to 82.6% in the reactors during the study period demonstrated the AOB activity.

Figure 3. Nitrogen transformations in the batch reactors during the enrichment of AOBs

FA concentrations in the reactors were above 0.1 mg/L throughout the study, which is an inhibitory value for NOBs [2] (Figure 3). The peak free ammonia concentration was only 58 mg/L and never reached the inhibitory value (150 mg/L) for AOBs. FNA concentrations maximum reached in the reactors was 0.034 mg/L (Figure 4) which was not an inhibitory range for AOBs and NOBs (0.2 mg/L) [2]. Hence, NOBs are mainly inhibited due to free ammonia and not due to free nitrous acid concentrations in the reactors.

Figure 4. Performance of the batch reactors during the enrichment of AOBs
Availibility of bicarbonates (alkalinity) is essential for AOBs, as the conversion of ammonia to nitrite is an acidifying process neutralised by bicarbonates [8]. Figure 4 showed adequate amount of bicarbonates were available in the reactors for AOB organisms to grow which was substantiated by biomass development. Biomass was measured by MLSS concentrations. Initially, MLSS concentration was 15,000 mg/L and reached to 6350 mg/L during the study period exhibited that under optimum reactor conditions only AOBs were developed. AOBs activity was confirmed by the occurrence of hydroxylamine and hydrazine concentrations in the reactors (Figure 4). Accumulation of intermediates was observed within a week in the reactors. The range of hydroxylamine and hydrazine was 0.001 to 0.010 mg/L and 0.005 to 0.035 mg/L, respectively [12]. Further, the confirmation of AOB bacterial population was investigated by conducting the MPN analysis specific for AOB [14, 13]. MPN obtained for the reactor biomass sample were 3.8 ×10^5 MPN/100mL, illustrated the adequate quantity of AOB population existed in the reactor. The results authenticated that ample concentrations of intermediates with biomass development and AOB specific MPN population acquired in the batch reactors established the enrichment of AOBs. The replicability of the experiment was validated by Pearson’s correlation analysis showed significantly strong correlation with 85 – 95% in the reactors.

Enrichment of Anaerobic Ammonium Oxidising Bacteria (AnAOB) in batch reactors

The anammox bacterial enrichment was observed in terms of operational parameters (pH, temperature), nitrogen transformations, biomass development and presence of trace amounts of intermediates [5, 18]. pH and temperature of the AnAOB reactors was 7.3 – 8.4 and 30 – 32°C, demonstrating that the reactors were under optimum conditions for the anammox bacterial growth in the reactors [21]. The conductivity and salinity of the reactors was 2.7 to 3.0 mS/cm and 1.4 – 1.7 psu, respectively.

Nitrogen transformations during the enrichment of AnAOBs in the reactors are depicted in the Figure 5. The maximum level of ammonia concentration was 98 mg/L reached within a week and ammonia removal efficiency increased up to 78% during the end of the study period. Simultaneous reduction in the nitrite concentration from 60.2 to 1.5 mg/L in 37 days demonstrated the growth of anammox bacteria with the consumption of nitrite as electron acceptor. Concentrations of nitrate prevailed in the reactor was in the range of 2.3 to 17.7 mg/L. Nitrate production illustrated that the nitrite converted to nitrate for producing the reducing equivalents for CO_2 fixation in anammox process [17]. The presence of nitrite and nitrate concentrations in the reactors also revealed existence of AOBs and NOBs along with the anammox bacteria. Free ammonia and free nitrous acid concentrations in the reactors are depicted in the Figure 5 and 6. FA concentrations in the reactors were above 0.1 mg/L showed NOBs are inhibited. Free nitrous acid concentrations in the reactors elevated to 0.007 mg/L and never reached the inhibitory value of 0.2 mg/L in the study period [2].
Alkalinity can be used as the indicator parameter for the anammox activity, as anammox bacteria consumes bicarbonate as their carbon source [21]. Variations in the bicarbonate concentrations in the reactors are depicted in the Figure 6. As sufficient bicarbonates were available in the reactor, the nitrogen removal by anammox process was increased. The anammox development in the reactor can be demonstrated by the Specific anammox activity. Initially, SAA was 0.005 mg NH₄-N/mg MLVSS/d and in 37 days of reactor operation it reached to 0.10 mg NH₄-N/mg MLVSS/d, which is comparable with the outcomes of the study conducted by Wang et al. [19].

The colour of the biomass in the reactor was also changed from black colour to brownish colour in the end of the study period due to the development of anammox bacteria. The biomass growth with the occurrence of trace concentrations of hydrazine and hydroxylamine authenticated the AnAOB enrichment (Figure 6). The hydrazine concentration was in the range of 0.001 to 0.003 mg/L and hydroxylamine was around 0.0002 to 0.001 mg/L in the reactors. The trace occurrences of intermediates (hydrazine and hydroxylamine) with the specific anammox activity proved the enrichment of AnAOB [15]. The reactors are validated by the duplicate reactors, which gave 90 – 95% correlation.

CONCLUSION

The study confirmed the feasibility for using mined MSW as seed for enriching the aerobic and anaerobic ammonium-oxidising bacteria in batch reactors. The enrichment of AOB and AnAOB operated for 37 days with nitrogen loading of 0.5 kg/N/d was authenticated by nitrogen transformation profiles, biomass accumulation, alkalinity, FA, FNA concentrations and occurrences of intermediates such as hydrazine and hydroxylamine in the reactors. Further, AOB activity was validated by 82.6% of Partial Nitritation Efficiency with AOB populations of 3.8 x10⁸ MPN/100mL and AnAOB activity by 78% removal of ammoniacal nitrogen with specific anammox activity of 0.10 mg NH₄-N/mg MLVSS/d.

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REFERENCES


