B-17 Recovery of dissolved methane with a degassing membrane in a UASB reactor

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1. INTRODUCTION

There is an increasing interest in developing renewable energy technologies. Anaerobic digestion has been known as a renewable energy-producing process. The methane (CH₄) produced in this process can be used as a fuel. In especial, an upflow anaerobic sludge blanket (UASB) reactor is considered to be effective in treating high-strength wastewaters. In a UASB reactor, the biogas is produced in the liquid phase (bulk liquid) and then evolved to the gas phase (headspace) while a part of CH4 is still dissolved in the bulk liquid. The dissolved CH4 should be considered as it is not possible to be captured in the usual processes. Significance of the dissolved CH4 has been recognized by some researchers. For example, a comparative analysis by Hartly et al. 1) confirmed that CH₄ super-saturation resulted in significant loss of CH4 in the effluent. The CH4 super-saturation is more significant at lower temperatures. Therefore, we operated a bench-scale UASB reactor equipped with a degassing membrane to recover the dissolved CH4. The effect of temperature on CH4 recovery efficiency was investigated.

2. MATERIALS AND METHODS

A bench-scale UASB reactor was operated²⁾ at various temperatures and trans-membrane pressures (Table 1). A three-layer composite hollow fiber membrane (MHF) (Mitsubishi Rayon Engineering Co., Ltd., Tokyo, Japan) was installed in a degassing membrane reactor (DMR), which followed the UASB reactor to recover dissolved CH₄ in the bulk liquid. The UASB reactor was covered with a water jacket to maintain the temperature, on the other hand, the DMR was operated at room temperature. The UASB reactor was fed with a synthetic wastewater²⁾ at a HRT of 10 h. After the gas production reached steady-state, dissolved gas in the bulk liquid was recovered through the degassing membrane using an air pump (Model

Table 1 Reactor operating condition.

Phase	Temperature (°C)	Trans-membrane pressure (kPa)		
1	35	0		
2	35	-50		
3	35	0		
4	35	-50		
5	35	-80		
6	35	0		
7	25	o oto		
. 8	25	-80		
9	25	0		
10	15	0		
11	15	-80		
12	15	0		
13	35	1980 to 0.00 o 1 mile		

APN-110KV-1, Iwaki Co., Ltd., Tokyo, Japan). The experiment was divided into 13 experimental phases based on operating conditions (i.e., trans-membrane pressure and temperature) (Table 1). The gas compositions in the headspace of the UASB reactor and the membrane were measured by a gas chromatography system (GC-14B; Shimadzu Co., Kyoto, Japan). The dissolved gas compositions were measured by the headspace method2). The biogas volumes were measured at ambient temperature. The total CH4 was defined as the sum of CH4 evolved into headspace, recovered from membrane and discharged into effluent. The CH₄ recovery efficiency was defined as a ratio of headspace CH₄ plus CH₄ in the membrane to headspace CH4. The concentrations of total COD (T-COD) and dissolved fraction of COD were measured using a Hach method (Method 8000). Particulate COD (P-COD) was calculated as T-COD minus D-COD. concentrations were determined by a high-performance liquid chromatography system.

3. RESULTS AND DISCUSSION

(1) COD removal.

The study was divided into 13 experimental phases based on operating conditions. Reactor performance (i.e., COD concentrations and gas production rate), temperature and pH for 13 phases are presented in Figure 1 and average (± standard deviation) values are summarized in Table 2. Effluent T- and D-COD concentrations were low at the beginning of the operation with T-COD removal efficiency of more than 85%. After adjusting pH to around 7 at day 51, gas production rates into headspace became stable. These rates were significantly correlated positively with temperature. Decrease in gas production rates at lower temperatures reflected lower COD removal efficiency due to decrease in microbial activity and could be explained by increase in solubility of biogas in bulk liquid at lower temperatures. In general, most reactions in the biodegradation of organic matters require more energy to proceed at lower temperatures than at the optimum temperature of 37°C for anaerobes³⁾. Hence, lowering the operational temperature leads to a decrease in the maximum specific growth and substrate utilization rates of the anaerobes. As a result, biological reactions proceed much slower under psychophysics conditions than under mesophilic conditions. Identical COD removal and gas production rates in the normal operation conditions between changes in the operating conditions (phases 1, 3, 6 and 13) showed good reproducibility of the reactor operation. Interestingly, the removal efficiency of P-COD rather than D-COD at lower temperatures tended to be improved during degassing periods. Lettinga et al. 3) indicated that in psychrophilic reactors, particles would settle more slowly because of the deterioration of liquid-solid separation probably attributed to increase in the viscosity of bulk liquid at

lower temperatures. This result opens up the possibility to improve COD removal efficiency by degassing of bulk liquid in a UASB reactor operated at lower temperatures.

(2) Gas production.

Figure 2 shows the headspace gas composition profiles, the production rates of CH₄ gas evolved into headspace, recovered from membrane and discharged into effluent, and CH₄ recovery efficiency in the UASB reactor. Average (± standard deviation) values of these rates and CH₄ recovery efficiency throughout the operation are summarized in Table 3. Degasification did not significantly affect gas compositions in the headspace of the

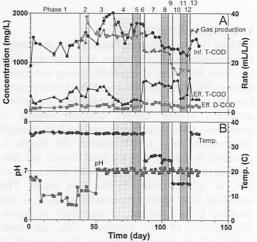


Figure 1 Concentrations of T-COD in influent and Tand D-COD in effluent, and gas production rate (A) and temperature and pH (B) in the UASB reactor. The gray area and the dotted lines represent degassing periods and the timing of change in temperature, respectively.

Table 2 Summary of average (± standard deviation) of the rates of T-COD lading, P-COD discharged and D-COD discharged, and the removal efficiencies of T-COD and D-COD in the UASB reactor. The gray area represents degassing operation.

Phase	T-COD loading (mg/L/h)	P-COD discharged (mg/L/h)	D-COD discharged (mg/L/h)	T-COD removal (%)	D-COD remova (%)
1	121 ± 13	18 ± 7	9±2	78 ± 9	93 ± 2
2	140 ± 9	23 ± 8	10 ± 1	76 ± 6	93 ± 1
3	173 ± 28	28 ± 8	12 ± 2	77 ± 7	93 ± 1
4	179 ± 44	12 ± 3	8±2	89 ± 3	96 ± 2
5	174 ± 15	15 ± 2	10 ± 1	86 ± 1	94 ± 1
6	169	41	18	65	89
7	152 ± 8	40 ± 5	18 ± 2	62 ± 3	88 ± 1
8	132 ± 5	40 ± 4	15 ± 1	58 ± 1	89 ± 1
9	125	39	10	61	92
10	124 ± 7	49 ± 2	13 ± 1	50 ± 1	90 ± 1
11	118 ± 6	21 ± 3	10 ± 1	74 ± 4	91 ± 1
12	129	57	12	46	90
13	143 ± 7	15 ± 2	12 ± 2	81 ± 1	92 ± 1

UASB reactor (Figure 2A). Exceptionally, CH4 concentration slightly increased and concentration decreased at 15°C. CH4 evolution rate into headspace decreased with decrease in temperature (Table 3) due to decrease in gas production rate (Figure 1A). Dissolved CH4 concentration increased from 63 ± 1 mg-COD/L at Phase 6 to 81 \pm 6 mg-COD/L at Phase 7 and 103 \pm 5 mg-COD/L at Phase 10 due to increase in solubility of CH₄ in bulk liquid at lower temperatures, resulting in more significant loss of dissolved CH4 from the UASB reactor (Figure 2B). By degasification, CH₄ could be recovered from bulk liquid in the DMR and dissolved CH₄ concentration decreased (Table 3). The CH₄ recovery rates from membrane were higher at lower temperatures (Phases 8 and 11) as compared with that at 35°C (Phase 5) because of higher dissolved CH₄ concentrations at lower temperatures. Consequently, the CH₄ recovery rate from the headspace and the membrane at lower temperatures (Phases 8 and 11) were comparable to that during normal operation at 35°C (Phases 1, 3 and 6)). CH₄ recovery efficiencies during degassing periods were between 1.09 and 1.19 at 35°C, 1.32 at 25°C and 1.53 at 15°C. Thus, degasification was more efficient at lower temperatures. Particularly under temperate climate conditions, many low and medium strength wastewaters are discharged at low ambient temperatures. On the other hand, practically all full-scale applications of anaerobic wastewater treatment so far were restricted to wastewater with temperatures exceeding 18°C3). Hence, in general, a significant amount of energy is required to bring the wastewater temperature up to the optimal mesophilic range (30 to 40°C). This puts a heavy burden on the economy of the wastewater system. We concluded that the UASB reactor equipped with the DMR was a promising alternative to the conventional UASB reactor.

CONCLUSIONS

A UASB reactor equipped with the degassing membrane reactor was operated to recover dissolved CH₄ in bulk liquid. Decrease in temperature resulted in decrease in CH₄ evolution rate to the headspace. Degasification could improve total CH₄ recovery efficiency especially at lower temperatures.

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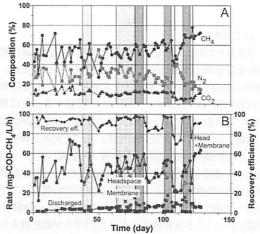


Figure 2 Headspace gas composition profiles (A), and the production rates of CH₄ gas evolved into headspace, recovered from membrane and discharged in effluent, and CH₄ recovery efficiency in the UASB reactor (B).

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Table 3 Summary of average (± standard deviation) of the production rates of CH₄ gas evolved into headspace, recovered from membrane and discharged in effluents from the UASB and the degassing membrane reactors, and CH₄ recovery efficiency in the UASB reactor. The gray area represents degassing operation.

Phase	CH ₄ production rate to head space (mg-COD/L/h)	CH ₄ loss from the UASB reactor (mg-COD/L/h)	CH ₄ recovery rate from membrane (mg-COD/L/h)	CH ₄ loss from the membrane reactor (mg-COD/L/h)	CH ₄ recovery efficiency (-)
1	46 ± 18	5 ± 1	TERRITOR OF	3±1	III THE SHALL SHAL
2	44 ± 14	5±0	4±2	2±0	1.09 ± 0
3	40 ± 9	6 ± 1		5 ± 1	
4	42 ± 4	6±0	7 ± 4	2±0	1.18 ± 0
5	47 ± 4	6±0	9±1	1±0	1.19 ± 0
6	42 ± 10	6±0	363-77F	5 ± 0	
7	37 ± 4	8 ± 1		7 ± 1	
8	39 ± 3	9±1	12 ± 3	1±0	1.32 ± 0
9	24 ± 4	8 ± 0		7 ± 1	
10	22 ± 1	10 ± 1		9 ± 1	
11	29 ± 3	· 10 ± 0	15 ± 4	1±0	1.53 ± 0
12	30	11 ± 1	The state of the s	9 ± 0	
13	58 ± 7	7 ± 1		6±0	