Contents lists available at ScienceDirect

Biochemical Engineering Journal

journal homepage: www.elsevier.com/locate/bej



Anaerobic treatment of low-strength wastewater with a high fraction of particulate matter in an unconventional two-phase ASBRs system

Andres Donoso-Bravo*, Gonzalo Ruiz-Filippi, Rolando Chamy

Escuela de Ingeniería Bioquímica, Facultad de Ingeniería, Pontificia Universidad Católica de Valparaíso, General Cruz 34, Valparaiso, Chile

ARTICLE INFO

Article history: Received 23 May 2008 Received in revised form 22 September 2008 Accepted 21 October 2008

Keywords: Anaerobic processes Two-phase system Anaerobic sequencing batch reactor Domestic wastewater Wastewater treatment Acetic acid

1. Introduction

The anaerobic treatment of low-strength wastewater, such as domestic wastewater, has not been extensively implemented because the process has a lower performance under these conditions. This is because the concentration of organic matter (substrate) is in the same range or near the value of the saturation constant of the anaerobic biomass (*Ks*). Hence, biomass is not growing at the maximum specific growth rate and, at the same time, is consuming the substrate at a low-degradation rate (with respect to the Monod kinetics). Moreover, if a high fraction of particulate or suspended organic matter is present (\geq 0.45 µm), the overall reaction slows down even further since hydrolysis of the complex composite particulates must first occur [1], which can become the limiting step of the whole process.

The application of a two-phase anaerobic system to treat this type of effluent may present significant advantages over a onephase system. The separation of the main reactions of anaerobic digestion, acidogenesis and methanogenesis, allows the selection and enhancement of microbial populations in each reactor in terms of temperature and pH as well as control of the intermediate products [2]. Pohland and Ghosh [3] proposed the separation of acid-producers and acid-consumers in two reactors in order to opti-

ABSTRACT

The results of a two-phase anaerobic system using anaerobic sequencing batch reactors (ASBRs), treating low-strength wastewater (COD \sim 500 mg/L) with a high fraction of particulate organic matter (70%, COD basis), are presented. Two reactors in series were used; the first one was hydrolytic–acidogenic, while the second one was methanogenic. This configuration was proposed to promote high efficiency solids removal. During the experiment, 69% and 50% efficiencies of total COD removal were obtained for OLRs of 0.63 and 1.22 kgCOD/(m³ d), respectively. Values of the solubilized organic fraction (SOF) achieved in the hydrolytic–acidogenic reactor were within the range of 0.3–0.6 gCOD_{solubilized}/gpCOD_{removed}, and the average acidified organic fraction (AOF) was 0.6 gCOD_{VFA-produced}/gsCOD_{fed}. The methanogenic reactor had a VFA removal fraction (VFARF) between 0.4 and 0.6 gCOD_{VFA-removed}/gCOD_{VFA-fed} for the OLR of 0.63 and 1.22. The two-phase ASBR system is suitable, and can be implemented, for the anaerobic treatment of this kind of wastewater.

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mize the activity of each population, improving their particular environmental and operational conditions.

The implementation of these systems has focused on the treatment of high-strength waste and wastewater using continuous anaerobic reactors. Different configurations have been evaluated: CSTR + UASB [4,5], CSTR + CSTR [6–10], and CSTR + UAF [11].

The anaerobic sequencing batch reactors (ASBRs), developed by Sung and Dague [12], is an anaerobic digestion system that works through consecutive cycles of operation, each of which has the following stages: feeding, reaction, settling, discharge and idle time. At the beginning of the reaction stage, the concentration of organic matter is at its maximum level, which in turn maximizes the degradation rate since the substrate is greater than the *Ks*. This situation presents an important kinetic advantage over continuous systems, where the concentration of substrate in the reactor is equal to the effluent, which is lower than the *Ks* [13]. Thus, the application of a two-phase ASBR system, besides enriching the development of both populations, can increase their growth rate and hence their degradation ability, improving the acidogenesis and hydrolysis rates in the first reactor and methanogenesis rate in the second one, and the overall performance of the system.

Few studies have evaluated or tested a two-phase system with ASBRs. Dugba and Zhang [14] worked with two ASBR in series, but in this case the objective was to study the effect of temperature (thermofilic/mesofilic, thermofilic/thermofilc) using an industrial effluent. Bouallagui et al. [15] reported a study with two-phase ASBR system (acidogenic and methanogenic), in which fruit and

^{*} Corresponding author. Tel.: +56 32 2273819; fax: +56 32 2273803. *E-mail addresses*: adonosobravo@gmail.com,

andres.donoso.b@mail.ucv.cl (A. Donoso-Bravo).

¹³⁶⁹⁻⁷⁰³X/\$ – see front matter 0 2008 Elsevier B.V. All rights reserved. doi:10.1016/j.bej.2008.10.011

Nomencl	lature
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	ASBBR	anaerobic sequencing batch biofilm reactor
	ASBR	anaerobic sequencing batch reactor
	COD _{VFA} i	COD of VFA of the ASBR1 influent
	COD _{VFA} 1	COD of VFA of the ASBR1 effluent
	COD _{VFA} 2	COD of VFA of the ASBR2 effluent
	$k_{ m H}$	hydrolysis catalytic constant
	Ks ₁	saturation constant of acidogenic biomass
	Ks ₂	saturation constant of methanogenic biomass
	OLR	organic load rate
	pCOD	particulate COD
	pCODi	particulate COD of the ASBR1 influent
	pCOD1	particulate COD of the ASBR1 effluent
	sCOD	soluble COD
	sCODi	soluble COD of the influent
	sCOD1	soluble COD of the ASBR1 effluent
	tCOD	total COD
	VFA	volatile fatty acids
		5
Greek letters		
	$\mu_{ m M1}$	maximum specific growth rate of acidogenic
		biomass
	$\mu_{{ m M2}}$	maximum specific growth rate of methanogenic
		biomass

vegetable wastes were treated. The focus was to optimize each of the reactions, which resulted in over 80% hydrolysis in the first reactor and high percentages of biogas production (320 L/kgCOD_{affluent}) in the second reactor. Both studies were carried out with high-strength substrates.

The objective of this study was to implement and evaluate the performance of a two-phase ASBR system in the treatment of lowstrength wastewater with a high fraction of particulate organic matter, simulating domestic wastewater.

2. Materials and methods

2.1. Experimental setup

Two laboratory scale reactors of acrylic, both with a total volume of 6L and effective volume of 5L were used (ASBR1, hydrolytic–acidogenic and ASBR2, methanogenic). Both reactors were jacketed and the temperature kept at approximately 35 °C. The

reactor was mixed by liquid recirculation during the reaction step. Peristaltic pumps were used for feedings, effluent discharge and mixing. The automated operation of the system (pumps, valves and mixer) was programmed using timers and counters in Microwin STEP7 using a PLC Siemmens S7200. Fig. 1 shows a diagram of the system used in this study. The time distribution for the stages in each reactor was as follows: feeding 15 min, settling 30 min, discharge 20 min, and idle time 5 min. The reaction times were 410 and 180 min for OLR1 and OLR2, respectively.

The ASBR1 was fed with synthetic wastewater that simulated domestic wastewater with the following composition: Potato solution 2.0 g/L, ovoalbumin 0.12 g/L, vegetable oil 0.025 g/L, urea 0.033 g/L. In addition to the macronutrients NH₄Cl 0.074 g/L and KH₂PO₄ 0.01 g/L, the following micronutrients were added: FeCl₃·4H₂O 4.0 mg/L, ZnCl₂ 0.1 mg/L, MnCl₂·4H₂O 1.0 mg/L, CoCl₂·6H₂O 4.0 mg/L, CuCl₂·2H₂O 0.06 mg/L, NiCl₂·6H₂O 0.1 mg/L, H₃BO₃ 0.1 mg/L, Na₂SeO₃·2H₂O 0.2 mg/L and (NH₄)6MoO₂·4H₂O 0.18 mg/L. To maintain the alkalinity of the system, 0.5 g/L NaHCO₃ was added. The ASBR1 was seeded with acidogenic sludge, with negligible methanogenic activity. The ASBR2 was fed with the effluent of the ASBR1 and it was seeded with an anaerobic granular sludge, with a methanogenic activity of $0.62 \text{ gCOD}_{CH_4}/(\text{gVSS d})$. Each of the reactors was inoculated in order to obtain an initial biomass concentration of 5 gVSS/L.

2.2. Start-up and operation of the two-phase system

The start-up was carried out with the operation of ASBR1 at an OLR of $1.21 \pm 0.12 \text{ kgCOD}/(\text{m}^3 \text{ d})$ (3 cycles/day) and a pCODi/tCODi ratio of 0.66 ± 0.04 . Once start-up of the ASBR1 was complete, the ASBR2 was connected to the system and fed with the discharge of ASBR1. A system OLR of $0.63 \pm 0.064 \text{ kgCOD}/(\text{m}^3 \text{ d})$ with a pCODi/tCODi ratio of 0.67 ± 0.04 was applied, as well as a second system OLR of $1.22 \pm 0.16 \text{ kgCOD}/(\text{m}^3 \text{ d})$ (6 cycles/day) with a pCODi/tCODi ratio of 0.70 ± 0.04 . The dynamic behavior of the reactor was analyzed by follow-up procedures, which were done by taking consecutive samples during the reaction stage in both reactors. The first samples were taken from the ASBR1. Samples were then taken from ASBR2 when the effluent was fed from the ASBR1.

2.3. Analytical methods

During the operation of the system, the influent to the system, the ASBR1 and the ASBR2 effluents were sampled for the following analysis: total COD (tCOD) and soluble COD (sCOD) were measured by the Closed Reflux-Titrimetric Method [16]; samples for sCOD



Fig. 1. Schematic diagram of the two-phase system implemented for the anaerobic treatment of low-strength wastewater with a high fraction of particulate organic matter.

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