



# ENHANCING H<sub>2</sub> PRODUCTION BY ANAEROBIC CO-DIGESTION OF ORGANIC SOLID WASTES AND CHEESE WHEY

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**Introduction.** Anaerobic Digestion (AD) has been used as a suitable way for the treatment of organic solid wastes (OSW). Additionally, in this process it is possible to obtain hydrogen (H<sub>2</sub>), which is considered as a potential biofuel due to its energetic content. There are several reports on H<sub>2</sub> production using pure cultures and model substrates [1]. However, when OSW are used as feedstock the application of axenic cultures is not possible. Therefore the use of mixed cultures is a suitable option for this process.

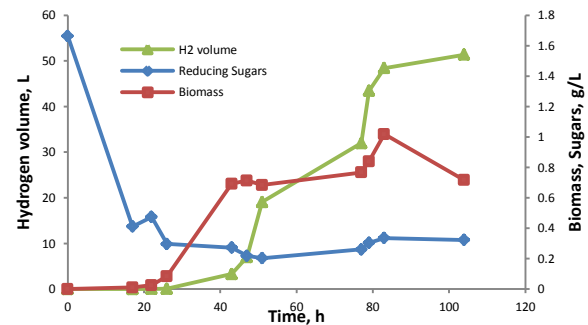
One of the challenges of the AD is the increment of H<sub>2</sub> production yields. Co-digestion of OSW with other residues or wastewaters that provide complementary nutrients has been reported to increase H<sub>2</sub> production. Moreover, the presence of proteins allows the release of NH<sub>3</sub><sup>+</sup> when hydrolyzed, so that a better control on pH can be attached [2].

In this work, co-digestion of organic wastes and cheese whey was tested for increasing the H<sub>2</sub> production using a mixed anaerobic culture.

**Methods.** Two batch cultures were performed in a 30 L bioreactor (15 L operation volume). The inoculum for the experiments was taken from an anaerobe digester, and heat pre-treated to eliminate methanogenic bacteria. A mixture of OSW was combined with cheese whey in a 1:1 proportion and then half diluted in water. After inoculation, the bioreactor was flushed with N<sub>2</sub> to guarantee anaerobic conditions. Operation conditions were 35 °C, pH 5.5, 150 rpm. Concentration of protein, reducing sugars, organic acids and hydrogen were measured by Bradford, DNS, HPLC and TCD chromatography, respectively. NaOH 4N and H<sub>2</sub>SO<sub>4</sub> 20% v/v solutions were automatically added to maintain pH.

**Results.** Fig. 1 shows the performance of the co-digestion process during the H<sub>2</sub> production. The initial concentration of reducing sugars (1.6 g/L) were 90% during the first 20 h and then maintained below 0.4 g/L. Biogas was produced from the beginning of the process (data not shown) but H<sub>2</sub> production started after 28 h. The same occurred for cellular growth. This lag phase can be associated to an initial hydrolysis

process. At this stage, the NaOH solution was added automatically to maintain pH. After 40 h, H<sub>2</sub> production rate increased and the growth rate decreased. At 60 h of culture no more NaOH solution was required for pH control but H<sub>2</sub>SO<sub>4</sub> was consumed. This fact could be explained by an increase in the protein hydrolysis rate, which produced alkaline condition due to constant releasing of NH<sub>4</sub><sup>+</sup> groups. At the beginning of the culture, bacteria consumed mainly free sugars. Then, when availability of sugars dropped hydrolysis of proteins increased (60 h). At this time, the highest hydrogen content in biogas was detected (51%). After 110 h, 51 L of H<sub>2</sub> were produced and concentration of reducing sugars was above 0.3 g/L. The highest H<sub>2</sub> production was associated with a better pH a control due to the NH<sub>3</sub><sup>+</sup> release.



**Fig.1** Hydrogen production during co-digestion process.

**Conclusions.** During this process, a lag phase between inoculation and the start of hydrogen production was identified. Although H<sub>2</sub> production started at 28 h, the highest production rate occurred when the availability of sugars decreased and proteins started to be hydrolyzed. The highest H<sub>2</sub> content in the biogas was determined in a relatively short time (60-80 h) makes the co-digestion a suitable process for the H<sub>2</sub> production.

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**References.** 1. Tao Y, Liu X, Zhou Z, Lee J, Yang, C. (2011). *Journal Bacteriol.* 194(2):274:283  
2. Martínez-García G, Johnson A, Bachmann R. (2007). *Inter Biodeterioration & Biodegradation.* 59(4):273:288.