Original Research

Producing Hydrogen in Sequential Dark and Photofermentation from Four Different Distillery Wastewaters

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Received: 1 August 2019 Accepted: 1 September 2019

Abstract

Four different distillery wastewaters were applied as the substrates for dark and photofermentation – wheat stillage (WTL), wheat syrup (WSR), maize stillage (MTL) and maize syrups (MSR). WTL was found to be the most effective substrate in dark fermentation, with a hydrogen production of 0.88 L H_2/L_{medium} and a yield of 1.17 L H_2/L_{WTL} . In the photofermentation experiments, the highest hydrogen production (1.7 L H_2/L_{medium}) and the highest yield (8.6 L H_2/L_{WSR}) were obtained with WSR as the substrate A sequential two-step treatment, comprising dark fermentation followed by photofermentation, was applied to the same four substrates. WTL and MTL were the substrates for dark fermentation and the effluent after 5-fold dilution was used as the substrate in photofermentation. The maximum hydrogen production in the two-step system was 4.47 L H_2/L_{WTL} . Simultaneously, the organic compounds from the waste and those formed in dark fermentation were completely utilized, except for ethanol and methanol, which were 19% and 10% utilized, respectively.

Keywords: distillery wastewaters, dark fermentation, photofermentation, biohydrogen, two-step system

Introduction

Large-scale fermentative production of ethyl alcohol is mainly based on cane sugar, corn, potatoes and different kinds of grains serving as the sources of hydrocarbons. Production of ethyl alcohol from these substrates generates a significant amount of wastewater. In 2017 the production of ethanol in the United States was 1.02 million barrels per day and has been increasing year by year [1]. In Poland the annual production of ethyl alcohol reached one million hectoliters [2]. It was estimated that the world demand for ethyl alcohol would increase by 70% in the next 10 years because of its application as a gasoline component [1]. Wastewaters coming from alcohol production, after drying, can be used for the cattle feed. However, the cost of the pretreatment is very high. Therefore, cheaper methods of utilization of the distillery wastewaters would have to be applied. Fermentative hydrogen production processes seem to be an attractive option, because they offer a dual function of waste reduction and hydrogen energy production [3, 4].

Although the application of different wastewaters from the food industry for hydrogen production in dark fermentation is well described in literature [5-9], the search for better yields, higher reaction rates

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and lower biomass production is still a challenge. The wide selection of substrates and microorganisms that can be used in the dark fermentation process opens a great chance for development of this method on a large scale. Recent papers [10-12] as well as our previous studies [13] can be a basis for microbiological hydrogen production with distillery wastewaters on a large scale. Wastewaters from the food industry can be used successfully also in photofermentative processes, where light-dependent heterotrophic bacteria decompose organic compounds, mainly volatile fatty acids, into hydrogen and carbon dioxide. Numerous papers [14-16] show that Rhodobacter sphaeroides can efficiently produce hydrogen from wastewaters coming from food industries like dairies, breweries, distilleries and others. Increasing the production of ethanol and bioethanol will result in higher amounts of waste, which will require efficient, environmentally friendly and economical methods of utilization. Combined systems, which link dark fermentation with photofermentation in a one- or two-step process, can lead to total biodegradation of organic compounds in waste and higher efficiency of hydrogen production than in separate processes.

The aim of this study was to compare four types of wastewaters coming from a local distillery (producing 113000 m³ waste/day) as the substrates for hydrogen production in dark fermentation, photofermentation and the two-step system. These wastewaters originated from different stages of ethanol production based on wheat and maize. Many publications have presented the production of hydrogen from distillery wastewaters [17-20], but none have compared different types of distillery wastewaters. We would like to stress there have so far not been papers describing systems as mentioned above. This paper proves that in the two-step system the vast majority of organic compounds from wastewaters degrade completely to H_2 and CO_2 with high efficiency.

Table 1. Characteristic of wastewater.

	Wastewater				
Properties	W	heat	Maize		
	WTL	WSR	MTL	MSR	
Total solid (TS) [g/mL]	0.128	0.136	0.130	0.234	
Elemen	of TS				
C [%]	48.8	45.8	48.9	47.1	
H [%]	6.9	7.7	8.5	9.2	
N [%]	6.5	7.1	4.1	2.6	
S [%]	0.3	0	0.003	0.175	
Total COD [g O ₂ /L]	114	197	251	325	
COD in solution [g O ₂ /L]	43	86	13	58	
VSS [g/mL]	0.122	0.126	0.123	0.214	
рН	4.0	3.8	3.8	3.4	

Experimental

Wastewaters

Four different wastewaters originating from the processing of wheat and maize in the local distillery (Murowana Goślina, Poland) were substrates in the fermentation processes. These wastes were wheat stillage (WTL), wheat syrup (WSR), maize stillage (MST) and maize syrup (MSR). Stillages are a waste obtained after first distillation while syrups are obtained after decantation and condensation of stillages. Characteristics of the applied wastes are given in Tables 1 and 2.

Inoculum

In dark fermentation an anaerobic digested sludge originating from a municipal purification unit was used as an inoculum. In the pretreatment process it was boiled for 15 minutes to eliminate methanogenic bacteria. The total nitrogen content in inoculum (measured by Kjeldahl method) was 0.9 g/L, carbon content was 12.5 g/L and the amount of volatile suspended solids (VSS) was 29.1 g/L.

In photofermentation *Rhodobacter sphaeroides* O.U.001 (ATCC 49419) bacteria were used. Bacteria were cultivated on Van Niel's medium containing K_2HPO_4 (1.0 g/L), MgSO₄ (0.5 g/L), yeast extract (10 g/L) and tap water filled up to 1 L and then activated to hydrogen production on modified Biebl and Pfennig medium according to the procedure already described [21].

Medium and Procedure

The medium used in dark fermentation contained 10, 20, 40, 60 or 75% v/v of distillery wastewaters. The medium was enriched with the following components (g/L): 1.0 NaHCO_3 , $0.5 \text{ NH}_4 \text{Cl}$, $0.25 \text{ KH}_2 \text{PO}_4$,

Table 2. Organic compounds in wastewater.

i	Wastewater					
Compound	Wh	leat	Maize			
[5, 2]	WTL	WSR	MTL	MSR		
Glucose	1.5	0	0.15	0		
Lactic acid	2.0	9.3	1.3	10.0		
Acetic acid	0.2	0.34	0.09	0.3		
Butyric acid	0	2.4	0	0.06		
Malic acid	0	3.8	0.3	2.3		
Glycerol	5.2	21.0	4.0	27.0		
Succinic acid	0.6	2.0	0.4	2.5		
Methanol	2.0	4.6	1.6	4.5		
Ethanol	0.08	0.04	0.04	0.2		

0.25 K_2HPO_4 , and 0.32 MgSO₄*7H₂O; and microelements 0.005 FeCl₃, 0.0032 NiSO₄, 0.005 CaCl₂, 0.0007 Na₂B₄O₇*H₂O, 0.0014 (NH₄)₆Mo₇O₂₄*4H₂O, 0.0023 ZnCl₂, 0.0021 CoCl₂*6H₂O, 0.001 CuCl₂*2H₂O and 0.003 MnCl₂*4H₂O. The medium with 75% v/v of wastes was not enriched with macro and micronutrients and consisted exclusively of non-diluted wastewaters and inoculum.

The experiments, run in 60 mL glass reactors (working capacity of 30 mL), were performed in five-fold repetitions. The samples were inoculated with 10 or 25% v/v of inoculum (biomass content in inoculum: 2.5 or 6.4 g VS/L, respectively). Initial pH was adjusted to 5.5. After deaeration of the samples with argon, all of them were sealed with butyl rubber membranes and incubated at 37°C with agitation (150 rpm). Medium containing 7g/L of glucose (standard medium) was applied as a reference.

In the photofermentation process distillery wastewaters replaced malic acid in modified Biebl and Pfennig medium and were introduced in 10, 20, 40, 60 or 70% v/v. The process was carried out in 25 mL vials (working capacity of 12.5 mL) made of borosilicate glass [19]. The medium was inoculated with 30% v/v of *R. sphaeroides* bacteria (0.36 g dry wt/L). Process was performed at $30\pm2^{\circ}$ C. The initial pH was set to 7.0. The medium with 70% v/v of waste was not enriched with macro and micronutrients and consisted exclusively of non-diluted waste and inoculum.

In the two-step system, the effluent originating from dark fermentation was centrifuged (12500 rpm), sterilized, flushed with argon and sealed with butyl rubber septa; the initial pH was adjusted to 7.0-7.2. The samples prepared in this way were used in the photofermentation reactor. A mercury-tungsten lamp (Ultra-Vitalux 300W from Osram) with light intensity of 116 W/m² at the surface of reactors was used in all the experiments.

Analytical Methods

Gases were collected with a gas-tight syringe over the surface of liquid. Gas chromatography (Varian 3800 equipped with TCD and capillary column CARBOPLOT P7) was applied for measurements of hydrogen and carbon dioxide content in the final biogas. The operational temperatures at the injection port, in the oven and at the detector were 120, 80 and 120°C, respectively. Argon was used as a carrier gas flowing with the rate of 8 mL/min.

The concentrations of volatile fatty acids (VFAs) and alcohols was analyzed with HPLC (Ultimate 3000 from Dionex, ThermoScientific, SHODEX sugar column SH1011, RI detector, 1 mL/min flow; eluent 5 mM H_2SO_4). Each sample was centrifuged for 15 minutes at 12500 rpm before the HPLC analysis. The loss of organic compounds (Δ COD) was measured with a dichromate method [22]. The amount of total solids (TS) and volatile solids (VS) were determined

according to the Standard Methods [22]. Ammonium ions concentration was measured applying MERCK spectroquant method (absorption at 690 nm).

The biomass content was established spectrophotometrically by measuring the optical density at 660 nm (DU640 UV-VIS spectrophotometer from Beckmann). Elementary analysis for C, H, N, S was performed by a Vario EL III elementary analyzer.

Substrate yield of hydrogen production was calculated as:

substrate yield =	wield -	amount of H2 produced in bioreactor (L)
	amount of wastewater in bioreactor (L)	

Specific yield of hydrogen production was calculated based on the following formula:

 $specific yield = \frac{amount of H2 produced (L)}{amount of COD removed (g COD)}$

Results and Discussion

The analysis of wastewaters showed high COD in sediments as well as in the solutions, so both fractions could potentially serve as the substrates for dark fermentation. HPLC analysis showed that the syrups did not contain glucose but contained glycerin, organic acids and alcohols in concentrations substantially higher than the stillages (Table 2). Organic acids cannot serve as the substrates in dark fermentation, whereas they are the effective substrates in photofermentation. On the other hand, glucose and glycerin can be successfully used in dark fermentation.

Hydrogen Production in Dark Fermentation

In the experiments with distillery wastewaters coming from alcohol production based on wheat, inoculum in concentrations of 10 and 25% v/v were tested and the latter led to higher hydrogen production, therefore it was used in further studies.

Fig. 1 shows hydrogen production for different initial waste concentrations in the media for all tested wastewaters. The reference test with medium containing only inoculum (without any other nutrients) showed hydrogen production of 0.02 mL/L, which is represented in Fig. 1 as 'inoculum'. This value is very low so does not contribute to overall hydrogen production.

It can be clearly seen that when WTL was used, hydrogen production was 3- to 4-fold higher than with other wastes. This effect is caused by the glucose content in WTL waste (there is no or a very low concentration of glucose in all other tested wastewaters – Table 2). Glucose is easily assimilated by bacteria from digested sludge, carrying out dark fermentation, and easier than glycerin and methanol (in which concentrations are high in the syrups). Even though theoretically 4 moles of hydrogen can be produced from 1 mole of glucose,



Fig. 1. Hydrogen production in dark fermentation carried out with distillery waste.

in practice maximum obtained values are around 2.7 moles, as glucose is also a growth substrate [5, 23]. WTL contains 1.5 g/L of glucose. When 75% v/v of this waste was used in the medium, 0.88 L H_2/L_{medium} was produced. Assuming that the average value of hydrogen produced from 1 mole of glucose is 2.7 moles, 0.38 L H_2/L_{medium} could be produced from glucose and the remaining 0.50 L H_2/L_{medium} from glycerin.

Previous studies have shown that the maximum consumption of glycerin by bacteria from digested sludge is 17 g/L [24]. Glycerin can be transformed to organic acids and alcohols in two pathways: oxidative, where pyruvate and afterwards formate, ethanol, butyric acid and acetic acid are formed; and reducing, where 1,3-propandiol is the only product [25]. In our studies both pathways can be observed (Table 3), and additionally glucose and lactose are transformed

to butyric and acetic acid as well as to ethanol and methanol.

Analysis of liquid metabolites, when WTL waste was used, showed that acetic, butyric and succinic acid as well as 1,3-propandiol were formed (Table 3). The final concentrations of these products indicated that they were formed mainly from the waste sediments (COD of liquid medium was only 38% of total COD of the waste). Glycerin and glucose contained in the liquid phase of the waste were completely consumed. However, lactic acid remained in the medium in up to 10% of initial concentration, while methanol was not consumed during the process. The concentration of succinic acid in the medium rose 5-fold and ethanol content increased over 3-fold.

Based on the above results, we can conclude that the main soluble substrates for microorganisms were glucose and glycerin, which were transformed to butyric and acetic acid as well as to 1,3-propanodiol. Analysis of the effect of waste content on hydrogen production showed that the amount of hydrogen increased significantly with the rising waste content in the medium in the range of 10-40% v/v (Fig.1, Table 4). A further increase of waste content resulted in only an insignificant rise of hydrogen production (up to 12 %). Therefore, the yield of the process (L H_2/L_{waste}) decreased when the waste content in the medium was over 40% v/v. Hydrogen content in final biogas varied from 48% v/v for MTL to 34% v/v for WSR (average value for dark fermentation around 50% [6, 26]).

The maximum amount of produced hydrogen (0.88 L H_2/L_{medium}) was observed when 75% v/v WTL was introduced to the medium, which represented 70% of hydrogen production in standard medium (1.2 L H_2/L from 7g glucose/L). Although four different distillery wastewaters were used in this study, only WTL gave

Table 3. Concentrations of main organic components in the liquid medium before and after fermentation of WTL waste with different dilutions.

	WTL content [% v/v]							
Metabolite	20%		40%		60%		75%	
[g/L]	Before	After	Before	After	Before	After	Before	After
Glucose	0.29	0	0.7	0	1.1	0	1.125	0.04
Lactic acid	0.44	0.04	1.05	0.05	1.5	0.08	2.0	0.4
Acetic acid	0.05	0.97	0.09	1.11	0.14	1.47	0.2	2.1
Butyric acid	0	0.73	0	1.2	0	2.13	0	2.7
Malic acid	0	0.03	0	0.11	0	0.13	0	0.2
Succinic acid	0.12	0.60	0.30	1.08	0.40	2.14	0.45	3.56
Glycerol	1.13	0.03	2.69	0	4.1	0	3.9	0
1,3-propanodiol	0	0.59	0	1.48	0	4.42	0	6.30
Methanol	0.48	0.61	0.61	0.39	1.42	1.58	2.47	2.32
Ethanol	0.02	0.07	0.03	0.03	0.05	0.18	0.08	0.37

Waste content [% v/v]	H ₂ [L/L _{medium}]	Substrate yield $[L H_2/L_{waste}]$	H ₂ [L/L _{medium}]	Substrate yield [L H ₂ /L _{waste}]
	W	TL	W	SR
10	0.22±0.02	2.2	0.15±0.02	1.5
20	0.43±0.15	2.18	0.13±0.02	0.63
40	0.82±0.10	2.05	0.10±0.01	0.25
60	0.85±0.09	1.42	0.09±0.01	0.15
75	0.88±0.08	1.17	0.10±0.01	0.13
	М	TL	MSR	
10	0.07±0.01	0.7	0.13±0.02	1.3
20	0.18±0.02	0.92	0.14±0.02	0.71
40	0.21±0.02	0.52	0.15±0.02	0.37
60	0.24±0.02	0.4	0.15±0.02	0.24
75	0.23±0.02	0.31	0.15±0.02	0.2

Table 4. Results of dark fermentation of wheat and maize distillery wastewater.

satisfactory results during the dark fermentation step. This showed that not all distillery wastewaters are a good substrate for biohydrogen production, because they can contain toxic compounds such as phenols and melanoidins, which inhibit dark fermentative bacteria [17, 20].

In our experiments, distillery wastewater with high COD content was used (at 75% dilution COD of WTL was 85 g O_2/L). The observed hydrogen production was 1.17 L/L_{medium} (equivalent of 7.4 mmol $H_2/$ g COD_{reduced}). For comparison, Mishra et al. [12] used distillery wastewater with COD of 40 g O_2/L and obtained 2.6 mmol $H_2/$ g COD_{reduced}, while Gadhe et al. [11] applied distillery wastewater with COD of 56 g O_2/L and observed maximum H_2 production of 5.45 mmol $H_2/$ g COD_{reduced}. This showed that in the case of WTL high hydrogen yields were obtained despite high COD value of the medium.

Hydrogen Production in Photofermentation

Media containing different concentrations of wheat and maize wastes (from 10-70%) were tested in the photofermentation process with *Rhodobacter sphaeroides* O.U.001. The results of this set of experiments are shown in Table 5.

In the case of WTL photofermentation, the amount of produced hydrogen reached the maximum when 20% v/v WTL was applied and did not change above this concentration (Table 5). This could have been caused by increased COD and produced biomass, which could lead to worse light access inside the bioreactor. However, the most efficient option, as far as produced hydrogen (1.1 L H_2/L_{medium}) and COD loss (5.79 g O_2/L) are concerned, was medium containing 40% of WTL.

Table 5. Results of photoferme	entation with wheat	t and maize distille	ry wastewater.
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Waste content [% v/v]	H ₂ [L/L _{medium}]	Biomass [g/L _{medium}]	Substrate yield $[L H_2/L_{waste}]$	H ₂ [L/L _{medium}]	Biomass [g/L _{medium}]	Substrate yield $[L H_2/L_{waste}]$
		WTL			WSR	
10	0.83±0.06	1.54±0.17	8.3	1.46±0.15	1.85±0.17	14.6
20	1.06±0.20	1.70±0.19	5.0	1.72±0.18	2.02±0.18	8.6
40	1.10±0.22	1.94±0.21	2.8	1.73±0.20	2.17±0.19	4.3
70	1.00±0.07	1.90±0.17	1.4	0.64±0.04	2.47±0.27	0.9
		MTL			MSR	
10	0.63±0.07	1.56±0.14	6.3	1.67±0.23	1.96±0.19	6.7
20	1.03±0.12	1.86±0.20	7.6	1.41±0.22	2.04±0.18	7.0
40	1.09±0.09	1.82±0.20	2.7	1.37±0.16	2.42±0.29	3.4
70	1.03±0.03	1.98±0.16	1.5	1.30±0.03	1.98±0.13	1.5

When WSR was applied, the maximum hydrogen production was observed when 20 and 40% v/v of WSR were introduced to the medium and with 70% v/v WSR the amount of generated hydrogen dropped by over 60% (Table 5). Similarly as for WTL, increasing COD-produced biomass as well as increasing methanol content could be the reasons [27]. In conclusion, the most efficient option as far as produced hydrogen (1.73 L H_2/L_{medium}) and COD loss (21.6 g O_2/L) are concerned, was diluted medium contained 40% of WSR.

In most cases the distillery wastewaters need to be diluted before they can be used for photofermentation – mainly due to high concentrations of ammonium ions [15], which were found to inhibit nitrogenase and hence hydrogen production and increase the biomass concentration at the same time. Furthermore, high biomass content additionally reduces hydrogen production as it limits light penetration in the reactor. On the other hand, nitrogen limited conditions also reduce hydrogen production, and therefore it is important to achieve the balance between the amount of nitrogen necessary for bacterial growth and the amount that causes the reduction of hydrogen production.

Substrate yield and specific yield dropped significantly with rising waste content in the medium for both WTL and WSR. However, substrate yields were higher for photofermentation than for dark fermentation. Organic acids contained in the wheat wastes were more effective substrates in photofermentation than glucose and glycerin for dark fermentation. Hydrogen content in final biogas was in the range of 84-87%, except when 70% v/v WSR was applied, where hydrogen content dropped to 65%.

When MTL (Table 5) was applied, hydrogen production reached the maximum when 20% v/v MTL was applied and did not change above this concentration (similarly to WTL). The most efficient option, as far as produced hydrogen (1 L H_2/L_{medium}) and COD loss

(5.88 g O_2/L) are concerned, was medium containing 70% of MTL.

When MSR (Table 5) was applied, hydrogen production gradually decreased with rising waste content in the medium (in the range of 10-70%). The reason for this effect could be very high COD in the crude waste (325 g O₂/L) and high methanol content (11.45 g/L). The highest hydrogen production was observed when MSR waste was 10-fold diluted $(1.67 \text{ L H}_2/\text{L}_{\text{medium}})$. High content of lactic acid in crude waste (13.4 g/L) enabled high hydrogen production with only 10% of MSR in the medium. The highest COD loss (34.3 g O₂/L) was obtained for 70% v/v MSR with simultaneous high hydrogen production -1.03 L H₂/L_{medium}. hydrogen content in final biogas was in the range of 80-87%, except when 10% of MTL was applied, where hydrogen content dropped to 58%.

Conversion of organic compounds in both wheatand maize-originating wastes in photofermentation are shown in Fig. 2. The complete transformation of succinic, lactic, acetic and malic acid was observed. Glucose, contained only in WTL, was 27% consumed. Glycerol concentration decreased in wheat wastes by 9-10% and in maize wastes remained almost unchanged. Methanol conversion in all four wastes was insignificant.

Two-Step System

Taking into account the content of the crude wastes and the products of dark fermentation, the two-step system, based on dark fermentation and photofermentation, was applied.

In the first step, where dark fermentation took place, hydrogen was produced from glucose and glycerol, which were 100% consumed. Simultaneously acetic, butyric, succinic and malic acid as well as 1,3-propandiol were formed, which served as the



Fig. 2. Conversion of organic compounds during photofermentation: a) wheat distillery wastewater and b) maize distillery wastewater.

Table 6. Hydrogen production in the two-step system; effluent from dark fermentation was 5-fold diluted before the photofermentation process.

	Dark fermentation I st step	Photofermentation II nd step	Total			
		[L H ₂ /L _{medium}]				
75% v/v WTL	0.88	0.66	1.54			
75% v/v MTL	0.23	0.48	0.71			
		[L H ₂ /L _{waste}]				
75% v/v WTL	1.17	3.30	4.47			
75% v/v MTL	0.31	2.40	2.71			

substrates for hydrogen production in photofermentation (second step of the hybrid system).

WTL and MTL led to the highest hydrogen production in dark fermentation and therefore these wastes were introduced to the medium (both in concentrations of 75% v/v) in the first step. The effluent from the dark fermentation reactor was inoculated with *Rhodobacter sphaeroides* O.U.001 in 30% v/v. The experiments showed that effluent had to be diluted 5-fold in order for hydrogen production to take place. The reason for that could be the content of ammonium ions in the effluent: 1.08 g/L coming from the digested sludge and 0.19 g/L originating from the wastes. Sabourin-Provost et al. described the inhibiting effect

of ammonium ions on photofermentation, proving that already as small a concentration as 2 mM decreases hydrogen production [28, 29]. Other studies with distillery wastewaters in the two-step hybrid system showed that the effluent from dark fermentation has to be diluted at least five times before introducing it to the photofermentation reactor [17, 19]. The experiments with higher concentrations of the effluent from dark fermentation led to lower bacterial growth, hydrogen production and COD reduction. Laurinavichene et al. [19] reported maximum efficiency of hydrogen production (17.6 L H₂/L_{waste}) with 5-fold diluted effluent, while Lazaro et al. [17] used 10-fold dilution to obtain maximum efficiency (5.5 mmol H₂/g COD_{reduced}).

Average hydrogen production in dark fermentation of 75% v/v WTL was 0.88 L H₂/L_{medium}, while in the second step (photofermentation) additionally 0.66 L H₂/L_{medium} was generated. Therefore, the total amount of produced hydrogen in a two-step system was 1.54 L H₂/L_{medium} (4.47 L H₂/L_{WTL}, Table 6). When MTL in 75% v/v was applied, 0.23 L H₂/L_{medium} was produced in dark fermentation and 0.48 L H₂/L_{medium} in photofermentation, which gave a total amount of 0.71 L H₂/L_{medium} (2.71 L H₂/L_{MTL}).

In the separate processes of dark and photofermentation of WTL, 1.17 L H_2/L_{WTL} and 1.4 L H_2/L_{WTL} were produced, respectively (2.57 L H_2/L_{WTL} in total). Whereas in a two-step system 4.47 L H_2/L_{WTL} was produced in total, which was nearly 2 times higher than taking together the amount generated in two separate processes. When MTL was used for dark and photofermentation in the separate processes, total hydrogen production was 1.81 L H_2/L_{MTL} (0.31 L H_2/L_{MTL} in dark fermentation and

Table 7. Examples of two-step systems in microbiological hydrogen production from waste.

Bacteria used in dark fermentation	Bacteria used in photofermen- tation	Origin of waste (main substrates)	Main substrates in photofermentation	Max H ₂ production	Ref.
Heat-treated anaerobic digested sludge	Rhodobacter sphaeroides O.U.001	Waste from chewing gum production (glycerol, Talha gum)	Xylitol, VFAs	6.7 L/L	[30]
Heat-treated anaerobic digestion sludge (dominant strain <i>Clostridium butyri-</i> <i>cum</i>)	Mixed photosynthetic bacteria (dominant strain <i>Rhodopseu- domonas palustris</i>)	Cassava ethanol wastewater	Acetate, butyrate	5.2 L/L	[31]
Clostridium butyricum LS2	Rhodopseudomonas palustris (1.8929)	Palm oil mill effluent	VFAs (butyrate, acetate, propionate), ethanol	3.1 L/L	[32]
Heat-treated sewage sludge	Mixed bacterial culture HAU- M1	Hydrolyzed corn stover	VFAs	7.0 L/L/d	[33]
Saccharolytic consortium (silage pit liquid)	Rhodobacter sphaeroides B-3059, Rhodobacter capsu- latus B10	Distillery wastewater (saccharides, proteins, VFAs - mainly lactate)	VFAs (butyrate, ac- etate, propionate)	17.6 L/L	[19]
Caldicellulosiruptor sac- charolyticus	Rhodopseudomonas palustris, Rhodobacter capsulatus	Sugar beet molasses (sucrose)	Acetate, lactate, ethanol	3.0 L/L	[34]
Heat-treated anaerobic digested sludge	Rhodobacter sphaeroides O.U.001	Distillery wastewater (glucose, glycerol)	VFAs (succinate, butyrate, acetate), ethanol, methanol	4.47 L/L	[this study]

1.5 L H_2/L_{MTL} in photofermentation). In the two-step process H_2 production reached 2.71 L H_2/L_{MTL} . In the case of both WTL and MTL, total hydrogen production was significantly higher in the two-step system than for the separate processes [19, 30-34].

Table 7 presents the results on hydrogen production for two-step systems for different wastes with various substrates obtained by other researchers. It is difficult to compare the results when different substrates and different bacterial cultures were applied. However, we can refer to the systems where the dominant strain was R. palustris and acetate with butyrate were the main substrates for the photofermentation step [31, 32, 34]. In these systems, hydrogen production was in the range of 3.0 L/L to 5.2 L/L, while 4.47 L/L was obtained in our experiments. Lin et al. [31] using mixed bacterial cultures and cassava ethanol wastewater as a substrate obtained 0.43 L H_2/L and 4.77 L H_2/L in the dark and photo steps, respectively (5.2 L H_2/L in total). High hydrogen production in the second step could be explained by high concentration of VFAs in dark fermentation effluent. A similar effect is observed in our experiments. In turn, Chandra et al. [18] obtained 1.51 L H₂/L from distillery wastewater during one step of hybrid dark-photo fermentation with mixed bacterial cultures as inoculum.

It is known that in dark fermentation, sugars are converted to VFAs, mainly to acetic and butyric acids [35]. High hydrogen production is observed when the ratio HAc/HBu is high, namely when the acetic acid pathway is dominant [36]. Butyric and propionic acid pathways cause a decrease of hydrogen production [17]. In our experiments the main VFAs were acetic and butyric acids with an HAc/HBu ratio varying between 1.3:1 to 0.7:1 (Table 3). Wang et al. [37] showed that hydrogen production rates and substrate conversion efficiency during photofermentation were improved for mixtures of acetic acid and butyric acids compared to processes with a single substrate. Therefore, mixtures of acetate and butyrate obtained after dark fermentation are beneficial for the second step of the hybrid system.

Indeed, an important advantage of the two-step process was a high utilization of organic compounds contained in the wastes, and also those formed in dark fermentation (1,3-propandiol, succinic, butyric, acetic and malic acid). When two separate processes of dark and photofermentation took place, 1,3-propandiol, organic acids, methanol and ethanol remained in the medium after dark fermentation, whereas glucose, glycerol and methanol remained in the medium after photofermentation. In the two-step system all the above-mentioned organic compounds were 100% utilized, except ethanol and methanol, which were 18.9% and 9.5% utilized, respectively (Fig. 3). Similarly, Lin et al. [31] and Laurinavichene et al. [19] obtained 99% utilization of VFAs during the two-step process with distillery wastewater as a substrate. However, only 78% reduction of butyrate and increased acetate



Fig. 3. Conversion of organic compounds in two-step system with WTL as a substrate.

concentration was observed in other two-step process with distillery wastewater [17], which was explained by the presence of toxic compounds and contamination by sulfur-reducing bacteria.

Combined biohydrogen production systems from distillery wastewaters presented by different studies (also single-step [18]) showed improved results compared to separate dark and photofermentation: higher hydrogen production as well as higher COD and VFA removal. Therefore, optimization of hybrid systems seems to be essential for application on a large scale. Our experiments were carried out in small bioreactors, and appropriate scale-up would be necessary to make the process economically viable. Most dark fermentation studies have been carried out at laboratory scale [38]. Limited studies have been performed on pilot-scale applications. More research on scalability is essential to provide information on feasible operating conditions, economic viability, reactor and plant design, as well as waste management. Dark fermentation processes have a prospect of being rapidly scaled-up owing to its similarity to the wellknown anaerobic digestion process. Recent studies on scaling-up dark fermentation usually showed showed only a slightly lower yield when compared to laboratory scale [38]. Considering the photofermentative process, one of its main disadvantages is the more complex design of reactors, owing to the need to maintain a suitable proportion between reactor surface area and volume when scaling-up. Moreover, the self-shading effect of bacteria cultures would affect abundance and homogeneous distribution of light. However, even in the case of photofermentation, efficient hydrogen production pilot bioreactors were presented [39].

Conclusions

Among four different distillery wastewaters only WTL gave high hydrogen yields, indicating that not all distillery wastewaters are suitable for dark fermentative hydrogen production. Based on our experiments, stillages were more effective substrates than syrups in dark fermentation as far as hydrogen production is concerned (by 80% in the case of wheat wastewater and by 40% in the case of maize wastewater). In contrast, the application of syrups during photofermentation led to 40% higher hydrogen production and over 70% higher COD loss in comparison with processes where stillages were used.

The combination of both dark and photofermentation in the two-step system gave two main advantages: 2-fold higher hydrogen production than for the separate processes and complete utilization of organic compounds contained in the wastes (glycerol, glucose and lactic acid) together with compounds formed in dark fermentation (1,3-propandiol, succinic, butyric, acetic and malic acid). The exception was ethanol and methanol, which were 18.9% and 9.5% utilized, respectively. Total hydrogen production in the two-step system reached 4.47 L H₂/L_{WTL} and 2.71 L H₂/L_{MTL}.

Acknowledgements

This work was financed by the National Science Centre (grant No. UMO-2017/26/D/ST8/00149).

Conflict of Interest

The authors declare no conflict of interest.

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