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Review

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Reactor systems for syngas fermentation processes: a review

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Highlights

- Operation principles of the main bioreactor configurations in syngas fermentation
- Operational parameters maximizing bioreactors' productivities
- Comparison of the mass transfer efficiency of different bioreactor setups
- Current status in commercialization of syngas fermentation

Abstract

Implementation of biofuels as an alternative to fossil fuels has been established as an answer to climate change by limiting GHG emissions. Syngas fermentation has emerged as a promising process for the conversion of waste biomasses to valuable products with bioethanol being on the main focus. However, the bottleneck of the mass transfer of syngas compounds H₂ and CO along with low production yields has set barriers to the development of an industrial scale plant. Recent research indicates that many different methodologies spring up in order to face this important challenge. The aim of this review is to assemble all these techniques applied in syngas fermentation, focusing on the different bioreactor configurations

operated in continuous mode for the production of liquid and gas biofuels. This article also outlines the so far entrepreneurial initiatives and the progress made towards the commercialization of the process.

Keywords: Synthesis gas, Syngas Fermentation, Bioreactor Configurations, Biochemicals, Biofuels, Mass Transfer

1. Introduction

The continuous increase of global energy needs due to the constant rise of global population and the intensification of industrial activities renders the use of renewable energy necessary [1,2]. Not only because of the gradual depletion of fossil fuel reserves, but also because humanity has to confront the major problem of the greenhouse effect and the resulting global warming [3–5]. Towards this direction, syngas fermentation to biofuels has gained increasing scientific attention as an alternative methodology for the production of renewable energy over the last decade [6–17].

Syngas or synthesis gas is a mixture of gases (mainly CO, CO₂ and H₂) that can be produced from the gasification of biomass. The proportion of each gas depends on the type of the biomass used as a feedstock, the configuration of the gasifier and the operational parameters of the gasification process [18]. The industrially used gasification technology primarily includes fixed bed and fluidized bed gasifiers and the gas effluent often undergoes additional cleaning processes before it is fed to a reactor [19]. There is also on-going research on the optimization of the gasification process and novel technologies are often introduced [20–23].

Furthermore, syngas is quite enthralling from an environmental point of view because to-date it remains an important by-product of several industrial processes such as steel milling, petroleum refining, steam reforming and others involving combustion and partial oxidation. European Union has set a long-term target (2050) to decrease its greenhouse gas emissions by 80-95% compared to 1990 [24] and in order to achieve that, exploitation of inorganic carbon sources is bound to have a fundamental role. Integration of

syngas producing industrial activities and syngas fermentation units in a pre-commercial stage has already been attempted for bioethanol production in China [25].

Conventional catalytic processes for the production of liquid fuels like Fischer-Tropsch (FT) present high operational costs as they require high temperatures and pressures, high supply cost for the catalysts, fixed H_2/CO ratios and pretreatment of the gaseous mixture for the removal of compounds that are poisonous to the catalysts [26]. Those disadvantages highlight the need for alternative ways to be sought. One promising alternative is the combination of thermochemical and biological processing of biomass [27]. The greatest merits of the fermentation processes are the mild conditions needed that entail low energy and infrastructure costs along with the high selectivity of the microbes. Apart from that, the biocatalysts (microbes) are cheap, they do not demand fixed H_2/CO ratio and the process is odorless. In addition, health hazards are precluded and environmental pollution is markedly abated [28]. However, syngas bioconversion to liquid fuels faces important challenges that should be circumvented before the process is scaled up. The main issues are the mass transfer of sparingly soluble syngas compounds (CO, H_2) to the water-based microbial cultures and the relatively low growth rate of the microbes that leads to relatively low productivity rates [29].

The bottlenecks of syngas fermentation have been perceived since three decades ago but, because of the use of alternative methodologies for the production of liquid fuels such as fermentation of sugars or gasification and FT, little effort was given to surpass them up to the last decade. Concisely, the bioreactor setups for syngas fermentation that are dominant in the literature are continuous stirred tank reactors (CSTR) and membrane reactors. However, often new ideas spring up due to limitations and challenges those two main configurations have. CSTRs demand a lot of energy consumption for high agitation speeds so as to increase gas-to-liquid mass transfer [30] and, on the other hand membrane bioreactors, where biofilm formation takes place, face fouling issues from high cell concentrations and cell washout phenomena at high hydraulic dilution rates [31].

A key targeted product of the combined biomass gasification – syngas fermentation process is bioethanol due to its high octane number that allows it to be used in fuel blends in the transportation sector. Today bioethanol is produced in a commercial scale through the fermentation of carbohydrates deriving mainly from corn, sugarcane, sugarbeet and wheat. It is, however, highly debated that these crops, which can also be a source of food in a world that has not eliminated hunger, should play a major role in bioethanol production. This conflict rose extensive research towards second generation bioethanol (2GB) produced from lignocellulosic biomass that cannot be used as a food source [32]. The key step for the production of 2GB is the pretreatment of the biomass, so as to break down the lignocellulosic structure and make cellulose and hemicellulose more amenable to the subsequent enzymatic hydrolysis step. Dependent on the type of the biomass many different pretreatment approaches have been studied (most common are: acid pretreatment, alkaline pretreatment, wet oxidation, organosolv pretreatment, ozonolysis, steam explosion, ammonia fiber explosion and biological treatment) each of them facing different economic and environmental drawbacks [33]. The main advantage of the thermochemical - biological processing is that the whole organic material of the lignocellulosic biomass (including lignin and hemicellulose) can be converted to hydrogen, carbon dioxide and carbon monoxide and then fermented to bioethanol, whereas in 2GB production through hydrolysis, carbon in lignin and in a fraction of hemicellulose (25 – 30% of the total feedstock carbon) cannot be utilized resulting thus in lower yields. In addition to that, the syngas fermentation platform is more flexible since it can process simultaneously waste gases from other sources besides lignocellulosic biomass [34].

Besides ethanol, other important products from the fermentation of syngas, are methane [8], acetic acid [10] and higher alcohols, i.e. butanol [35–37] and acids [11]. Methane is a valuable molecule for energy storage and can be utilized as a fuel for heat and electricity production as well as in Natural Gas Vehicle in the transportation sector. Acetic acid is an important precursor for the production of adhesives, inks, paints and coatings. It has also medical applications, for example, as an antiseptic against pseudomonas infections. Higher carbon chain VFAs such as propionic acid and butyric acid are used industrially as building

blocks for the production of high added value chemicals and as additives for the preservation of food while butanol is mainly used as a solvent in many industrial applications. The aforementioned chemicals are principally produced chemically from fossil resources and, thus, there is high scientific interest on the possibility of sustainable biological production through syngas fermentation.

The aim of this review is to present the so far reported bioreactor configurations operated in continuous mode for the bioconversion of syngas to the abovementioned added value products and compare their effectiveness in achieving high productivity rates and high product concentrations.

2. Basic Reactor Principles

One of the most critical decisions to be considered for the growth of microorganisms is the configuration of the reactor that will be used as well as the operation mode of the process (continuous, batch, semi-continuous, combinations of processes). In an industrial scale several factors affect the final choice of the reactor and the operation mode, i.e. secondary products that may require the microorganisms to be in a stationary phase, genetic instability, toxicity of the desired product which results to inhibition of the microbes [38,39].

The basic principle of stirred tank reactors is the agitation that leads to uniform conditions of concentration and temperature throughout the reactor volume. One agitator or more are mounted on a shaft which is connected to a gear box and a motor, used for the manipulation of the rotational speed. The power input for the rotation of the agitator is a crucial economic parameter that determines the economic sustainability of the bioprocess [40].

Bubble columns reactors (BCR) are cylindrical vessels, which are filled with a liquid phase, and a gas phase is supplied at their bottom. For the design of a bubble column attention should be paid to two factors: diameter to length ratio and the type of the gas sparger forming the bubbles. The gas is spread in the liquid volume of the column through convective flow which is driven by the incremental density differences

[41,42]. An additional reactor configuration is the gaslift. Depending on their structure, gaslift reactors can be divided in two main categories; external loop and internal loop. The external loop airlift reactors consist of distinct conduits and the fluid flow is circulating from the one conduit to the other. On the other hand internal loop airlift reactors consist of concentric tubes and the fluid flow is circulating from the inner to the outer tube. In both configurations gas is sparged from the bottom of the reactor [42,43].

Hollow fiber membrane reactors (HFR) are gas to liquid transfer membrane systems providing a high specific surface area. They can also serve as support for microbial growth and biofilm formation but special attention should be given to the thickness of the biofilm because it can also have a negative effect when microbial growth is high enough [44,45]. This phenomenon is called biofouling and a lot of research is currently applied towards illuminating its causes, so as to design more efficient reactors where biofouling can be limited and controlled [46,47]. In order to mitigate biofouling in membrane bioreactors several strategies have been attempted including both physical and chemical methods[48]. Depending on the application, the membranes can be either submerged to the liquid medium in the reactor or externally connected in series with the reactor [49–51]. Applications of various types of hollow fiber membrane modules have been reviewed by Kumar et al. [52].

Trickle bed reactors (TBR) consist of a packed bed column on which biofilm grows and gas is flowing co-currently or counter-currently to the liquid [53]. The name of this type of reactors comes from the trickling of the liquid medium through the pores of the inert material and through the gaps amongst the inert material [54]. This kind of setups has been used for degradation of pollutants such as hydrogen sulfide, volatile organic compounds (VOCs), dichloromethane and ammonia [55–59].

3. Mass Transfer Calculations under Abiotic Conditions

Several bioreactor configurations can be found in the literature tested under abiotic conditions for the calculation of the mass transfer coefficient (K_La) of syngas compounds to water. However, any comparison

of the values of mass transfer coefficients reported in different publications should be done with caution because of the different methodologies used for the calculation of K_La , the different ways of volume calculation (working volume, total volume, packed bed volume) and the different operational parameters tested [60]. A collection of reported K_La values and the operational conditions under which they were achieved can be seen in table 1.

One of the highest reported values of K_La is 946 h^{-1} for a composite hollow fiber (CHF) membrane module made of polyethylene and polyurethane [61]. The inlet was pure CO at a flow of $500 \text{ mL} \cdot \text{min}^{-1} \cdot L_{RWV}^{-1}$ (RWV = Reactor Working Volume) and the CHF module was operated without a gas outlet. In a similar configuration with hydrophobic polypropylene hollow fiber membranes, K_La for CO was calculated at 385 h^{-1} when a syngas composition of 50% CO, 30% H_2 and 20% CO_2 , and an inlet flow of $55.34 \text{ mL} \cdot \text{min}^{-1} \cdot L_{RWV}^{-1}$ were applied [50]. It was also shown that higher gas supply rates, higher liquid recirculation rates and a bigger number of fibers result to the improvement of the mass transfer rate. When experiments were carried out with a gas-lift reactor coupled with a $20 \text{ }\mu\text{m}$ bulb diffuser and pure CO in the inlet at a flow of $1667 \text{ mL} \cdot \text{min}^{-1} \cdot L_{RWV}^{-1}$ the K_La value was 129.6 h^{-1} for CO [62]. In a previous experimental study, Munasinghe and Khanal reported values ranging from 0.4 to 91 h^{-1} for eight different bioreactor configurations tested under the same conditions [63].

Orgill et al. [60] compared the mass transfer coefficients of a trickle bed reactor, a hollow fiber membrane reactor and a stirred tank reactor under various gas flow and liquid flow rates. In that study the gas applied was air and the calculation of the K_La was determined through the %DO (dissolved oxygen) in the liquid phase [60]. Regarding the trickle bed reactor, two different sizes of packing material were assessed (3 mm and 6 mm soda lime glass beads) and for the hollow fiber reactor, the effect of five different hollow fiber modules on mass transfer was examined. The maximum mass transfer values of the TBR, the HFR and the CSTR were 421, 1062 and 114 h^{-1} respectively. Finally, for a study regarding the type of the impeller

employed in a CSTR the outcome was that improved mass transfer without increased power supply can be achieved with dual impeller schemes with an axial flow impeller as the top impeller [64].

Table 1. Mass transfer coefficient K_La calculated under abiotic conditions for several bioreactor configurations and operational parameters

Reactor Configuration	Gas Composition	Working Volume (L)	Packing or Hollow Fiber Material	Gas Flow (mL·min ⁻¹)	Agitation (rpm)	K _L a (h ⁻¹)	References
Hollow Fiber Membrane (HFMBR)	50% CO, 30% H ₂ , and 20% CO ₂	HFM 0.13 , Reservoir 2.4	Hydrophobic polypropylene	140	90 (reservoir)	385 CO	[50]
Composite Hollow Fiber Membrane (CHF module)	99.99% CO	3	MHF0504MBFT (polyethylene and polyurethane)	1500	NA	946	[61]
Gas-Lift	>99.99% purity of CO and H ₂ in each experiment	3	NA	5000	NA	129.6 CO, 97.2 H ₂	[62]
Column with Column Diffuser	99.99% purity of CO	3	NA	5000	NA	40	[63]
Column with 20-μm Bulb Diffuser					NA	70.8	
Column only with Sparger					NA	50.4	
Column with Sparger with Mechanical Mixing					300	55.8	
Column with Submerged Composite Hollow Fiber Membrane					NA	1.1	
Air-Lift Combined with a 20-μm Bulb Diffuser					NA	91.1	
Air-Lift Combined with Single Point Gas Entry					NA	45	
Trickle Bed	air	1 (packed bed volume)	3 mm soda lime glass beads	130.9	NA	171	[60]
			6 mm soda lime glass beads	106.4		421	
5 (water holding tank)		Hydrophilic porous polystyrene	1000-2000 (effect of gas flow was considered negligible)	NA	55		
		Hydrophilic porous polyether sulfone			20		
		Hydrophobic porous polypropylene			240		
		Hydrophobic non-			1062		

			porous polydimethylsiloxane				
			Hydrophilic porous fresenius polysulfone			40	
CSTR		3 (glass vessel)	NA	400	900	114	

*NA – Not Applied

4. Continuous Processes

4.1 Production of Ethanol and/or Acetate

Ethanol constitutes the main targeted product of syngas fermentation because of its availability for a wide spectrum of applications, hence several factors of the fermentation process have been studied and are in depth reviewed [15,27,28,34,65–67]. In short, the most crucial ones are: bioreactor configuration and operational parameters such as pressure, temperature and pH. In addition the yield and the productivity of ethanol can be affected by the type of liquid media used, surfactants, even the use of nanoparticles [68–77]. Regarding the effects of temperature on syngas fermentation, there has recently been an effort to enhance alcohol productivity and avoid acid crash phenomena by applying lower values than the optimum ones with positive results in batch experiments[78].

The majority of bioreactor configurations tested for the production of ethanol, or acetate as a precursor, is stirred tank vessels. *Peptostreptococcus productus* was one of the first bacteria ever studied for the fermentation of syngas compounds to liquid fuels [79–81]. The maximum productivity of acetate measured was $4.8 \text{ mmol} \cdot \text{L}_{\text{RWV}}^{-1} \cdot \text{h}^{-1}$ with an agitation speed of 400 rpm and a gas flow of $19.54 \text{ mL} \cdot \text{min}^{-1} \cdot \text{L}_{\text{RWV}}^{-1}$ [79]. Those results were promising and advocated to the continuation of experimental research over the subject. However, to our knowledge no other studies with *P. productus* (later classified as *Ruminococcus productus* [82] and finally as *Blautia producta* [83]) can be found in the literature for continuous syngas fermentation. The first pH based study for syngas fermentation was performed by Grethlein et al. [84] with bacterium *Butyrivacterium methylotrophicum* in a CSTR set-up. The scientists observed that lowering the pH resulted in an alteration of the metabolism of CO towards more reduced products, which is quite critical since a route to alcohols' production opened. Researchers had earlier managed to produce acetate and butyrate with *B. methylotrophicum* [85–87].

Another landmark in the research of syngas fermentation is the introduction of microbubble sparging [88,89]. The researchers designated the augmented mass transfer of CO into the liquid phase when the gas phase was sparged in the form of microbubbles with a 5.5 times increase of the K_La value compared to conventional sparging (91 h^{-1} and 14 h^{-1} respectively). This was a rational observation since smaller bubbles have more beneficial surface/volume ratio [90]. The operational parameters were the same for both experiments and only the effect of the type of the sparger was tested. Acetate productivity from *Butyribacterium methylotrophicum* was measured at $2.71 \text{ mmol} \cdot \text{L}_{\text{RWV}}^{-1} \cdot \text{h}^{-1}$ which was less than the one with *P. productus*. A schematic of the bioreactor set-up, as it was reported from Bredwell et al. [88], is presented in Figure 1.

Figure 1. CSTR set-up with external microbubble generator [88]. The microbubble generator contained a spinning disc rotated at 4000 rpm for the microbubble formation.

Chang et al. [75] used a bioreactor configuration (Figure 2) made up from a bubble column reactor and a hollow fiber membrane module as a cell recycle system. The purpose of the study was to ascertain *Eubacterium limosum* as a means of syngas conversion to liquid fuels under several CO partial pressures.

The acetate productivity reached almost $3.65 \text{ mmol} \cdot \text{L}_{\text{RWW}}^{-1} \cdot \text{h}^{-1}$ and K_{La} was calculated at 72 h^{-1} with a CO partial pressure of 41.5 kPa .

Figure 2. Schematic bioreactor set-up used by Chang et al. [75] for CO fermentation. CO was fed from the bottom of the bubble column with a gas recycle loop recirculating the non-converted CO in the top. The liquid media were stored in a reservoir vessel flowing into the bubble column from the top. Liquid recirculation was also employed with a hollow fiber membrane module.

A different approach to the bioconversion process of syngas to ethanol was given by Kundiyana et al. [77] and Richter et al. [91] by suggesting a two stage system where in the first stage microbial growth coupled to acetogenesis occurs while in the second phase solventogenesis takes place. Kundiyana et al. [77] used two CSTR fermentors connected in series and installed a hollow fiber membrane module for cell recycle in each fermentor. The flow of syngas was tested both in a co-current and a counter-current orientation. In the first case syngas was flowing from stage 1 reactor to stage 2 and in the second from stage 2 to stage 1.

Conversely, Richter et al. [91] developed a setup with a CSTR and a bubble column reactor applying also a cell retention system and a gas recycle loop (Figure 3). Ethanol production was taking place in the bubble column at a productivity of $8.04 \text{ mmol} \cdot \text{L}_{\text{RWW}}^{-1} \cdot \text{h}^{-1}$ and a concentration of $20.7 \text{ g} \cdot \text{L}^{-1}$ which is promising for this kind of systems as this value is much higher than single CSTR fermentation with *C. ljungdahlii* reported by Mohammadi et al. [92] and Younesi et al. [93] who achieved maximum titer of $6.5 \text{ g} \cdot \text{L}^{-1}$ and $11 \text{ g} \cdot \text{L}^{-1}$,

Figure 3. Two stage syngas fermentation to ethanol scheme by Richter et al. [91]. Solid lines represent the flow of the liquid media and dotted lines the flow of the gas phase. Abbreviations: Ag, agitation; BP, bypass; E, effluent reservoir; Ex, exhaust; FT, foam trap; G1, G2, gas recycle loops; HF, hollow fiber module for cell recycle; M, media reservoir; Per, permeate; Ret, retentate. Liquid media flowed to the CSTR from a reservoir vessel and subsequently were pumped to the bubble column. Recirculation of the liquid media was applied through a hollow fiber membrane module. The gas was sparged from the bottom of the bubble column and was recirculated for better conversion rates. Similarly, syngas was flowing to the agitated vessel.

respectively. However the researchers reported 2% ethanol content in the effluent which is relatively low and demands advanced distillation techniques.

In order to overcome mass transfer limitations, Shen et al. [94–96] tested three biofilm formation modules in regard to their K_La and ethanol productivity. The syngas composition (20% CO, 5% H₂, 15% CO₂, and 60% N₂) and the biocatalyst (*Clostridium carboxidivorans* P7) was the same for all the experiments whereas the operational parameters were optimized for each case. First, a hollow fiber membrane module made of microporous hydrophobic polypropylene was tested [94]. The module was not submerged within the bioreactor, but instead, connected to an agitated reservoir vessel. Ethanol productivity was $3.12 \text{ mmol} \cdot \text{L}_{\text{RWV}}^{-1} \cdot \text{h}^{-1}$ with an ethanol to acetate ratio of 4.79 when the syngas flow, the liquid recirculation and the dilution rate were $37.5 \text{ mL} \cdot \text{min}^{-1} \cdot \text{L}_{\text{RWV}}^{-1}$, $25 \text{ mL} \cdot \text{min}^{-1} \cdot \text{L}_{\text{RWV}}^{-1}$ and 0.96 d^{-1} respectively. Second, the author employed a cordierite-based ceramic monolith cylinder housed in a plexiglass column connected with an agitated reservoir vessel akin to the previous set up [95]. The confinements in the range of values of the operational parameters in this case are due to unwanted abrading and biofouling phenomena taking place in the biofilm. The maximum ethanol productivity of $2.13 \text{ mmol} \cdot \text{L}_{\text{RWV}}^{-1} \cdot \text{h}^{-1}$ was achieved at $37.5 \text{ mL} \cdot \text{min}^{-1} \cdot \text{L}_{\text{RWV}}^{-1}$ syngas flow rate and 0.48 d^{-1} dilution rate. Ethanol to acetate ratio was measured at 2.1. A schematic of the reactor configuration is presented in Figure 4. The third set up was different from the previous ones: a horizontal mesh cage packed with AnoxKaldnesTM carriers was placed into a glass tank and was rotated by a gearmotor [96]. The rotated packed bed bioreactor provided a 45% cage submersion when the cage was in its vertical state to the surface of the liquid. Syngas was entrained in the liquid media through a submerged gas distributor in the bottom of the vessel. While the rotational speed of the gearmotor was 50 rpm, the headspace absolute pressure 29.7 psi and the dilution rate 0.96 d^{-1} , the productivity of ethanol reached $6.06 \text{ mmol} \cdot \text{L}_{\text{RWV}}^{-1} \cdot \text{h}^{-1}$ which was the highest amongst the configurations tested with the headspace pressure being a parameter affecting the productivity to a significant degree.

Table 2. Bioreactor configurations for the production of ethanol and acetate

Reactor Configuration	Gas Composition	Working Volume (L)	Packing or Hollow Fiber Material	Gas flow (mL·min ⁻¹)	Agitation (rpm)	K _L a (h ⁻¹)	Microorganisms	Temperature (°C)	Product	Productivity / Titer	References
CSTR	63.43% CO, 15.96% CO ₂ and 20.61% CH ₄	0.325	NA*	6.35	400	36.6 h	<i>Peptostreptococcus productus</i>	37	acetate	4.8 mmol·L _{RWV} ⁻¹ ·h ⁻¹	[97]
CSTR	conventional sparging 20% H ₂ , 15% Ar, 55% CO, and 10% CO ₂	1.5	NA	300	200	14.2 h	<i>Butyribacterium methylotrophicum</i>	37	acetate	2.71 mmol·L _{RWV} ⁻¹ ·h ⁻¹	[88]
	microbubble sparging 20% H ₂ , 15% Ar, 55% CO, and 10% CO ₂					90.64 h					
Bubble column	CO at different partial pressures	0.2	NA	80	NA	72 h	<i>Eubacterium limosum</i>	37	acetate	3.65 mmol·L _{RWV} ⁻¹ ·h ⁻¹	[75]
Two stage: CSTR - CSTR	30% H ₂ , 30% CO ₂ , 40% CO	2	NA	100	100	NM**	<i>Clostridium ragsdalei</i>	37	ethanol, acetate	NM	[77]
Two stage: CSTR - Bubble Column	60% CO, 35% H ₂ , and 5% CO ₂	CSTR: 1 , Bubble column: 4	NA	CSTR: 25 Bubble Column: 133	CSTR: 200	NM	<i>Clostridium ljungdahlii</i>	35	ethanol	8.04 mmol·L _{RWV} ⁻¹ ·h ⁻¹	[91]
CSTR	55% CO, 20% H ₂ , 10% CO ₂ and 15% argon	2	NA	14	500	NM	<i>Clostridium ljungdahlii</i>	37	ethanol, acetate	1.7 mmol·L _{RWV} ⁻¹ ·h ⁻¹ ethanol and 1.1 mmol·L _{RWV} ⁻¹ ·h ⁻¹ acetate	[92]
CSTR	syngas consisted of 70% CO	2	NA	14	550	135 h	<i>Clostridium ljungdahlii</i>	37	ethanol and acetate	total concentration 11 g·L ⁻¹	[93]

RPB-BR Rotating Packed Bed Bioreactor	20% CO, 5% H ₂ , 15% CO ₂ , and 60% N ₂	1.5	AnoxKaldnes™ K1 carriers	300	50 (rotation of packed bed)	70 (100 rpm and 1 vvm)	<i>Clostridium carboxidivorans P7</i>	37	ethanol	6.06 mmol·L _{RWW} ⁻¹ ·h ⁻¹	[96]
HFMBR	20% CO, 5% H ₂ , 15% CO ₂ , and 60% N ₂	8	microporous hydrophobic polypropylene hollow fibers	300	NA	1096.2 h	<i>Clostridium carboxidivorans P7</i>	37	ethanol	3.12 mmol·L _{RWW} ⁻¹ ·h ⁻¹	[94]
Monolithic Biofilm reactor	20% CO, 5% H ₂ , 15% CO ₂ and 60% N ₂	8	cordierite based monolithic cylinder	300	NA	450 h	<i>Clostridium carboxidivorans P7</i>	37	ethanol	2.13 mmol·L _{RWW} ⁻¹ ·h ⁻¹	[95]
Trickle bed	38% CO, 28.5% CO ₂ , 28.5% H ₂ and 5% N ₂	0.5	6-mm soda lime glass beads	4.6	NA	NM	<i>Clostridium ragdalei</i>	37	ethanol, acetate	5.7 g·L ⁻¹ , 12.3 g·L ⁻¹ respectively	[98]
Continuous gas fed	CO 100%	1.2	NA	NM	250	NM	<i>Clostridium autoethanogenum</i>	30	ethanol, acetate	0.91 g·L ⁻¹ and 0.92 g·L ⁻¹	[99]
CSTR	20% CO, 15% CO ₂ , 5% H ₂ and 60% N ₂	3	NA	NM	150	NM	<i>Clostridium strain P11</i>	37	ethanol	0.58 mmol·L _{RWW} ⁻¹ ·h ⁻¹	[100]
Trickle bed	38% CO, 5% N ₂ , 28.5% CO ₂ and 28.5% H ₂	1	6 mm soda lime glass beads	18.9	NA	NM	<i>Clostridium ragdalei</i>	37	ethanol	3.43 mmol·L _{RWW} ⁻¹ ·h ⁻¹	[101]

*NA – Not Applied **NM – Not Mentioned

Figure 4. Schematics of the monolith biofilm reactor designed by Shen et al. [95]. The red lines depict the syngas flow and the blue lines the liquid flow. The liquid media flow in a loop from the agitated vessel to the bottom of the column, pass through the cordierite monolith (biofilm formed) where fermentation takes place and finally the fermentation broth flows back to the reservoir vessel from the top of the column. Syngas is inserted through a microporous diffuser in the bottom of the column and after its bioconversion in the biofilm it flows from the headspace of the column to the headspace of the reservoir agitated vessel.

Contrary to the current research status that opts for ethanol production Kantzow et al. [102] tried to maximize the acetate productivity taking advantage of the innate acetogenic nature of *Acetobacterium woodii*. For the experiments they used a CSTR reactor with a submerged hollow fiber membrane module to stimulate cell concentration. The membranes were constructed from hydrophilic polysulfone and they were used for full cell retention. It should be highlighted that CO was not included in the gas feed but a gas

mixture of H_2 , CO_2 and N_2 was applied instead. The reactor was operated at 1200 rpm stirring speed with a dilution rate of $0.35\ h^{-1}$ and a productivity of $103.8\ mmol \cdot L_{RWV}^{-1} \cdot h^{-1}$ was achieved. This value was 7.1 times higher than the one without cell retention operated under the same conditions depicting the noteworthy contribution of the hollow fiber membrane module to gas fermentation.

Ethanol and acetate production from syngas has also been tested in trickle bed bioreactors. Devarapalli et al. harnessed the attribute of this configuration and filled a borosilicate glass column with 6 mm soda lime glass beads [98]. This bioreactor setup had already been proved to function for CO fermentation purposes [103]. After 1662 h of semi-continuous operation with biocatalyst *Clostridium ragsdalei* the concentrations of ethanol and acetate were 5.7 and $12.3\ g \cdot L^{-1}$ respectively when liquid recirculation rate was $400\ mL \cdot min^{-1} \cdot L_{RWV}^{-1}$, syngas flow $9.2\ mL \cdot min^{-1} \cdot L_{RWV}^{-1}$, and the flow in co-current mode. Counter-current mode needs further optimization as the author reported flooding of the bed and foam formation at high liquid recirculation rates. The same researchers, studied the effects of dilution and gas flow rates on ethanol productivity by changing the setup from semi-continuous to continuous under co-current and counter-current configurations [101]. They observed that ethanol productivity was higher during co-current mode with a maximum of $3.43\ mmol \cdot L_{RWV}^{-1} \cdot h^{-1}$ at $0.012\ h^{-1}$ dilution, gas flow $18.9\ mL \cdot min^{-1} \cdot L_{RWV}^{-1}$ and liquid recirculation rate of $200\ mL \cdot min^{-1} \cdot L_{RWV}^{-1}$. Flooding issues under counter-current mode were observed again resulting to over two times less ethanol production compared to co-current mode.

In an effort to diminish the cost of media broth, Maddipati et al. [100] replaced $1\ g \cdot L^{-1}$ yeast extract with the much cheaper $20\ g \cdot L^{-1}$ corn steep liquor (CSL) and run experiments in continuous mode in a chemostat. *Clostridium* strain P11 was used as a biocatalyst and the agitation speed was relatively low at 150 rpm. Albeit higher ethanol concentration values were achieved with $20\ g \cdot L^{-1}$ CSL in the medium, the productivity of ethanol was very low ($0.58\ mmol \cdot L_{RWV}^{-1} \cdot h^{-1}$) due to the poor conversion efficiency of CO and H_2 . In another study the addition of $0.75\ \mu m$ tungsten to the media solution for continuous syngas fermentation with *Clostridium autoethanogenum* led to an interesting observation, that is at pH 4.75 no accumulation of

acetate took place [99]. The aforementioned bioreactor set-ups are summed up in table 2 where crucial information on operational parameters is also presented.

4.2 Production of Long Chain Alcohols (Butanol – Hexanol)

Butanol is a liquid fuel that can supplant the use of gasoline and it is emerging as a game-changer in the current view of biofuels [104–106]. Although studies about ethanol production from syngas fermentation can be ubiquitously found, there is limited information regarding continuous production of butanol with syngas compounds as a carbon and energy source. The main reason is that there are few acidogens discovered that can produce butanol naturally [35,36] and therefore metabolic engineering approaches are needed for high butanol selectivity [107–109]. In addition to that, butanol has been reported to be inhibitory to carboxydophilic clostridia [110] and mixed microbial consortia demand a high hydraulic retention times in order to implement carbon elongation [111].

Clostridium carboxidivorans is a microbe that has recently drawn a lot of attention in terms of its inherent ability to produce long chain alcohols [110,112–115]. A try for consolidated butanol – ethanol production was attempted with *Clostridium carboxidivorans* P7 in a CSTR reactor with agitation speed of 250 rpm and temperature of 33 °C [116]. Two experiments were performed under the same conditions except for the pH control. In the first experiment pH was constant at 5.75 for 408 h and then its value was changed to 4.75. The maximum ethanol and butanol concentrations achieved were 5.55 g·L⁻¹ and 2.66 g·L⁻¹ respectively. In the second experiment pH was gradually decreased from 5.75 to 4.75 after 72 h of operation and the maximum concentrations succeeded were 2.90 g·L⁻¹ for ethanol and 1.60 g·L⁻¹ for butanol. The reason for the lower concentrations in the second experiment was that biomass formation was not favored in low pH. *Clostridium carboxidivorans* P7 converts CO first to fatty acids at high pH (acidogenic phase combined with cell growth) and then the formed fatty acids are converted to alcohols at lower pH.

In a syngas fermentation study with bacterium *Alkalibaculum bacchi* an unexpected production of butanol and propanol was observed [117]. The researchers performed 16S rRNA screening of the liquid medium

looking for possible contaminations and the results showed the existence of *Clostridium propionicum* in the fermentation broth. Driven by the incentive to produce higher alcohols with the aforementioned co-culture they performed semi-continuous experiments in a CSTR reactor [118]. The concentrations of alcohols achieved were not high but the results were promising for further research in co-culture syngas fermentations for the production of higher alcohols. Richter et al. [119] deployed a CSTR reactor, inoculated with a co-culture of *Clostridium ljungdahlii* and *Clostridium kluyveri*, connected in line with downstream processing for the production of long chain alcohols from syngas. The reactor volume was 1 L, agitated at 400 rpm, the temperature was at 37 °C and the syngas fed in the reactor had a composition of 60% CO (vol/vol), 35% H₂ and 5% CO₂. While the pH was controlled at 6 and the dilution rate was set at 40 mL·h⁻¹ the net productivities of n-butanol and n-hexanol were 0.408 mmol·L_{RWV}⁻¹·h⁻¹ and 0.220 mmol·L_{RWV}⁻¹·h⁻¹, respectively. It should be highlighted that under the same operational conditions traces of n-octanol were reported at the condensate of the reactor. Another co-culture study was performed in anaerobic bottles by Diender et al. [120], who showed that *C. autoethanogenum* and *C. kluyveri* can work together and convert CO or syngas to medium chain fatty acids and their respective alcohols. The researchers proposed also a model for the metabolic interactions of the two microbes leading to the production of long chain alcohols.

4.3 Methane Production

The first efforts to ferment gas to CH₄ in a continuous process are dated back to 1978 [121]. Wise et al. examined the efficiency of a CSTR reactor functioning under mesophilic and thermophilic conditions, inoculated with mixed microbial consortia from an anaerobic sludge digester. Mixed microbial consortia from anoxic environments are hosts to a vast variety of microorganisms including methanogenic archaea that can reduce C1 sources to CH₄ [122]. Following the stoichiometry, that is for the production of 1 mol of CH₄, 4 mol of H₂ and 1 mol of CO₂ are needed [123], the gas mixture used as a substrate was H₂ and CO₂ in a 4:1 ratio. A cell recycle system was also applied to increase cell concentration. At mesophilic conditions (37 °C) and atmospheric pressure the maximum CH₄ productivity achieved was 178.4 mmol·L_{RWV}⁻¹·h⁻¹ with the

reactor operated at 3.5 d of liquid retention time. Another interesting outcome of this study was that although gas solubility decreases in elevated temperatures better CH₄ productivity could be achieved (446 mmol·L_{RWV}⁻¹·h⁻¹ at 2-2.5 d liquid retention time and 60 °C). According to a study for gas diffusivity in water, the diffusion rate of CO doubles from 35 °C to 60 °C acting as an offset to the decreased solubility [124]. Introduction of 3% CO to the gas feed resulted to a washing out of the cells at a liquid retention time of 8 d indicating the toxic effects of CO.

CH₄ production from syngas was attempted successfully with a triculture consisting of the photosynthetic bacterium *R. rubrum* and two methanogens, *M. formicicum* and *M. barkeri* [125,126]. The research group examined four bioreactor setups, three of which were trickle bed reactors and one a packed bubble column. In all cases the syngas composition was the same but the working volume of the reactors was different. The two trickle bed reactors with volumes as low as 0.736 L and 1.051 L, packed with Intalox saddles presented the highest CH₄ productivities of 3.4 and 3.3 mmol·L_{RWV}⁻¹·h⁻¹ respectively while a scaled up trickle bed reactor with a volume of 26 L exhibited an almost 10 fold decrease in maximum CH₄ productivity. A point to be taken into account though, is that the packing material used in the scaled-up reactor was different. The maximum CH₄ productivity of 0.4 mmol·L_{RWV}⁻¹·h⁻¹ achieved with the packed bubble column bioreactor was significantly lower than the low volume trickle bed reactors indicating that the latter bioreactor configuration is more efficient. Burkhardt et al. [127] used a pilot scale trickle bed reactor (61 L) to convert H₂ and CO₂ to CH₄ and achieved a high product quality (gas of 98% CH₄) with a productivity of 2.77 mmol·L_{RWV}⁻¹·h⁻¹ when the retention time of the gaseous substrate in the packed bed was 4 h.

On the other hand Guiot et al. assessed the potential of microbial populations derived from an industrial granular sludge to convert CO to CH₄ [128]. For their experiments they used a 30 L gas-lift reactor under mesophilic conditions (30 °C) and N₂ – CO diluted gas inflow. The maximal CH₄ specific productivity

achieved was $0.126 \text{ mmol} \cdot \text{g}_{\text{VSS}}^{-1} \cdot \text{h}^{-1}$, when the gas recirculation rate was $600 \text{ mL} \cdot \text{min}^{-1}$ and gas retention time 8.6 d, with the yield of CH_4 from CO reaching 95 % of the theoretical.

A similar idea deploying sewage sludge for simultaneous anaerobic digestion and CO biomethanation was developed by Luo et al. [129]. CO was fed to a stirred tank bioreactor through a hollow fiber membrane module while agitation at 150 rpm was also applied. The 400 mL working volume fermentor, operated under thermophilic conditions (55°C) and neutral pH, presented a high CH_4 productivity of $3.7 \text{ mmol} \cdot \text{L}_{\text{RWV}}^{-1} \cdot \text{h}^{-1}$. However, the percentage of CH_4 in the gas effluent was only 19.2% because of the high CO flow rate of $6.6 \text{ mL} \cdot \text{min}^{-1} \cdot \text{L}_{\text{RWV}}^{-1}$ and, consequently, the low gas retention time of 2.5 h. The CO fraction in the biogas effluent was 35.2%. Sequencing tests showed that the main species involved in CO bioconversion were species close to *Methanosarcina barkeri* and *Methanothermobacter thermautotrophicus*. Detailed information on more operational parameters is presented in Table 3.

A reverse membrane bioreactor configuration was employed by Westman et al. [130] opting for full cell retention within the fermentation vessel. Flat plain hydrophilic PVDF membranes were used for the cell encasement, permeable for the substrate and the product but impermeable for the cells. The sachets containing the inoculum (anaerobic culture from municipal solid sludge digester) were heat-sealed and subsequently inserted in the liquid media. The bioreactor was operated under thermophilic conditions (55°C) with a syngas feed of CO (55% mol), H_2 (20% mol), and CO_2 (10% mol) resulting to a maximum CH_4 productivity of $0.35 \text{ mmol} \cdot \text{L}_{\text{RWV}}^{-1} \cdot \text{h}^{-1}$.

4.4 Biodiesel Production

Despite the broad range of bioreactor configurations proposed for syngas bioconversion, there is only one publication connecting syngas fermentation with continuous biodiesel production [131]. Hu et al. developed an integrated process for the conversion of syngas to lipids in two stages, a bubble column and a stirred tank reactor, linked in series. In the first stage the thermophilic anaerobic acetogen *Moorella*

Table 3. Bioreactor configurations for methane production

Reactor Configuration	Gas Composition	Working Volume (L)	Packing or Hollow Fiber Material	Gas Retention(h)	Microorganisms	Temperature (°C)	Productivity (mmol·L _{RWV} ⁻¹ ·h ⁻¹)	References
CSTR	H ₂ :CO ₂ = 4:1	2	NA	NM	mixed microbial consortia from anaerobic sewage digester (mesophilic)	37	178.4	[121]
CSTR	H ₂ :CO ₂ = 4:1	1	NA	NM	mixed microbial consortia from anaerobic sewage digester (thermophilic)	60	446	[121]
Trickle bed	H ₂ :CO ₂ = 4:1	61	Bioflow 40 (RAUSCHERT)	4	Anaerobic sludge from sewage plant	37	2.77	[127]
Gas-lift	CO	30	NA	NM	industrial granular sludge	37	0.126 ^a	[128]
Reverse membrane bioreactor	55% CO , 20% H ₂ and 10% CO ₂	0.6	Flat plain hydrophilic PVDF	NM	anaerobic culture from municipal solid sludge digester	55	0.35	[130]
Trickle bed	14.8 % Ar, 9.9 % CO ₂ , 55.6% CO and 20.4% H ₂	0.736	6.35 mm Intalox saddles	2.45	triculture (<i>R. rubrum</i> , <i>M. formicicum</i> , <i>M. barkeri</i>)	37	3.4	[125]
Packed bubble column	15 % Ar, 9.6 % CO ₂ , 55% CO and 20.4% H ₂	3.26	Glass Raschig rings (6 mm x 6 mm)	3.40	triculture (<i>R. rubrum</i> , <i>M. formicicum</i> , <i>M. barkeri</i>)	34	0.4	[125]
Trickle bed	14.74 % Ar, 9.72 % CO ₂ , 54.42% CO	26.052	Pall rings	13	triculture (<i>R. rubrum</i> , <i>M.</i>	37	0.45	[126]

	and 21.11% H ₂				<i>formicum, M. barkeri)</i>			
Trickle bed	14.82 % Ar, 9.67 % CO ₂ , 55.62% CO and 19.68% H ₂	1.051	Intalox saddles	1.05	triculture (<i>R. rubrum, M. formicum, M. barkeri)</i>	37	3.3	[126]
CSTR with submerged HFM	CO 100%	0.4	Bundle of 600 microporous polypropylene HFMs with 40% porosity and 0.04 um pore size (Membrana)	2.53	digested sewage sludge	55	2.54	[129]

NA – Not Applied **NM – Not Mentioned

a: mmol·g_{vss}⁻¹·h⁻¹

thermoacetica fixed CO₂ to acetate. The produced acetate acid was in turn pumped through a hollow membrane filter into an aerobic bioreactor, where it was used as a substrate by the genetically engineered oleaginous yeast, *Yarrowia lipolytica*, for the production of lipids. The overall productivity achieved with the two-stage lab scale system was 0.19 g·L_{RWV}⁻¹·h⁻¹ [131]. The same researchers' team had managed to produce acetate at a titer of 31 g·L⁻¹ and a productivity of 9.16 mmol·L_{RWV}⁻¹·h⁻¹ under thermophilic conditions in a bubble column fermentor [132].

4.5 Hydrogen Production

The biological production of H₂ from syngas has drawn less attention than other biofuels due to storage and delivery challenges [133]. In addition to that, syngas compared to other feedstocks presents high microbial toxicity due to its content in CO, which results in diminished productivities. The microbes and the mechanisms involved in biological H₂ production from CO have been recently reviewed in depth from Rittman et al. [134]. In short, carboxydophilic hydrogenogens under strict anaerobic conditions convert CO and H₂O to H₂ and CO₂ via the biological water-gas shift reaction.

Younesi et al. [135] used a CSTR configuration to produce H₂ from syngas with photosynthetic bacterium *Rhodospirillum rubrum* as a biocatalyst. At an optimized syngas flow rate of 14 mL·min⁻¹ and an agitation speed of 500 rpm the maximum H₂ productivity achieved was 24 mmol·L_{RWV}⁻¹·h⁻¹ which was higher than the one reported by Klasson et al. (4.7 mmol·L_{RWV}⁻¹·h⁻¹) using the same bacterium and the same syngas composition (20% H₂, 15% Ar, 55% CO and 10% CO₂), but different source of carbon for cell growth. [136]. An increased productivity of H₂ from CO was achieved in a HFMBR with a pure culture of the extremely thermophilic bacterium *Carboxydotherrmus hydrogenoformans* [137]. Evaluation of a combination of several different operational conditions led to the observation that maximum CO conversion occurs at 70 °C, with a CO partial pressure of 2 atm and liquid velocity through the hollow fiber membrane module of 130 m·h⁻¹, resulting to a H₂ productivity of 125 mmol·L_{RWV}⁻¹·h⁻¹. The same microorganism was also tested for its CO conversion bioactivity in a 30 L gas-lift reactor [138]. Due to a low biomass concentration (highest

value achieved was $0.106 \text{ g}_{\text{VSS}} \cdot \text{L}_{\text{RWV}}^{-1}$) the maximum H_2 productivity was $6.7 \text{ mmol} \cdot \text{L}_{\text{RWV}}^{-1} \cdot \text{h}^{-1}$, which is a value 18 times lower than the one with the HFMBR module.

A different strategy was followed from Oh et al. [139] who cultivated *Rhodopseudomonas palustris* P4 in a CSTR (5 L working volume) system under aerobic conditions with sucrose as a substrate before switching to anaerobic conditions with 10% CO in the gas phase. The pH was maintained at 7, the temperature at 30 °C and the agitation at 700 rpm. When the gas retention time was at 5 min and the inlet CO fraction 20%, the maximum H_2 productivity was $41 \text{ mmol} \cdot \text{L}_{\text{RWV}}^{-1} \cdot \text{h}^{-1}$ with a 61% CO conversion efficiency. The same concept of aerobic growth with sucrose before the initiation of anaerobic CO fermentation was conducted in a CSTR reactor (3 L working volume) by Jung et al. [140] with *Citobacter* sp. Y19. The temperature was maintained at 30 °C, the pH at 7 and the agitation at 500 rpm with a H_2 productivity estimated at $5.71 \text{ mmol} \cdot \text{L}_{\text{RWV}}^{-1} \cdot \text{h}^{-1}$.

Kim et al. [141] studied the effectiveness of a genetically engineered *Thermococcus onnurineus* NA1 strain to convert a 100% CO gas phase into H_2 and compared it to the wild type strain. For their experiments they used a CSTR reactor (2 L working volume) at 80 °C, with the pH controlled at 6.1 – 6.2 and the agitation speed set at 300 rpm. When CO was continuously fed with a flow rate of 240 mL min^{-1} , the H_2 productivity achieved was $123.5 \text{ mmol} \cdot \text{L}_{\text{RWV}}^{-1} \cdot \text{h}^{-1}$, whereas for the wild type strain was $31.8 \text{ mmol} \cdot \text{L}_{\text{RWV}}^{-1} \cdot \text{h}^{-1}$. The aforementioned continuous experiments are collected in Table 4

Table 4. Bioreactor configurations for hydrogen production

Reactor Configuration	Gas Composition	Working Volume (L)	Gas Retention (h)	Microorganisms	Temperature (°C)	Productivity (mmol·L _{RWV} ⁻¹ ·h ⁻¹)	References
CSTR	H ₂ , Ar, CO, and CO ₂ (20/15/55/10)	2	2.4	<i>Rhodospirillum rubrum</i>	30	24	[135]
CSTR	H ₂ , Ar, CO, and CO ₂ (20/15/55/10)	1.25	NM	<i>Rhodospirillum rubrum</i>	30	4.7	[136]
HFMBR	CO	0.16	NM	<i>Carboxydotherrmus hydrogenoformans</i>	70	125	[137]
Gas-Lift	CO	30	NM	<i>Carboxydotherrmus hydrogenoformans</i>	70	6.7	[138]
CSTR	N ₂ and CO (80/20)	5	0.08	<i>Rhodopseudomonas palustris P4</i>	30	41	[139]
CSTR	N ₂ and CO (90/10)	3	0.25	<i>Citobacter sp. Y19</i>	30	5.71	[140]
CSTR	CO	2	0.14	genetically engineered <i>Thermococcus onnurineus NA1</i>	80	123.5	[141]

5. Commercialization of syngas fermentation

Ineos Bio is a company established in 2008 but its bioethanol project started from 2001[142]. A great deal of research including pilot scale tests (where gasification, fermentation and distillation were integrated) led to the industrialization of the process in late 2012 with a capacity of 8 MMGY (million gallons per year) ethanol and 6 MW electricity [143]. The company deployed proprietary naturally occurring biocatalysts with high selectivity to ethanol and tolerant to most impurities produced in the gasification process [142]. The ethanol produced had 99.7 % purity so that it could be mixed with gasoline. However, the plant was idled in 2015 and on September 2016 the company announced its intention to sell the ethanol business due to changes in the US ethanol market that did not conform to their strategic objectives [143]. On July 2017 Alliance Bio-Products Inc. gained the approval of USDA (United States Department of Agriculture) to purchase the Ineos Bio plant with the company planning to use the facility to enhance their cellulose-to-sugar process [144].

Lanzatech is close to commercializing a process converting steel mill off gases to ethanol with two plants being currently under construction; one in China with an ethanol capacity of 16 MMGY and one in Belgium with 21 MMGY [145]. The company uses a proprietary microbial strain for the bioconversion of synthesis gas and a hybrid separation system with water recycle for the purification of the product and the collection of the co-products. The process is already tested in their pre-commercial scale plant in China. Lanzatech presents a high activity, with more than 40 international partnerships with the academia and the private sector, targeting the development of platforms that will convert syngas to a bigger variety of products such as jet fuel, butadiene, 2,3-butanediol, isopropanol and isobutylene [145].

Another company doing business with syngas fermentation was Coskata that developed a semi-commercial process for the fermentation of syngas to fuel chemicals. However, the company went out of business in 2015 and its technology was acquired by the company Synata Bio [146].

6. Conclusions and future perspectives

The evolution of syngas fermentation platforms has an orientation towards packed bed bioreactors and membrane modules combined with biofilm formation. The main reason for the thriving research effort in biofilm reactors is the increased mass transfer that they can provide compared to the traditional bubble column and CSTR systems as well as the enhanced cell retention within the reactor. Recent literature findings show promising ethanol productivities with a rotating packed bed bioreactor and a trickle bed bioreactor configuration while new metabolic engineering techniques have been mainly applied towards an efficient production of higher alcohols and hydrogen. Novel ideas such as reverse membrane bioreactors and hollow fiber membrane bioreactors spring up for syngas biomethanation, yet, more experimental experience is required for the industrial success of these types of set-ups. Future research on the development of bioreactor configurations for continuous syngas fermentation should focus on the 1) design of novel modules that would further enhance gas-to-liquid mass transfer, 2) optimization of the operational parameters in the already used and studied set-ups, 3) combinations of bioreactors in series, 4) study of the versatility of a bioreactor for the production of different bioproducts, the economic assessment and the scale-up potential of the proposed platforms.

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