Biological conversion of methane to polyhydroxyalkanoates: Current advances, challenges, and perspectives

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1. Introduction

Plastics have become indispensable in our daily lives and are widely used in packing materials, household appliances, transportation equipment, and electronic devices [1]. The estimated annual plastics production is 311 million tons [2], and is predicted to surpass 500 million metric tons by 2050 [3]. Plastics are non-degradable, which is the primary disadvantage [4] and has led to the accumulation of plastics in many environments [5]. Further, only a small portion of plastics can be effectively recycled. The cumulative quantity of plastic waste entering the oceans has been estimated at up to 12.7 million metric tons and is predicted to increase by an order of magnitude by 2025 in the absence of waste-management infrastructure enhancements [6]. More importantly, microplastics transfer pollutants via food chains [7] and have negative effects on human health due to their small size and wide distribution in water bodies [7,8]. Moreover, conventional plastics generate two greenhouse gases (GHGs), methane and ethylene, when exposed to ambient solar radiation [5]. Methane is the second-most prevalent GHG after carbon dioxide (CO₂), contributing 18% of the total atmospheric radiation forcing [10]. The global warming potential of methane has increased from 25 to 34 times that of CO₂ over the past 100 years [11], indicating that methane emissions have a significant environmental impact [12]. More than 63% of global methane emissions are anthropogenic [13] and are generated from fossil fuels, anaerobic wastewater treatment, landfilling, coal mining, and natural gas refineries [14]. Moreover, methane accounts for 90% of natural gas, worldwide demand for which is estimated to increase by 44.0% by 2040 [15,16].

Polyhydroxyalkanoates (PHAs) are a kind of polyester produced biologically from carbohydrates and fatty acids through the action of microorganisms [17] with potential applications as a substitute material for conventional plastics [18]. Methanotrophs could accumulate PHB when their nutrient supply is imbalanced [19], and the theoretical yield of PHB synthesis is 67% [20]. Recently, based on its properties of biodegradability and biocompatibility [21,22], demand for PHB products has expanded from its initial applications in packaging materials [22,23] to industrial and agricultural applications [24] and to the biomedical and pharmaceutical sectors [5,25]. Employing PHB into commodities does not require new technologies, as existing equipment originally developed for processing polyethylene and polypropylene can be used [26]. However, the high production costs of PHB, which are 3–4 times higher than those of conventional plastics and similar biopolymer plastics, limits its commercial utilization [27–29]. A major

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proportion of the total costs of PHB (30–40%) is attributed to feedstock [27,30], such as palm oil, glucose, sucrose, and corn starch [19,31,32]. The utilization of methane could reduce production costs by at least 30–35% and make the PHB production process more economically and environmentally friendly [33,34]. A study of the economic feasibility of methane-based PHB production showed that the biosynthesis cost was $8.5/kg PHB when produced at a relatively small scale (500 tons/a) [34]. If the scale of production would be expanded to 100,000 tons/a, costs would be reduced to $4.1–$6.8/kg PHB [33]. Methane emissions from landfills and anaerobic digesters could be used to synthesize PHB, which could theoretically replace 20–30% of the total annual plastics market [35,36]. Therefore, the introduction of methane-based PHB not only opens up a new path for PHB production, but also mitigates the issues related to non-degradable plastics and GHG emissions.

As methane-based PHB has economic and environmental advantages over petroleum-based plastics, this review explores PHB biosynthetic pathways, methanotrophs, bioreactors, and material performance of PHB with an emphasis on overcoming the challenges such as low efficiency of methanotrophs, low gas-liquid mass transfer efficiency, and poor material properties and strategies to overcome them. Further, development opportunities, including feast-famine regimes, engineered microorganisms, gas-permeable membrane bioreactors, two-phase partitioning bioreactors, poly-β-hydroxybutyrate-co-hydroxyvalerate, and high molecular weight of PHB are outlooked.

2. PHB biosynthetic pathway

The oxidation of methane to methanol is catalyzed by methane monooxygenase (MMO) enzymes in both particulate and soluble forms (pMMO and sMMO, respectively) (Fig. 1)[39,40]. Methanol is oxidized to formaldehyde, and formaldehyde can either be converted into biomass or further oxidized to formic acid and CO₂ [39]. Throughout these oxidation steps, cellular carbon and the energy required to continuously uptake methane are obtained [39].

Furthermore, PHB synthesis occurs in three sequential enzymatic steps [41]: two acetyl-CoA molecules condense in the presence of the enzyme β-ketothiolase (phbA) to form acetyl-CoA acetyl-CoA, which is then reduced by nicotinamide adenine dinucleotide (NADH)-dependent acetoacetyl-CoA reductase (phbB) [42]; finally, the polymerization of monomers into PHB polymers is catalyzed by PHB synthase (phbC) [43]. Serine is metabolized to acetyl-CoA, which is directed to the tricarboxylic acid cycle (TCA) cycle under nutrient-sufficient conditions or the PHB cycle under nutrient-deficient conditions [41]. Methane accumulates in the form of PHB by type II methanotrophs via the serine pathway, but ribulose monophosphate methanotrophs may be incapable of PHB synthesis. Type II genera were found to sequence phbC and accumulate PHB, whereas in type I genera, no phbC or measurable PHB was detected [44].

3. Methanotrophs for PHB production

3.1. Methanotrophs

Methanotrophs are classified into two major groups: Group I (α-Proteobacteria; referred to as Type II) [40] and Group II (γ-Proteobacteria; referred to as Type I and Type X) [45]. Type II methanotrophs are the most effective PHB producers and utilize the serine pathway [46]. Isolating efficient methanotrophs with high cell concentrations [20], rapid growth [47], and high PHB content [48] is crucial for improving PHB yields and reducing PHB production costs. Methanotrophic strains utilized in PHB synthesis mainly include Methylosinus trichosporium OB3b [49], Methylosinus trichosporium IMV3011 [38], and Methylocystis hirsute [46]. These methanotrophs are mainly sourced from the Culture Collections and contain PHB contents of 30%, 40%, and 42.5%, respectively. The

![Fig. 1. Methane metabolism in methanotrophs. Adapted from Ref. [37,38].](image-url)
CZ-2 was identified as *Methylobacterium organophilum* and PHB accumulation in the consortium and the isolate in a bioreactor reached 34% and 38% (w/w), respectively [50]. Notably, nitrogen limitation promoted PHB accumulation by up to 57% (w/w) in bioreactors [50]. Furthermore, a high-throughput screening technique combined with a Nile Red flow-cytometric assay dramatically reduced handling time for each experiment in the PHB accumulation studies, creating an integrated system to optimize the methanotrophic consortium [51]. Optimization of calcium and copper levels significantly enhanced PHB production to 49.4% in pure cultures of *Methylcoccus parvus* OBBP while decreasing doubling time from 10.6 h to 8.6 h [51]. These results indicate that the directional isolation of methanotrophs could promote higher PHB production capacities and that highly efficient methanotrophs are very scarce. To gain better insight into PHB processes, future investigations should attempt to isolate highly efficient methanotrophs, and pure-culture fundamental studies should be conducted.

### 3.2. Isolation of highly efficient methanotrophs

Feast-famine strategies have been proven to allow the most effective selection of PHB-accumulating organisms [52]. Providing sufficient carbon sources and nutrients during the feast stage is conducive to the rapid consumption of substrate and accelerates biomass growth, while the growth of biomass is limited by restricting the supply of nutrients in the famine stage, thus inducing carbon storage in the form of PHB [53,54]. During a longer famine stage, non-target microorganisms were selectively died out due to the specific culture conditions for methanotrophs [55,56]. Further, PHB storage capacity and PHB storage rates can be increased by repeated feast-famine cycles in a mixed microbial culture [57,58]: for example, a community of acetate-fed microorganisms rapidly produced up to 89% (w/w) PHA content under these conditions [58]. During the famine phase, PHB yields and PHB concentrations reached 30% and 0.6 g/L, respectively [38]. Adding 0.30 g/L citric acid as an inhibitor of the TCA cycle caused PHB content and concentrations to reach ~40% and 0.26 g/L, respectively [38]. After combining high-throughput screening techniques with a Nile Red assay and optimizing calcium and copper concentrations, the PHB content increased from 18.1% to 49.4% by using *Methylcoccus parvus* OBBP [51]. The combination of high-throughput screening technology and the feast-famine strategy may significantly improve the efficiency of isolating high-producing methanotrophs.

### 3.3. Engineered microorganisms

Methanotrophs could generate various potential target products through genetic engineering; for instance, *Methylmicromobium buryatense* has been genetically engineered to produce C-4 carboxylic acid [59], lactic acid [60] and 2,3-butanediol [61], demonstrating the potential of methanotrophs to produce methane-based PHB via this approach. Through the metabolic modification of microorganisms, regulation of gene expressions in metabolic pathways, and adjustment of cellular metabolic networks, PHB yields could be increased [62]. The overexpression of PHB synthesis pathways involving three genes (phbA, phbB, and phbC) could achieve increased PHB accumulation [63]. Thus, the simultaneous adjustment of the expression of three enzymes could directly optimize the accumulation rates, molecular weight, and performance of PHB [63]. Notably, PHB biosynthesis is actively connected to core metabolic networks and methods to improve metabolic flux in PHB synthesis have been extensively studied [48,64,65], including oxygen limitation [66,67], overexpression of NADH synthesis enzymes [68], substrate limitation [69], and overexpression of PHB synthesis operons [70]. PHB biosynthesis competes with several alternative metabolites and intermediates, making it vital to attenuate competitive pathways for PHB synthesis pathways [62] via, for example, reducing acetyl-CoA consumption during the TCA cycle [71] or preventing β-oxidation [72,73]. Meanwhile, the clustered regularly interspaced short palindromic repeats interference (CRISPRi) has been employed in PHA biosynthesis to induce direct metabolic flux [74,75]. The phbC activity can be regulated by designing CRISPRi binding sites using various sgRNA [62,74]. Interestingly, phbC activity was found to influence PHB content and molecular weights in direct and reverse proportions, respectively [76], allowing PHB contents to be controlled at 1.47–75.21% (w/w) levels and molecular weights at 2–6 million Dalton (MDa) [74]. These results suggest the expanding potential of manipulating and optimizing methanotrophic PHB synthesis by engineered microorganisms.

### 4. Bioreactor configuration and operation

#### 4.1. Bioreactors

Methane-based PHB production faces unique challenges due to the low solubility of methane and oxygen in aqueous solutions (22 mg/L and 9 mg/L at 20 °C and 1 atm, respectively) [77]. Low gas mass transfer rates result in low cell density and PHB synthesis rates, greatly limiting the large-scale production of PHB by methanotrophs [78,79] used pressure bioreactors to increase the mass transfer efficiency, and PHB content reached 51% [20]. Reduced the liquid medium from 0.5 L to 0.1 L and enhanced the cell density of methanotrophs from 1.4 g/L to 5.0 g/L. Although high pressure would significantly increase the solubility of methane, creating such conditions can be expensive and detrimental for microorganisms [80]. In another study, vigorous agitation reduced the size of bubbles and achieved a large gas-liquid volumetric mass transfer coefficient (k_a) of 102.9 h⁻¹ [77], but it increased the power consumption of the operation impeller and led to high shear stress that resulted in cell damage [81]. Biosynthesis of PHB by *Methylcoccus hirsuta* from natural gas using bubble column bioreactors (BCBs) and vertical tubular loop bioreactors, respectively [46]. Under optimum conditions, the high cell density cultivation of *Methylcoccus hirsuta* using natural gas in a BCB lead to PHB accumulation of up to 73.4% [82]. A 25% w/w PHB content was obtained under optimized operating conditions and engineering parameters in a BCB [83]. Suspended-growth BCBs can overcome mass transfer limitations through the use of ultrafine bubble diffusers with micropores size <0.5 μm [84].

#### 4.2. Optimization of bioreactor design

##### 4.2.1. Gas-permeable bioreactors

The low solubility of methane and oxygen limits the production efficiency of methane-based PHB to some extent. A novel membrane biofilm reactor application could deliver gaseous substrates (e.g. methane and oxygen) to microorganisms [85,86], similar to a H₂-based membrane biofilm reactor. Gas-permeable hollow fibers could be used to efficiently deliver methane, which would enable the gas diffusion free of bubbling [87]. The nitrate removal rate was increased to ~700 mg N/L/d when methane was employed as an electron donor through gas-permeable hollow fiber membranes [88,89]. Furthermore, hollow fiber membrane bioreactors are considered as an effective method to promote methane utilization, as they shorten the methane transfer pathway [90] and mitigate limitations associated with methane transfer [91–93]. Therefore, it seems feasible that the efficiency of the gas-liquid mass transfer of
methane and oxygen could be significantly improved via gas permeable bioreactors [94]. Hollow fiber membranes to achieve a high kLa of methane (1501 h⁻¹) and cell growth rate for Methylosinus trichosporium OB3b was increased by 67.3%. Our laboratory has designed a double-membrane aerobic methanotrophs-culture device (ZL201821323775.2), shown in Fig. 2. The small pore size and high contact area of membranes would significantly enhance the mass transfer rate of methane and oxygen. Based on their high gas transfer efficiency and homogeneous diffusion, gas permeable membranes appear to be an advantageous option [95]. However, gas-permeable membranes also have certain drawbacks, such as membrane fouling and membrane resistance to mass transfer [96]. Apart from their physical characteristics, gas-permeable membrane bioreactors have potential to allow a safe and efficient gas-supply technology and establish an efficient PHB-synthesis process.

### 4.3. Two-phase partitioning bioreactors

Based on the addition of a non-aqueous phase (NAP) with a high affinity for poorly water-soluble compounds (e.g., methane and oxygen) [97,98], two-phase partitioning bioreactors (TPPBs) have a higher potential to drive mass transfer [99] (Fig. 3). Furthermore, most NAPs used in TPPBs have shown a high affinity for oxygen; thus, the mass transfer rate of oxygen is also improved [100,101]. The methane conversion efficiency can be artificially improved by adding a methane vector such as silicone oil or paraffin oil. Methane average volumetric elimination capacity increased by 131% up to 51 g m⁻³ h⁻¹ with added 10% (v/v) silicone oil in TPPBs [102]. Under nitrogen-limited conditions, the CZ-2 achieved a PHB accumulation of 57% in TPPBs [50]. After the addition of 1-propanol, the cell growth rate and the maximum optical density of Methylosinus trichosporium OB3b enhanced by 700% and 730%, respectively [103]. TPPBs are also particularly suitable for the treatment of enhancing the kLa of methane and oxygen. The long-term stability of TPPBs is vital, because the NAP may be degraded or lost during continuous operation [104]. Moreover, the feasibility of the use of silicone oil as a NAP in TPPBs is limited due to its expense. However, the cost of industrial-grade silicone oil is ~55 times lower than that of high-purity silicone oil [101], and the efficacy of industrial-grade silicone oil should be experimentally assessed. Further, electrolytes (e.g., KCl, NaCl, K₂SO₄, Na₂SO₄) can suppress the coalescence of methane bubbles, increasing the gas-liquid kLa of methane from 103 h⁻¹ to 711 h⁻¹ [78].

### 5. Strategies for enhancing PHB-material performances

Current PHB applications are restricted due to PHB’s narrow melt processing window, thermal instability, and brittleness [105]. PHB quality has a decisive impact on the mechanical performance and processability of subsequent products [76]. Manipulation of the molecular weight (Mw) and the synthesis of poly-β-hydroxybutyrate-co-hydroxyvalerate (PHBV) seems to be effective methods to improve the material performance and explore its potential applications [25].

#### 5.1. Manipulation of high molecular weight PHB

The Mw of PHB determines its thermodynamic properties and crystallization characteristics, which strongly influence its mechanical properties and processability [106]. Increasing the Mw of PHB can promote β-form crystallization, and in turn, enhance the mechanical properties of PHB [107]. The Mw of PHB (10⁵–10⁶ Da) depends on the PHB’s cultivation conditions and the host microorganism. Normal PHB is characterized by brittleness and a secondary crystallization phenomenon [108], while the tensile strength of PHB with an ultra-high Mw can reach 1320 MPa, which is higher than petroleum-based products [109]. Strategies for high Mw of PHB production are summarized in Table 1.

Under potassium-limited conditions, the production of PHB with an Mw up to 3.1 MDa was accumulated [105]. When sulfur or iron was limited, the Mw of PHB decreased by 20.6% and 41.6%, respectively [105], indicating PHB synthesis is hampered under iron-deficient conditions. The addition of citric acid and methanol
(0.1%) increased PHB yields from 12% to 40% and increased PHB Mw up to 1.5 MDa [38]. The expression level or activity of phbC [76,112] and depolymerase [76] and the effectiveness of the chain transfer reactions [113] could also affect the Mw of PHB. To change the expression of PHB synthase, a rationally designed library of ribosomal binding sites with defined strengths for the PHB synthesis operon of phbA, phbB, and phbC were constructed in high or low copy-number plasmids in E. coli. High-quality PHA (Mw: 2.7–6.8 MDa) could be efficiently produced through the combination of a one-step library and proper screening [63].

5.2. PHBV synthesis

One strategy to enhance the processability of PHB is the introduction of 3-hydroxyvalerate (3HV) units into PHB polymers to produce PHBV. The melting point of PHBV lowers as the proportion of 3HV monomers increases, leading to a broader thermal processing window [114,115] and improved polymer ductility and flexibility [116]. Hence, PHBV has attracted increasing attention for integrated biomedical applications [117] and food-packaging applications [118] due to its improved resilience, flexibility, and strength. PHBV synthesis strategies are summarized in Table 2.

As the HV content increased to 40%, the melting temperature of PHBV significantly decreased from 180 °C to 75 °C [122]. Only PHB, not PHBV, was synthesized when methane was used as the sole carbon substrate by Methylocystis-enriched bacteria [119]. PHBV was synthesized and the mol% of 3HV was increased with added valerate [115]. Adding propionate and valerate can modify the mol% 3HV composition, where valerate addition leads to higher PHB content and mol% 3HV [119]. High HV content (25–35%) was obtained in methane-propionate experiments by the CZ-2 [121]. The copolymer had a 3HB:3HV:3HO ratio of 55:35:10 when the feedstock consisted of citrate plus methane, and had a HB: HV ratio of 75:25 with propionate feeding plus methane [121]. Tailored PHB-copolymer composition could be generated through the addition of different co-substrates [115]. Under cobalt-deficient conditions, Methylobacterium extorquens AM1 was able to synthesize PHBV from methanol alone [123]. Although methanotrophs can produce

**Table 1**

| Microorganisms       | Mw (MDa) | Reactor Type  | Technology       | PHB (%) | Y<sub>PHB/CH4</sub> (g g<sub>0</sub>/C0) | References |
|----------------------|----------|---------------|------------------|---------|---------------------------------------|------------|
| Methylocystis sp.    | ^GB 25 DSM 7674 | 3.10 | Pressure bioreactor | Potassium deficiency | 33.60 | 0.45 [105] |
| Methylocystis sp.    | ^GB 25 DSM 7674 | 2.46 | Pressure bioreactor | Sulfur deficiency | 32.60 | 0.40 [105] |
| Methylocystis sp.    | ^GB 25 DSM 7674 | 1.81 | Pressure bioreactor | Iron deficiency | 10.40 | 0.22 [105] |
| Methylosinus trichosporium | ^IMV3011 | 1.50 | Shake flasks | Citric acid | 40.00 | – [38] |
| Methylocystis sp.    | ^GB25 DSMZ 7674 | 2.50 | Pressure bioreactor | Ammonium deficiency | 51.30 | 0.52 [110] |
| Methylocystis sp.    | ^GB25 DSMZ 7674 | 2.50 | Pressure bioreactor | Phosphorus deficiency | 46.80 | 0.55 [110] |
| Methylocystis sp.    | ^GB25 DSMZ 7674 | 2.50 | Pressure bioreactor | Magnesium deficiency | 28.30 | 0.37 [110] |
| Methylocystis sp.    | ^GB25 DSM 7674 | 2.30 | Pressure bioreactor | Nitrogen deficiency | 51.00 | 0.54 [111] |

**Table 2**

| Microorganism                | HV (mol%) | PHAs content (w%) | Co-substrate | References |
|------------------------------|-----------|-------------------|--------------|------------|
| Methylocystis parvus OB85P   | 22        | 54                | Valerate (100 mg/L) | [119] |
| Methylocystis parvus OB85P   | 37        | –                 | Valerate (400 mg/L) | [119] |
| Methylocystis parvus OB85P   | 8         | 32                | Propionate (100 mg/L) | [119] |
| Methylocystis sp. WRRC1       | 60        | 78                | Valerate     | [120] |
| Methylocystis-dominated      | 18–22     | 43–45             | Valerate (100 mg/L) | [115] |
| Methylobacterium organophilum | ^CZ-2     | 35                | Citrate      | [121] |
| Methylobacterium organophilum | ^CZ-2     | 25                | Propionate   | [121] |

**Fig. 3.** Mass transfer models in (a) conventional systems (b) two-phase partitioning bioreactors. NAP: non-aqueous phase.
PHBV, it is still necessary to provide co-substrates and carbon source supplements, which increases the cost of PHBV production.

6. Conclusions
Methane-based PHB production has an excellent potential to benefit the environment as a substitute for petroleum-based materials and to reduce GHG emissions. This review provided an analysis of PHB biosynthetic pathways and identified the challenges and opportunities associated with methanotrophic PHB production. Obtaining high-efficiency methanotrophs through a feast-famine strategy and microorganism engineering, coupled with gas-permeable membrane bioreactors and two-phase partitioning bioreactors could help enhance gas-liquid transfer efficiency. Further, manipulating the molecular weight of PHB and generating PHBV are necessary to achieve high-quality products. Further investigations of methane-based PHB would help support the complex needs of the bioplastics market.

Declaration of competing interest
We declare that we have no financial and personal relationships with other people or organizations that could inappropriately influence the work, there is no professional or other personal interest of any nature or kind in any product, service and/or company that could be construed as influencing the position presented in, or the review of, the manuscript entitled.

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References
[1] M.V. Reddy, Y. Mawatari, R. Onodera, Y. Nakamura, Y. Yajima, Y.C. Chang, Bacterial conversion of waste into polyhydroxybutyrate (PHB): a new approach of bio-circular economy for treating waste and energy generation, Bioresource Technology Reports 7 (2019) 100246.
[2] U.T. Bornscheuer, Feeding on plastic, Science 351 (6278) (2016) 1154–1155.
[3] X. Chen, N. Yan, A brief overview of renewable plastics, Materials Today 16 (2013) 627–635.
[4] E. Akaraonye, T. Keshavarz, I. Roy, Production of polyhydroxyalkanoates: the approaches of bio-circular economy for treating waste and energy generation, Bioresource Technology Reports 7 (2019) 100246.
[5] J.C.C. Yeo, J.K. Muriuki, W. Thitsartarn, Z.L. Li, C. He, Recent advances in the development of biodegradable PHB-based toughening materials: approaches, advantages and applications, Mater. Sci. Eng. C 92 (2019) 1022–1116.
[6] Q. Liao, A. Tsui, S. Billington, C.W. Frank, Extruded foams from microbial poly(3-hydroxybutyrate-co-3-hydroxyvalerate) and its blends with cellulose acetate butyrate, Polym. Eng. Sci. 52 (7) (2012) 1495–1508.
[7] A. Tsui, X. Hu, D.L. Kaplan, C.W. Frank, Green Polymer Chemistry: Biocatalysis and Materials II, American Chemical Society, 2013, pp. 251–279.
[8] S. Kourmentza, J. Placido, N. Venetsaneas, A. Burniol Figols, C. Varrone, H.N. Gavala, M.A.M. Reis, Recent advances and challenges towards sustainable poly(3-hydroxyalkanoate) (PHA) production, Bioengineering 4 (2) (2017) 55.
[9] A.K. Singh, J.K. Srivastava, A.K. Chandel, L. Sharma, N. Mallick, S.P. Singh, Biomedical applications of microbially engineered polyhydroxyalkanoates: an insight into recent advances, bottlenecks, and solutions, Appl. Microbiol. Biotechnol. 103 (5) (2019) 2007–2032.
[10] F.I. Butt, N. Muhamad, A. Hamid, M. Moniruzzaman, F. Sharif, Recent progress in the utilization of bio-synthesized polyhydroxyalkanoates for biomedical applications – Review, Int. J. Biol. Macromol. 120 (2018) 1294–1305.
[11] J.P. Flores, M. Fazel, R.A. Simão, Preparation and characterization of thermosensitive starch composite reinforced by plasma-treated poly (hydroxybutyrate) PHB, Int. J. Biol. Macromol. 123 (2019) 609–621.
[12] K.G. Harding, J.S. Dennis, H. von Blöttzit, S.T.L. Harrison, Environmental analysis of plastic production processes: comparing petroleum-based polypropylene and polylethylene with biologically-based poly(3-hydroxybutyric) acid using life cycle analysis, J. Biotechnol. 130 (1) (2007) 57–66.
[13] I. Levett, G. Birkett, N. Davies, A. Bell, A. Langford, B. Laycock, P. Lant, S. Pratt, Techno-economic assessment of poly(3-hydroxybutyrate) (PHB) production from methane—the case for thermophilic bioprocessing, Journal of Environ. Chemical Engineering 4 (4) (2016) 3724–3733.
[14] H.F. Listewnik, K.D. Wendlandt, M. Jechorek, G. Mursel, Process design for the microbial synthesis of poly(3-hydroxybutyrate) (PHB) from natural gas, Eng. Life Sci. 7 (3) (2007) 278–282.
[15] B.E. DiGregorio, Biobased performance bioplastic: mirel, Chem. Biol. 16 (1) (2009) 1–2.
[16] K.H. Rostekowski, C.S. Criddle, M.D. Lepech, Cradle-to-Gate life cycle assessment for a cradle-to-cradle cycle: biogas-to-bioplastic (and back), Environ. Sci. Technol. 46 (18) (2012) 9822–9829.
[17] R. Hanson, T. Hanson, Methanotrophic bacteria, Microbiol. Rev. 60 (1996) 421–471.
[18] Y. Zhang, J. Xin, L. Chen, H. Song, C. Xia, Biosynthesis of poly-3-hydroxybutyrate with a high molecular weight by methanotroph from methane and methanol, J. Nat. Gas Chem. 17 (1) (2008) 103–108.
[19] M.G. Kalyuzhnaya, A.W. Puri, M.E. Lidstrom, Metabolic engineering in methanotrophic bacteria, Metab. Eng. 29 (2015) 142–152.
[20] J.D. Semrau, A.A. Dispirito, S. Yoon, Methanotrophs and copper, FEMS (Fed. Eur. Microbiol. Soc.) Microbiol. Rev. 34 (4) (2010) 496–531.
[21] P.J. Strong, B. Laycock, S.N.S. Mahamud, P.D. Jensen, P.A. Lant, G. Tyson, Intergovernmental Panel on Climate Change 1535, 2013.
trichosporium OB3b fermentation, J. Ind. Eng. Chem. 39 (2016a) 149–152.

[95] H. Li, Z. Zhang, X. Xu, J. Liang, S. Xia, Bioreduction of para-chloronitrobenzene in a hydrogen-based hollow-fiber membrane biofilm reactor: effects of nitrate and sulfate, Biodegradation 25 (2) (2014) 205–215.

[96] V. Gupta, R. Goel, Managing dissolved methane gas in anaerobic effluents using microbial resource management-based strategies, Bioreour. Technol. 289 (2019) 121601.

[97] C. Kennedy, E. Clifford, S. Gerrity, R. Walsh, M. Rogers, G. Collins, A horizontal flow biofilm reactor (HFB) technology for the removal of methane and hydrogen sulphide at low temperatures, Water Sci. Technol. 66 (9) (2012) 2007–2012.

[98] A. Orzdz, J.C. Lopez, I. Figueroa Gonzalez, R. Muinoz, G. Quijano, Assessment of methane biodegradation kinetics in two-phase partitioning bioreactors by pulse respirometry, Water Res. 67 (2014) 46–54.

[99] K.G. Clarke, P.C. Williams, M.S. Smit, S.T.L. Harrison, Enhancement and repression of the volumetric oxygen transfer coefficient through hydrocarbon addition and its influence on oxygen transfer rate in stirred tank bio-

[100] J. Balbono, M. Nikolauz, M. Kastner, M.C. Veiga, C. Kennes, Removal of dichloromethane from waste gases in one- and two-liquid-phase stirred tank bioreactors and biotrickling filters, Water Res. 43 (1) (2009) 11–20.

[101] K. Muinoz, A.J. Daugulis, M. Hernández, G. Quijano, Recent advances in two-phase partitioning bioreactors for the treatment of volatile organic com-

[102] J. Rocha Rios, S. Bordel, S. Hernández, S. Revah, Methane degradation in two-

[103] K. Kim, Y. Kim, J. Yang, K.S. Ha, H. Usta, J. Lee, C. Kim, Enhanced mass transfer rate and solubility of methane via addition of alcohols for Methylosinus trichosporium OB3b fermentation, J. Ind. Eng. Chem. 46 (2017) 350–355.

[104] R. Muinoz, S. Villaverde, B. Gueyishe, S. Revah, Two-phase partitioning bio-

[105] J. Helm, K.D. Wendlandt, M. Jechorek, U. Stottmieer, Potassium de

[106] J. Choi, S.Y. Lee, High level production of supra-molecular weight poly-3-

[107] T. Iwata, Strong fibers and films of microbial polyesters, Macromol. Biosci. 5 (2005) 689–701.

[108] K.D. Wendlandt, M. Jechorek, J. Helm, U. Stottmieer, Production of poly-3-

[109] K.D. Wendlandt, M. Jechorek, J. Helm, U. Stottmieer, Production of PHB food packaging applications, Food Packaging and Shelf Life 21 (2019) 100029.