Molecular Mechanism Associated With the Impact of Methane/Oxygen Gas Supply Ratios on Cell Growth of Methylomicrobium buryatense 5GB1 Through RNA-Seq

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The methane (CH4)/oxygen (O2) gas supply ratios significantly affect the cell growth and metabolic pathways of aerobic obligate methanotrophs. However, few studies have explored the CH4/O2 ratios of the inlet gas, especially for the CH4 concentrations within the explosion range (5~15% of CH4 in air). This study thoroughly investigated the molecular mechanisms associated with the impact of different CH4/O2 ratios on cell growth of a model type I methanotroph Methylomicrobium buryatense 5GB1 cultured at five different CH4/O2 supply molar ratios from 0.28 to 5.24, corresponding to CH4 content in gas mixture from 5% to 50%, using RNA-Seq transcriptomics approach. In the batch cultivation, the highest growth rate of 0.287 h⁻¹ was achieved when the CH4/O2 supply molar ratio was 0.93 (15% CH4 in air), and it is crucial to keep the availability of carbon and oxygen levels balanced for optimal growth. At this ratio, genes related to methane metabolism, phosphate uptake system, and nitrogen fixation were significantly upregulated. The results indicated that the optimal CH4/O2 ratio prompted cell growth by increasing genes involved in metabolic pathways of carbon, nitrogen and phosphate utilization in M. buryatense 5GB1. Our findings provided an effective gas supply strategy for methanotrophs, which could enhance the production of key intermediates and enzymes to improve the performance of bioconversion processes using CH4 as the only carbon and energy source. This research also helps identify genes associated with the optimal CH4/O2 ratio for balancing energy metabolism and carbon flux, which could be candidate targets for future metabolic engineering practice.

Keywords: methanotroph, Methylomicrobium buryatense, CH4/O2 gas supply ratio, transcriptomics, methane metabolism, nitrogen fixation
INTRODUCTION

As climate change and global warming become severe, much attention has been given to explore more efficient and economical approaches to utilize the GHGs. Methane (CH$_4$) as one of the major GHGs has a much greater global warming potential than CO$_2$. Currently, more than 50% of the CH$_4$ emission is contributed by natural gas from energy extractions and biogas from agriculture and enteric fermentations (Opio et al., 2013). Because of the development of horizontal drilling and hydraulic fracturing techniques, the unconventional natural gas (shale gas and tight gas) is produced at a surplus globally in last decade. The tremendous increase in shale gas also caused a huge amount gas flared annually at extraction sites around the world. Therefore, it is urgent to capture CH$_4$ and fulfill the value of this feedstock.

Methanotrophic bacteria are a subset of methylotrophs, which are capable of using CH$_4$ as the sole carbon source for cell growth and biosynthesis of metabolites. Thus, the biological conversion of CH$_4$ has been gathering more and more interest as a key role in global carbon cycle. So far, significant efforts have been undertaken to utilize CH$_4$ as the substrate to produce biofuels and biochemicals by methanotrophs.

*Methylomicrobium buryatense* 5GB1 is a type I methanotroph that assimilates CH$_4$ to methanol using methane monooxygenase (pMMO) first, and then metabolizes methanol mainly through RuMP pathway for pyruvate production and cell growth (Fu et al., 2017). Recently, *M. buryatense* 5GB1 has been proved as a promising industrial strain due to its robust growth as well as the established genetics and synthetic biology approaches for strain engineering. A series of genetic constructions and metabolic engineering works of *M. buryatense* 5GB1 have been carried out to enhance the carbon flux for the production of biomass, lipids, lactic acid, fatty acids, etc. (Dong et al., 2017; Fu et al., 2017; Hu et al., 2017; Garg et al., 2018). The highest lipid productivity of 45 mg/L/h along with the growth rate of 0.22 h$^{-1}$ and DCW of 21 g/L has been achieved by knocking out glycogen biosynthesis genes of *M. buryatense* 5GB1 in batch cultures (Fu et al., 2018). The flux balance model of *M. buryatense* 5GB1 also suggests that the maximum carbon conversion efficiency (CCE) can be as high as 60-70% on molar basis. Garg et al. (2018) have successfully improved the CCE 50 times by using suitable promoters and RBS combinations for lactic acid production by *M. buryatense* 5GB1.

*Abbreviations*: ANOVA, analysis of variance; CCE, carbon conversion efficiency; CH$_4$, methane; ClpX, ATPase and specificity subunit of ClpX-ClpP ATP-dependent serine protease; DCW, dry cell weight; DEGs, differentially expressed genes; DraG, ADP-ribosylhydrolase; DraT, ADP-ribosyltransferase; Fbp, fructose-1,6-bisphosphatase; GHGs, greenhouse gases; Gik, glucokinase; GlgG, response regulator; GlgK, nitrogen assimilation regulatory protein; GlgN, sensory histidine kinase; MoxF, methanol dehydrogenase [cytochrome c] subunit 1; NifA, Nif-specific regulatory protein; NifD, nitrogenase molybdenum-iron protein alpha chain; NifH, nitrogenase iron protein; NifHDK, nitrogenase; NifK, nitrogenase molybdenum iron protein beta chain; NifL, anti-activator of NifA; NMS, nitrate mineral salt; OadG, putative oxaloacetate decarboxylase gamma chain; OD, optical density; PhoB, DNA-binding response regulator in two-component regulatory system with PhoR; PhoU, phosphate uptake regulator; pMMOs, particulate methane monoxygenases; Por, pyruvate-ferrodoxin/flavodoxin oxidoreductase; PstA, high-affinity phosphate transport protein membrane subunit; PstB, high-affinity phosphate transporter ATP-binding subunit; PstC, high-affinity phosphate transport protein membrane subunit; PstS, periplasmic phosphate-binding protein; PstS1a, ABC-type phosphate transport system; RBS, ribosome-binding site; RuMP, ribulose-monophosphate.

*Methylomicrobium buryatense* 5GB1 is an aerobic obligate methanotroph, and the oxygen supply is an essential factor influencing cell growth, electron transfer, and energy supply. It has been projected that the growth rate is associated with the consumption rates of CH$_4$ and O$_2$, which may be caused by the differences of NADH production via oxidative phosphorylation. Similar phenomenon has been observed in cultures of *M. buryatense* 5GB1 with two different CH$_4$/O$_2$ ratios of inlet gas mixtures. It was found that a better growth and gas update rate were obtained in the cultures with CH$_4$/O$_2$ ratios of 4.4 on molar basis, compared with that of 0.14 (Zhu et al., 2017).

Moreover, transcriptomic studies suggested that the expression of several key genes associated with NADH formation (formaldehyde to carbon dioxide) was changed dramatically under O$_2$-limited conditions (Gilman et al., 2015). Akberdin et al. (2018) also demonstrated that *Methylomicrobium alcaliphilum* 20Z cultured under aerobic or micro-aerobic conditions affected the formation of pyruvate and biomass significantly. It has also been shown that *M. buryatense* 5GB1 exhibited a complex metabolism under O$_2$ starvation (Gilman et al., 2017). These findings suggest that under different CH$_4$/O$_2$ gas supply ratios, the expression of certain genes can be regulated between aerobic methanotrophic growth and denitrification phases, which is important for industrial applications of methanotrophs.

Although applying optimal CH$_4$/O$_2$ ratios of the inlet gas exhibits significant effects on the overall CCE of metabolic pathways related to cell growth, carbon flux, reducing power and energy generation, few studies have thoroughly explored the CH$_4$/O$_2$ ratios of the inlet gas, especially for the CH$_4$ concentrations within the explosion range (5~15% of CH$_4$ in air), which is mainly due to the safety concerns. We have already developed an automatic control system with CH$_4$ alarm and cut-off device along with gas mass flow meters to keep operation running smoothly and safely (Fei and Pienkos, 2018). From the standpoint of revealing the influence of the inlet gas ratio of CH$_4$/O$_2$ on cell growth, nutrients utilization and energy consumption, it is desirable to understand the effects of various inlet gas ratios of CH$_4$/O$_2$ on methanotrophic metabolism using a lab-scale, safe bioconversion system (Fei and Pienkos, 2018).

To explore the changes at transcriptional levels globally, many researchers have applied transcriptomics to their studies. Recently, Fu et al. (2019) reported significant differences between growth on CH$_4$ and methanol by transcriptional analysis in *M. buryatense* 5GB1. Similarly, a comparative transcriptome analysis was utilized in *Methylomonas* sp. DH-1 to explore the roles of secondary metabolite pathways during growth on CH$_4$ and methanol (Nguyen et al., 2019). However, there is no systematical research on *M. buryatense* 5GB1 under different CH$_4$/O$_2$ ratios.

In this study, five different inlet gas ratios of CH$_4$/O$_2$ (including three ratios within explosion range for the first time) were investigated in methanotrophic cultivation with continuous supply of CH$_4$ and O$_2$ in bioreactors. Cell growth and key metabolites were measured to analyze the impact of different CH$_4$/O$_2$ ratios on *M. buryatense* 5GB1, and transcriptomics study was performed to identify genes differentially expressed under these conditions to help elucidate the impact of CH$_4$/O$_2$ ratio.
on cell growth and metabolism of M. buryatense 5GB1. The effective gas supply ratio identified in this study could help enable the production of key intermediate such as pyruvate, which is an important precursor for various downstream biochemical biosynthesis, and therefore offers a potentially efficient path to produce biochemicals using CH₄ as the only carbon source by improving nutrient utilization and energy supply.

**MATERIALS AND METHODS**

**Strain and Culture Conditions**

*Methylocrobium buryatense* 5GB1 obtained from Prof. Mary Lidstrom was revived on NMS agar medium for 5 days, while seed culture was prepared in a gas cylinder filled with gas mixture of 20% CH₄ in air at 30°C. The medium composition of NMS used in batch cultures was mentioned previously (Fei et al., 2018). Aerobic fermentation was carried out in a 3 L fermentation cylinder (BaoXing Bio-engineering Equipment, Shanghai, China) containing 2 L of modified NMS2 (8 g/liter KNO₃ and 2 × trace elements) medium by mass-flow meter controlling mixed gas content with reliable explosion-proof equipment and good ventilated condition. The composition of the gas blend used in this experiment was as follows: 0.28 (#1, 5% CH₄ in air), 0.98 (#2, 10% CH₄ in air), 0.93 (#3, 15% CH₄ in air), 1.31 (#4, 20% CH₄ in air), and 5.24 (#5, 50% CH₄ in air).

**Cell Growth and Metabolite Measurements**

Optical density of cell was measured at 600 nm with a spectrophotometer (TU-1810, PERSEE, Beijing, China). As for DCW, sampling was performed every 12 h. 10 mL bacterial solution was placed in a 15-mL centrifuge tube each time and centrifuged at 7,000 rpm for 15 min. The supernatant was discarded, and cells were then resuspended with 5 mL deionized water. Centrifugation step was repeated once. Cells were preserved in the −80°C refrigerator for 4 h, which was then dried by a lyophilizer for 36~48 h before measuring the dry weight of the cells. Statistical analysis was performed in Microsoft Excel 2007, and P-values with statistical significance of P < 0.05 and P < 0.01 were obtained.

Total protein and pyruvate were determined using kits from Beyotime and Suzhou Comin Biotechnology. The total protein was measured using the Bradford Protein Assay Kit (Beyotime, Shanghai, China). Briefly, 1 mL bacterial culture was transferred to the 1.5 mL microcentrifuge tubes for cell lysis using the ultrasonic cell breaker (YH-1000Y, Ningbo, China). The procedure also involved in adding a single working reagent for pyruvate determination at 520 nm by UV-spectrophotometry.

**Enzymatic Assays**

Particulate methane monooxygenase activity was measured using propene epoxidation assay as described before (Miyaji et al., 2002). The intracellular NADH and NAD⁺ concentrations were measured by the enzyme cycling method (He et al., 2016). Briefly, 1-mL bacterial liquid was transferred to the 1.5-mL microcentrifuge tubes for NADH or NAD⁺ extraction. After centrifugation, cell pellets were resuspended with 300 µL 0.2 M NaOH for NADH quantification or 300 µL 0.2 M HCl for NAD⁺ determination. The samples were bathed in 50°C for 10 min and immediately placed in iced water. The cell lysate was neutralized by adding the same volume of HCl or NaOH. The supernatants were stored at −20°C and used to detect the NADH and NAD⁺ activity after centrifuging at 15,000 rpm for 5 min. The solution (total volume was 0.6 mL) contained 0.1 mL of each of followings: 1.0 M Bicine buffer (pH 8.0), absolute ethanol, 40 mM EDTA (pH 8.0), 4.2 mM MTT (thiazolyl blue), and 0.2 mL of 16.6 mM PES (phenazine methosulfate). 50 µL extract, 0.3 mL ddH₂O, 0.6 mL mixture and 50 µL alcohol dehydrogenase (500 U/mL) were then mixed to start the reaction, and the absorbance at 570 nm was measured with the spectrophotometer (TU-1810, PERSEE, Beijing, China) for 5 min. Commercial NADH and NAD⁺ (Meyer Chemical Technology, Shanghai, China) were used to make the standard curves.

**Transcriptomic Analysis**

Transcriptomics study was carried out following the method described in previous reports (He et al., 2018; Yang et al., 2018). Briefly, cell culture samples were collected within 24 h fermentation duration with different CH₄/O₂ gas mixture ratios. RNA-Seq were performed using the paired-end sequencing technology according to standard illumina protocols by GENEWIZ, Inc. (Suzhou, China). The quality of RNA-Seq fastq data was checked using FastQC program¹. Data passing the quality control were imported into CLC Genomics Workbench (version 11.0) for RNA-Seq analysis to get the RPKM values with the reference genome from the MicroScope database². Gene expression normalization, ONE-WAY ANOVA analysis, and hierarchical clustering analysis were conducted using JMP Genomics (version 9.0) to identify differentially expressed genes at different conditions. Genes were determined to be significantly differentially expressed with a selection threshold of P-value ≤ 0.01 and log₂-fold change ≥ 1 (significant induction) or ≤ −1 (significant repression). Duplicate samples were used for each condition.

**RESULTS AND DISCUSSION**

**CH₄/O₂ Ratios Affected Cell Growth Especially on Logarithmic Phase**

Cell growth using various gas mixtures with different CH₄/O₂ ratios (from 0.28 to 5.24 mole/mole) were evaluated in 3-L bioreactors in duplicate. Three ratios within the CH₄ explosion

¹http://www.bioinformatics.babraham.ac.uk/projects/fastqc/
²http://www.genoscope.cns.fr/aga/microscope
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**FIGURE 1** | Cell mass of *M. buryatense* 5GB1 in the cultures using different CH$_4$/$O_2$ ratios of 0.28, 0.58, 0.93, 1.31, and 5.24 at different time points of 12, 24, 36, 48, 60, and 72 h post inoculation, respectively. The * and ** indicated this ratio exhibited significant difference compared to that under the CH$_4$/$O_2$ ratio of 0.93 with statistical significance of $P < 0.05$ and $P < 0.01$, respectively. DCW, dry cell weight.

The highest specific growth rate of *M. buryatense* 5GB1 reported before was 0.231 h$^{-1}$ obtained in cultures with CH$_4$/O$_2$ of 1.31 (20% CH$_4$ in air) (Puri et al., 2015). Our result, for the first time exhibited that the CH$_4$/O$_2$ ratio of 0.93 is optimal for cell growth of 5GB1, providing a DCW of 15 g/L along with a productivity of 0.3 g/L/h at the time point of 48 h. More importantly, *M. buryatense* 5GB1 exhibited a good state of growth on logarithmic phase, since the biomass at the time points of 24, 36, and 48 h especially the first two time points with the ratio of 0.93 was higher than those with other ratios. It is clear that the growth was significantly limited when cultured under the condition of 5% CH$_4$ in the air, which is mainly due to the insufficient carbon source of CH$_4$ during cultivation. However, higher CH$_4$ supply (e.g., 50% CH$_4$ with the ratio of 5.5) did not provide the best cell growth neither (Figure 1 and Supplementary Figure S1). Although cell grew better than that using lower carbon source in cultures under the gas ratio of 0.28, they grew poorly compared with other three conditions with 10~20% CH$_4$, which could be due to limited electron acceptor O$_2$ in the gas mixture.

Our findings thus demonstrated the importance of CH$_4$/O$_2$ ratios in the gas mixtures for balanced cell growth and production cost. The molar ratio of 0.93 was the best among all ratios, which can reach an OD value two-fold of other conditions within 24 h post-inoculation and had the highest OD value within 36~48 h fermentation time. The shorter time cells took to reach its highest growth and the lower CH$_4$ in the gas mixture could mean a higher productivity with less carbon supplied for economic biochemical production.

**Different CH$_4$/O$_2$ Ratios of Inlet Gas Affected the Accumulation of Key Metabolites and the Level of Reducing Power**

Since CH$_4$/O$_2$ ratios affected cell growth and the ratio of carbon and oxygen was crucial for balanced cell growth, the impact of different CH$_4$/O$_2$ ratios on the redox and the production of metabolite intermediates were investigated. Several key metabolites and enzymes during CH$_4$ oxidation and metabolism were analyzed as shown in Table 1, when *M. buryatense* 5GB1 was cultured with different gas supply ratios. It is obvious that the highest values of pyruvate and pMMO activity were observed in cultures with the gas ratio of 0.93. The higher values of pyruvate amount and pMMO activity were also observed in cultures with the gas ratio of 0.93, which is consistent with the result of growth rate in the same condition. These findings could be due to the fact that some key metabolites were not sufficiently synthesized when either CH$_4$ or O$_2$ was not supplied sufficiently under other conditions. Higher pyruvate (key precursor for TCA cycle) produced in cultures with gas ratio of 0.93 was...
corresponding with the higher DCW, which is in good agreement with previous report showing that TCA cycle is essential for the production of both cell mass and reducing power in 5GB1 (Fu et al., 2017).

The changes of CH$_4$/O$_2$ ratios could affect NADH/NAD$^+$ ratio, which determined the fluxes of metabolic pathways as well as the transcriptional level of many genes (Zhou et al., 2011). In this work, under the gas ratios of 1.31 and 0.93, the values of NADH/NAD$^+$ ratio were higher, and the highest value was 1.61, almost double that under the gas ratio of 5.24 (Table 1). This could be due to the different content of metabolite accumulation and the complicated metabolism in *M. bur natuur* 5GB1 with different gas ratios. This process was accompanied by changes of many genes at transcriptional level, so genomic and transcriptomic analyses could help further identify gene and metabolic pathways differentially expressed at different CH$_4$/O$_2$ ratios.

### Identification of Differentially Expressed Genes at Different Gas Supply Conditions

Transcriptomic study was further carried out to investigate the impact of different CH$_4$/O$_2$ ratios on global gene expression and the underlying mechanism of CH$_4$/O$_2$ ratio on cell growth and metabolism. As shown in Figure 1, the cell growth under the gas ratio of 0.93 exhibited obvious difference over other gas conditions at 24 h, so we adopted the samples for transcriptome analyses. Since different CH$_4$/O$_2$ ratios significantly influenced the cell growth and central metabolism, which result in different phenotypes, the relationship of five conditions with different CH$_4$/O$_2$ ratios were analyzed by hierarchical clustering using significantly differentially expressed genes based on the RNA-Seq result (Figure 2).

The result demonstrated that these samples can be divided into three groups: #3, #2&#4, and #1&#5, which not only indicates that the metabolism of #3 was different from others, but also suggests that the metabolism in cells was different under #2, #4, and #5 conditions despite that they exhibited similar growth (Figure 2A and Supplementary Figure S1). There were total of 1429, 733, and 592 DEGs by comparing #3 with other three conditions of #2, #4, and #5, respectively (Figure 2B and Supplementary Table S2). The DEGs of #3 versus #2, #4 or #5 were analyzed, and the result indicated that 337 common genes were significantly differentially expressed (log$_2$ fold change $\geq 1$, $p$-value $\leq 0.01$) with 160 down-regulated and 177 up-regulated ones (Supplementary Table S3). Among these significantly differentially expressed genes, the function of 106 genes was unknown (Supplementary Table S3), which needs further studies to confirm.

Our results also demonstrated that different CH$_4$/O$_2$ ratios affected both carbon and energy metabolism of *M. but natur* 5GB1 at transcriptional level (Supplementary Figure S2). Genes up-regulated were mainly involved in signal transduction, membrane transport, cell process, carbohydrate and energy metabolism, while genes down-regulated were majorly involved in metabolism of cofactors and vitamins, nucleotide metabolism and amino acid metabolism. This result indicated that the different phenotypes at condition of 0.93 compared with others were possibly ascribed to the function and expression levels of DEGs involved in carbon and nitrogen utilization, although further investigation is needed.

### Genes Involved in Nitrogen Fixation and Utilization Upregulated to Supply Sufficient Nitrogen for Enhanced Growth

*Methylomicobium but natur* 5GB1 has been reported to use nitrate, ammonia, and urea as nitrogen sources (Kaluzhnaya et al., 2001). Some methane-oxidizing bacteria (methanotrophs) are known to be capable of expressing nitrogenase and utilizing N$_2$ as the nitrogen source (Auman et al., 2001; Matsen et al., 2013). From the genome annotation and metabolic modeling, *M. but natur* 5GB1 was also predicted to be able to assimilate N$_2$ into ammonia, which can be used as a nitrogen source (de la Torre et al., 2015; Garg et al., 2018). The intriguing discovery from our transcriptomic analysis was that thirty-four nitrogen assimilation related genes were significantly up-regulated in the #3 condition compared with other conditions, including nitrogen fixation structural genes molybdenum nitrogenase genes (*nifHDK*), nitrogenase MoFe maturation genes (*NifPZ*), cofactor biosynthesis genes (*nifBEMNXQTVUY*), ferredoxin genes (*dxBCD*), as well as genes involved in electron supply (*rnCH*, *nifF*, *rxaBDEG*), nitrogen fixation regulation (*nifLA*, *glnGLK*), and post-translational modification (*draTG*) (Figure 3 and Supplementary Table S2).

In *P. stutzeri* A1501, the nitrogen regulatory cascade, AmyB-GlnK-NtrBC, senses the nitrogen signal and controls the expression of *nif*-specific regulatory proteins NifLA, in which the NiFA is an activator and the NifL is an anti-activator to control the expression of all other *nif* genes (Lin et al., 2015). In our RNA-Seq result, genes encoding PII protein GlnK, *nif*-specific regulatory protein NifLA, as well as GlnLG homologous to NtrBC were up-regulated under the optimal CH$_4$/O$_2$ ratio condition #3. The upregulation of nitrogen fixation regulatory genes indicated that the *nif* gene expression in 5GB1 may be tightly regulated at the transcriptional level. Correspondingly, the nitrogenase genes *nifHDK* were up-regulated trigged by this regulatory network under the CH$_4$/O$_2$ ratio of 0.93, which shown enhanced DCW (Figure 1). These findings in physiological and transcriptional levels may suggest that extra nitrogen source from N$_2$ can be assimilated supporting cell growth and metabolic activities after the nitrate source was exhausted, although further study is needed in the future.

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**Table 1** Quantification of the amount of total protein, pyruvate, NADH, and pMMO activity under different CH$_4$/O$_2$ ratio conditions at time point 24 h post inoculation.

| CH$_4$/O$_2$ ratio (mol) | 0.58 (#2) | 0.93 (#3) | 1.31 (#4) | 5.24 (#5) |
|-------------------------|-----------|-----------|-----------|-----------|
| Protein (g/L)           | 1.18 ± 0.07 | 1.00 ± 0.07 | 0.75 ± 0.05 | 0.88 ± 0.00 |
| Pyruvate (mg/g protein) | 1.13 ± 0.35 | 2.91 ± 0.33 | 1.03 ± 0.26 | 2.23 ± 0.49 |
| NADH/NAD$^+$            | 0.94 ± 0.10 | 1.28 ± 0.05 | 1.61 ± 0.32 | 0.89 ± 0.02 |
| pMMO (ug/min/g protein) | 6.04 ± 1.25 | 7.0 ± 0.06 | 4.65 ± 0.40 | 6.75 ± 1.68 |
While nitrogen fixation is an energy-consuming biological process (Bebout et al., 1993), it is with a high probability to cause the deficiency of biomass yield because of the activation of nitrogen fixation (de la Torre et al., 2015). In contrary, better cell growth and higher DCW were achieved along with the higher expression of nitrogen assimilation related genes, which indicated that 5GB1 likely shifted its metabolic flux to overcome the energy barrier for nitrogen fixation. Meanwhile, the nitrogenase post-translational genes (draTG), which can switch off or on the nitrogenase activity through adding or removing
ADP-ribose group to a specific arginine residue on nitrogenase (Moure et al., 2015), were also up-regulated in response to quickly modulating the activity of nitrogenase when the environmental ammonium concentration was suddenly changed (Masepohl and Hallenbeck, 2010). Additionally, $RxsABCDE$ genes are essential for nitrogen fixation, which are involved in transferring electrons to nitrogenase (Schmehl et al., 1993; Koo et al., 2003). The upregulation of $RxA$, $RxB$, $RxD$, $RxE$, and $RxG$ suggested a high nitrogenase activity achieved under condition #3, which could then facilitate the use of $N_2$ as nitrogen source for cell growth. Although higher activity of nitrogenase from #3 condition would promote the nitrogen assimilation, the intensive electron cost during nitrogen fixation process could cost cellular energy (Hoffman et al., 2014; Inomura et al., 2018). Unfortunately, research on balancing the $N_2$ assimilation and energy supply is still very limited.

It has been reported that higher $CH_4/O_2$ ratio can significantly affect nitrite removal in aerobic methanotrophs, and the highest denitrifying activity correlated with two nitrite reductases (NirK and NirS) was also observed when the molar ratio of $CH_4/O_2$ was 4.4 (Zhu et al., 2017). However, nitrite reductase (NirB) of $M. buryatense$ 5GB1 did not show significant differential expression, which may be due to the low $CH_4/O_2$ ratio (0.93) used in the #3 condition.

### The Upregulation of Methane Metabolism Genes Boosted Cell Growth

As an aerobic obligate methanotroph, $M. buryatense$ 5GB1 can oxidize $CH_4$ to methanol and formate in the beginning before flowing into RuMP and/or serine pathway for assimilations, providing energy for cell growth and metabolic processes at the same time (Troitsenko and Murrell, 2008). It has been reported that the major pathway for pyruvate production in Group I methanotrophic bacteria is the glycolysis pathway (Kalyuzhnaya et al., 2013). Since cell grew better in #3 condition, carbon flux to glycolysis could be enhanced considering the fact that genes related to glycolysis pathway were also upregulated in #3 condition, which is consistent with our result that the expression of $fbp$ (MBURv2_60345) was up-regulated greater than two folds (Figure 4).

Fructose-1,6-bisphosphatase is an essential enzyme in glycolysis pathway to hydrolyze D-fructose-1,6-bisphosphate to D-fructose-6-phosphate (Rittmann et al., 2003). Previous

![FIGURE 4](https://example.com/figure4.png)

**FIGURE 4** | Overview of the differential expression of genes encoding enzymes involving in nitrogen fixation, methane metabolism, ClpX system, and Pst system in $M. buryatense$ 5GB1 in condition #3 (0.93) compared with #2 (0.58), #4 (1.31), and #5 (5.24). Dotted arrow indicates multi-step reaction, while solid arrow represents one-step reaction. The numbers in shadow at right-side of protein represent the log$_2$-based changes of differentially expressed gene under condition #3 compared to #2, #4, or #5, respectively. The red shadow means upregulation, while the blue shadow means downregulation.
study confirmed that Fbp regulated the level of fructose-1,6-bisphosphate, one of the most important intermediates associating with the ATP generation (Li et al., 2013). Besides, Rittmann et al. (2003) investigated the relationship between fbp gene and cell growth by deleting it from the chromosome, and discovered that C. glutamicum WTΔfbp could not grow as well as the wild-type strain. Although Entner–Doudoroff pathway was suggested to be a potential solution to increase carbon flux for methanotrophic bacteria (Kalyuzhnaya et al., 2015), our result suggests that the enhanced expression of fbp gene under the optimal CH4/O2 ratio can accelerate the conversion of CH4 into pyruvate and NADH in M. buryatense 5GB1.

**Genes Involved in Key Intermediate Synthesis Upregulated for Enhancing Precursor Supply to TCA Cycle**

In both aerobic and anaerobic microorganisms, ATP is an important energy source to sustain microbial growth (Khmelenina et al., 2011). Aerobic microorganisms mainly rely on TCA cycle and oxidative phosphorylation, which provide 95% ATP (Khmelenina et al., 2011). Comparing the transcriptional analysis of #3 with other conditions, two genes (por and oadG) involved in Acetyl-CoA (AcCoA) and oxaloacetate synthesis were up-regulated (Figure 4), which encode putative oxidoreductase and putative oxaloacetate decarboxylase gamma chain, respectively. Pyruvate was converted to oxaloacetate and AcCoA, which then entered into TCA cycle. The higher production of DCW and NADH observed in #3 condition was therefore consistent with the previous prediction that TCA cycle is responsive for both biomass synthesis and reducing power generation in M. buryatense 5GB1 (Fu et al., 2017). This result is also in agreement with pervious study reporting the additional NADH input can support the direct coupling between methane oxidation and methanol oxidation for better cell growth (de la Torre et al., 2015).

**Genes Involved in ClpX Upregulated to Promote Cell Growth by Controlling ATP Hydrolysis for Energy Supply**

As a AAA + protease (ATPase associated with a variety of cellular activities), ClpX hydrolyses ATP to provide energy for targeting the misfolded protein for subsequent degradation during protein synthesis in cell growth process (Keiler et al., 1996; Gottesman et al., 1998; Moore and Sauer, 2007). In addition, the energy produced by hydrolysis of ATP is the main energy source of the ClpXP protease complex in protein degradation. Therefore, upregulated ClpX activated more energy supporting in protein synthesis during cell growth under the #3 condition. Previous studies in Bacillus subtilis showed that the cells were damaged readily with the corresponding decrease of the growth rate in the absence of either ClpX or ClpP under stressful environmental conditions (Zhang and Zuber, 2007), which suggests the upregulation of ClpX could be beneficial for protecting cells from damage and promoting the cell growth as what we observed under #3 condition in this study.

**CONCLUSION**

In this work, the impact of different CH4/O2 ratios on the growth and global gene expression of M. buryatense 5GB1 was investigated, and our result indicated that the CH4/O2 mole ratio of 0.93 is optimal for cell growth, pyruvate accumulation, nutrient utilization and energy supply. Meanwhile, genes related to nitrogen fixation and methane metabolism were significantly up-regulated, which suggest that M. buryatense 5GB1 is capable of utilizing N2 as the nitrogen source to reduce the production cost. In addition, by controlling CH4/O2 ratios during the cultivation, the level of phosphate uptake for enhancing cell growth of methanotrophs was improved to maintain an optimal intracellular Pi pool for growth. Our work thus provides novel insights in terms of the influences of CH4/O2 ratios of gas supply on transcriptional level of key genes in methanotrophic bacteria, and candidate gene targets for future metabolic engineering in this promising industrial methanotrophic microorganism.
DATA AVAILABILITY STATEMENT

The authors declare that all the data supporting the findings of this study are available within the manuscript and its Supplementary Information files from the corresponding author on request. The RNA-Seq raw data was deposited at Sequence Read Archive (SRA) database with the BioProject accession number PRJNA597286.

AUTHOR CONTRIBUTIONS

QF and SY conceived the work, and provided conceptual advice with inputs from all authors. LH designed and performed the experiments, analyzed the data, and wrote the manuscript. YY handled and analyzed the transcriptomic data with help from YC. XY provided conceptual advice. LH, YY, QF, and SY wrote the manuscript, and prepared figures and tables. All authors contributed to data analyses, read, revised and approved the final manuscript.

FUNDING

This work was supported by the National Key R&D Programs of China (2018YFA0901500) and National Natural Science Foundation of China (21878241, 21978071, and U1932141). This work was also supported by Technical Innovation Special Fund of Hubei Province (2019AHB055 and 2018ACA149), and the Open Project Funding of State Key Laboratory of Biocatalysis and Enzyme Engineering of Hubei University.

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AUTHOR CONTRIBUTIONS

QF and SY conceived the work, and provided conceptual advice with inputs from all authors. LH designed and performed the experiments, analyzed the data, and wrote the manuscript. YY handled and analyzed the transcriptomic data with help from YC. XY provided conceptual advice. LH, YY, QF, and SY wrote the manuscript, and prepared figures and tables. All authors contributed to data analyses, read, revised and approved the final manuscript.

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FIGURE S1 | Growth curve of M. buryatense 5GB1 under different CH4/O2 ratios of 0.28, 0.58, 0.93, 1.31, and 5.24, respectively.

FIGURE S2 | KEGG enrichment of differentially expressed genes (DEGs) in M. buryatense 5GB1 by comparing 0.93/0.5 with 0.58/0.2, 1.31/0.4, and 5.24/0.5. (a) Common up-regulated genes in KEGG pathway. (b) common down-regulated genes in KEGG pathway.

TABLE S1 | Log2 OD growth data for specific growth rate under different CH4/air ratio conditions.

TABLE S2 | Differentially expressed genes of #3 (15% CH4 in air) versus #1 (10% CH4 in air).

TABLE S3 | Differentially expressed genes of #3 (15% CH4 in air) versus #1 (10% CH4 in air).

This work was also supported by Technical Innovation Special Fund of Hubei Province (2019AHB055 and 2018ACA149), and the Open Project Funding of State Key Laboratory of Biocatalysis and Enzyme Engineering of Hubei University.

SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fbioe.2020.00263/full#supplementary-material
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Conflict of Interest: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.