Metabolic Engineering Strategies for Co-Utilization of Carbon Sources in Microbes

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Abstract: Co-utilization of carbon sources in microbes is an important topic in metabolic engineering research. It is not only a way to reduce microbial production costs but also an attempt for either improving the yields of target products or decreasing the formation of byproducts. However, there are barriers in co-utilization of carbon sources in microbes, such as carbon catabolite repression. To overcome the barriers, different metabolic engineering strategies have been developed, such as inactivation of the phosphotransferase system and rewiring carbon assimilation pathways. This review summarizes the most recent developments of different strategies that support microbes to utilize two or more carbon sources simultaneously. The main content focuses on the co-utilization of glucose and pentoses, major sugars in lignocellulose.

Keywords: metabolic engineering strategies; co-utilization; lignocellulosic biomass

1. Introduction

Microbial production using renewable and economical carbon sources has been highly preferred to address the increasing concerns on the global energy shortage and climate change. Lignocellulosic biomass has become an attractive carbon source for microbial production in recent years, because its hydrolysates substantially offer several simple sugars, such as glucose and pentoses, that can be readily utilized by microorganisms [1]. In addition, other carbon sources in hydrolysates, such as acetate, can also be used for supporting microbial growth and production [2,3]. However, achieving high-yield and cost-effective microbial production of target products from lignocellulosic resources still have challenges from the laboratory scale to the industrial scale, one of which is how to achieve efficient utilization of the mixed sugars. Most microbes consume these mixed sugars sequentially or selectively due to the carbon catabolite repression (CCR), which prolongs the microbial production process and reduces the efficiency of carbon conversion. In order to solve this issue, co-utilization of mixed carbon sources in microbes is highly preferred.

For most microbes, CCR leads to preferential utilization of one carbon source when there are two or more carbon sources in the medium. CCR is an essential regulatory system which can guarantee microbes to obtain specific carbon sources efficiently and survive in competing environments [4]. However, the selective utilization of carbon sources caused by CCR is sometimes a barrier for simultaneous utilization of different carbon sources in bioengineering process. To overcome this barrier, metabolic engineering strategies have been developed against different mechanisms of CCR. This review will summarize the recent efforts devoted to the strategies for eliminating CCR caused by different mixed carbon sources with the emphasis on co-utilization of glucose and pentoses.
2. Glucose and Pentoses from Lignocellulosic Biomass

Lignocellulosic biomass is cheap and renewable for bio-based chemical production. The worldwide production of lignocellulose is estimated to be $1.8 \times 10^{11}$ tons/year [5]. Lignocellulose is composed of the heterogeneous complex of cellulose (30%–50%), hemicellulose (15%–35%), and lignin (10%–30%) [6]. Cellulose is a polymer of glucose, while hemicellulose is a heteropolymer of glucose and pentoses, such as xylose and arabinose [7]. Lignin is a complex of polymers of cross-linked phenylpropene units. After hydrolysis of lignocellulose, glucose and pentoses are the predominant sugars as the carbon sources. Co-utilization of glucose and pentoses can be realized via different strategies, depending on the features of microbes (Table 1).

*Escherichia coli*, a widely used industrial production species, possesses CCR that is mainly mediated by inducer exclusion coupled to the phosphoenolpyruvate-dependent sugar phosphotransferase system (PTS) [7–9]. Inducer exclusion is a process that the dephosphorylated $EIIA^{glc}$ could inhibit the activity of non-PTS sugar transport system in present of glucose (Figure 1). To disable this mechanism of CCR, *E. coli* PTS disruption strains were constructed, which could consume glucose and xylose simultaneously to produce various polyhydroxyalkanoates (PHAs) [10]. The $ptsG$ mutant showed better growth and better utilization than wild-type and the accumulation of short-chain-length PHAs was improved from 7.1% of cellular dry weight (CDW) to 11.5% of CDW after 48 hours [10]. In addition to the deletion of $ptsG$, disruption of $ptsHicrr$-encoding sugar non-specific protein constituents of the PTS could also relieve the CCR [11]. However, deficiency of PTS can affect the transportation and phosphorylation of glucose because it is the main method for glucose uptake in *E. coli* [12]. To address this problem, the $glf$-encoding glucose facilitator of *Z. mobilis* was integrated into the genome for enhancement of the glucose-utilizing rate. Meanwhile, increasing the activity of the pentose phosphate pathway strengthened the utilization of xylose. With this strategy, the resulting strain could produce 29 g/L ethanol in around 16 h, which accounted for 97% of the theoretical ethanol yield, by consuming glucose and xylose simultaneously [13]. In addition, global transcriptional regulators can also mediate CCR, like the cAMP receptor protein (CRP). Replacement of *E. coli*-native CRP with a cyclic AMP-independent mutant increased the xylose utilization in the presence of glucose. The molar ratio of xylitol was improved after deletion of the xylB gene and overexpression of heterologous xylose reductase or xylitol dehydrogenase (Figure 2). By overexpression of xylose reductase from *Candida boidinii*, the concentration of xylitol was about 38 g/L in a fermenter with a 10 L working volume [14]. Among the sugars in lignocellulosic hydrolysates, *E. coli* prefers to consume arabinose than xylose in the absence of glucose. This is because arabinose transcriptional regulator (AraC) can also inhibit xylose transcriptional regulator (XylR), leading to the repression of xylose metabolism (Figure 1). Hence, deletion of the araC gene can achieve the co-utilization of xylose and arabinose [15]. Kim *et al.* constructed an evolved strain, GX50, containing araC deletion and pentose operon constitutive expression, which could be used as a platform strain for co-utilization of glucose and xylose for xylitol production [15]. Another strategy to relieve arabinose-xylose diauxie is to optimize the XylR expression level in *E. coli*. The engineered strain could produce 36% more ethanol than the wild-type in 72 h [16].
Table 1. The strategies for co-utilization of glucose and pentoses from lignocellulosic biomass in different microbes.

| Microbe                  | Strategy                                                                 | Carbohydrates          | Product       | References |
|--------------------------|--------------------------------------------------------------------------|------------------------|---------------|------------|
| **Escherichia coli**     | Inactivation of ptsG gene                                                | Glucose and xylose     | Ethanol       | [13]       |
|                          | Deletion of ptsG gene                                                    | Glucose and xylose     | Polyhydroxyalkanoates | [10]   |
|                          | Replacement of native cyclic AMP receptor protein with a cyclic AMP-independent mutant | Glucose and xylose     | Xylitol       | [14]       |
|                          | Engineering of chb and asc operons and adaptive evolution                | Cellulbiose and xylose | -             | [37]       |
|                          | Expression of xylR at the appropriate level                             | Xylose and arabinose   | Ethanol       | [16]       |
|                          | Deletion of araC, constitutive expression of genes required for pentose metabolism and adaptive evolution | Glucose and xylose     | Xylitol       | [15]       |
|                          | Inactivation of ptsHlcr gene; overexpression of galP                    | Glucose, xylose, and arabinose | Cinnamic and \(\pm\)-hydroxycinnamic acid | [11] |
| **Saccharomyces cerevisiae** | Expression of the xylose isomerase; Overexpression of XKS1, RPE1, RKL1, TAL1, and TKL1; Deletion of GRE3 and COX4 genes; Adaptive evolution | Glucose and xylose     | Ethanol       | [38]       |
|                          | Construction of a growth-based screening system for mutant hexose transporters | Glucose and xylose     | -             | [19]       |
|                          | Deletion of D-ribulose-5-phosphate 3-epimerase                           | Glucose and xylose     | Ethanol       | [20]       |
|                          | Maintaining glucose in the useful concentration range in fed-batch reaction | Glucose and xylose     | Ethanol       | [39]       |
|                          | Expression of xylose reductase, xylitol dehydrogenase and xylulokinase; Engineering of hexose transporters | Glucose and xylose     | Ethanol       | [18]       |
|                          | Evolutionary engineering strategy based on repeated batch cultivation with repeated cycles of consecutive growth | Glucose, xylose, and arabinose | Ethanol       | [24]       |
|                          | Evolutionary engineering via continuous culture using xylose and arabinose as limiting carbon sources | Xylose and arabinose   | Ethanol       | [23]       |
|                          | Expression of a cellulodextrin transporter, intracellular \(\beta\)-glucosidase and xylose reductase and optimization of the expression | Cellulbiose and xylose | Xylitol Ethanol | [22] |
|                          | Integration of the fermentation pathways of cellulbiose and xylose and an acetic acid reduction pathway | Cellulbiose, xylose, and acetic acid | Ethanol       | [40]       |
| **Saccharomyces pastorianus** | Co-expression of all three classes of cellulase genes with the xyl1/xdh1/XKS genes | Xylose and cellulose   | Alcohol       | [41]       |
| **Clostridium acetobutylicum** | CcpA mutagenesis                                                       | Glucose and xylose     | Acetone, Butanol, Ethanol | [25,26] |
|                          | Inactivation of \(g\&G\) and overexpression of the rate-limiting steps in xylose pathway | Glucose, xylose, and arabinose | Acetone, Butanol, Ethanol | [27] |


Table 1. Cont.

| Microbe                     | Strategy                                      | Carbohydrates       | Product          | References |
|-----------------------------|-----------------------------------------------|---------------------|------------------|------------|
| *Clostridium sp.* Strain BOH3 | Expression of xylose isomerase and xylulokinase | Glucose and xylose  | Butanol Riboflavin | [28]       |
| *Clostridium tyrobutyricum* | Overexpression of *xylT*, *xylA*, and *xylB*   | Glucose and xylose  | n-Butanol        | [42]       |
| *Corynebacterium glutamicum*| Expression of *araBAD* operon and/or the *xylA* gene from *Escherichia coli* | Glucose, xylose, and arabinose | Amino Acid       | [43]       |
| *Enterococcus munditi QU 25* | Maintaining the glucose concentration lower than 25 g/L | Glucose and xylose  | L-Lactic acid    | [44]       |
| *Cryptococcus curvatus*     | Decreasing glucose concentration               | Glucose, xylose, and cellobiose | Microbial lipid  | [45]       |
was used to produce ethanol, and the affinity of different transporters affected the consumption of xylose and glucose during the fermentation by the strain lacking \textit{RPE1} genome, and xylose for ethanol production [20]. After the xylose utilization pathway was integrated into the genome, \textit{RPE1} was deleted to reduce the flow of carbon from glucose into the pentose phosphate pathway. The ratio of xylose and glucose consumption was about 1:10 during the fermentation by the strain lacking \textit{RPE1} [20]. To avoid the inhibition of glucose and insufficient supply of NAD(P)H, the expression of \textit{HXT1} was chosen to co-utilize with xylose. The cellobiose metabolic pathway and xylose reductase were usually begun with the overexpression of the xylose utilization pathway. The introduction of xylose reductase, xylitol dehydrogenase, and xylulokinase in \textit{Saccharomyces cerevisiae} is also a potential host for biotechnological production of fuels and chemicals [17]. It can utilize glucose, but not xylose, as a carbon source. Therefore, most strategies usually began with the overexpression of the xylose utilization pathway. The introduction of xylose reductase, xylitol dehydrogenase, and xylulokinase in \textit{S. cerevisiae} could allow the strain to co-utilize glucose and xylose after adjusting the expression of hexose transporter permeases [18]. This strategy was used to produce ethanol, and the affinity of different transporters affected the consumption of sugars and the production of ethanol. The strain with overexpression of the transporter encoded by \textit{HXT7} could achieve the maximal consumption of sugars and ethanol production rate in the glucose and xylose co-fermentation process [18]. The ethanol production was higher than that with either glucose or xylose. However, it cannot consume xylose significantly in a xylose-enriched medium, whereas the \textit{HXT7} permease would be a better choice to allow the xylose uptake rate to be the same as that of glucose [18]. In addition to adjusting the expression of hexose transporter permeases for decreasing glucose repression, the mutations on hexose transporters could also prevent the competitive inhibition from glucose. The mutation sites were designed on hexose transporters in order to transport xylose rather than glucose [19]. This research demonstrated that the mutant Gal2-N376F can transport xylose with the highest affinity and cannot transport any hexose at all [19]. Furthermore, the deletion of \textit{d-ribulose-5-phosphate 3-epimerase (RPE1)} is another way to achieve the co-utilization of glucose and xylose for ethanol production [20]. After the xylose utilization pathway was integrated into the genome, \textit{RPE1} was deleted to reduce the flow of carbon from glucose into the pentose phosphate pathway. The ratio of xylose and glucose consumption was about 1:10 during the fermentation by the strain lacking \textit{RPE1} [20]. To avoid the inhibition of glucose and insufficient supply of NAD(P)H, the co-utilization of glucose and xylose co-fermentation process [18]. The ethanol production was higher than that with either glucose or xylose. However, it cannot consume xylose significantly in a xylose-enriched medium, whereas the \textit{HXT7} permease would be a better choice to allow the xylose uptake rate to be the same as that of glucose [18]. In addition to adjusting the expression of hexose transporter permeases for decreasing glucose repression, the mutations on hexose transporters could also prevent the competitive inhibition from glucose. The mutation sites were designed on hexose transporters in order to transport xylose rather than glucose [19]. This research demonstrated that the mutant Gal2-N376F can transport xylose with the highest affinity and cannot transport any hexose at all [19]. Furthermore, the deletion of \textit{d-ribulose-5-phosphate 3-epimerase (RPE1)} is another way to achieve the co-utilization of glucose and xylose for ethanol production [20]. After the xylose utilization pathway was integrated into the genome, \textit{RPE1} was deleted to reduce the flow of carbon from glucose into the pentose phosphate pathway. The ratio of xylose and glucose consumption was about 1:10 during the fermentation by the strain lacking \textit{RPE1} [20]. To avoid the inhibition of glucose and insufficient supply of NAD(P)H.
for xylose reductase, cellulose, a disaccharide of glucose, was chosen to co-utilize with xylose. The cellulose metabolic pathway and xylose reductase were introduced into *S. cerevisiae* for utilization of cellulose and xylose simultaneously [21]. The engineered strain could produce 19 g/L xylitol with co-fermentation of cellulose and xylose within 48 h, while the titer of xylitol was only 13 g/L, with sequential utilization of glucose and xylose [21]. When the initial concentrations of the cellulose and xylose were higher, the titer of the xylitol was higher. By overexpression of NADPH-regenerating enzymes, the productivity of xylitol increased by 37%–63% [21]. Zha *et al.* used the same strategy to realize the co-fermentation of these two sugars. Through the optimization of exogenous gene expression, two sugars had similar consumption rates and the production of xylitol increased by 85.7% in 120 h [22]. Moreover, evolutionary engineering approaches were employed to make strains suitable for co-utilizing the mixed sugars in the medium. After the continuous culture for evolution, the evolved strain that carried the xylose and arabinoase pathways could produce ethanol with the yield of 0.29 g per consumed xylose (g) at 120 h [23]. Wisselink *et al.* improved the strain by a novel evolutionary engineering strategy which consists of “repeated batch cultivation with repeated cycles of consecutive growth in three media with different compositions”. The evolved strain had a higher yield of biomass and exhibited a high yield of ethanol, 0.44 g/g of total sugars [24].

**Figure 2.** Metabolic engineering of xylitol production in *E. coli*. Xylose is transferred into *E.coli* cells by xylose transporters. Xylose can be converted into xylulose by native xylose isomerase, which is subsequently phosphorylated to xylulose-5-phosphate by xylulose kinase. Xylulose-5-phosphate is an intermediate of the pentose phosphate pathway. Introduction of heterologous xylose reductase or reversible xylitol dehydrogenase can achieve the xylitol production. Xylose reductase reduces xylose to xylitol using NADH or NADPH, and xylitol dehydrogenase reduces xylulose to xylitol using NADH. Blue lines represent the native pathway in *E.coli* and red lines represent the heterologous pathway for xylitol production.

In addition, co-utilization of glucose and major pentoses from lignocellulosic biomass has also been applied in other microbes. For the industrial solvent-producing strain *Clostridium acetobutylicum*, the disruption of the gene *ccpA*, which encodes a catabolite control protein, could lead to the simultaneous utilization of glucose and xylose. With this strategy, the engineered strain could produce acetone, butanol, and ethanol with titers of 4.94, 12.05, and 1.04 g/L, respectively, in around 36 h [25]. Based on the CcpA molecular modulation, Wu constructed CcpA mutants to overcome CCR and
overexpressed sol genes to accelerate the sugar consumption and product formation. Finally, the engineered strain could produce 5.01 g/L acetone, 8.61 g/L butanol, and 0.94 g/L ethanol within 60 h [26]. Another way to release CCR in C. acetobutylicum was inactivation of glcG gene-encoding enzyme II in PTS [27]. By disruption of the glcG gene and overexpression of xylose pathway, the strain could achieve 16.06 g/L as the total titer of acetone, butanol, and ethanol in the mixtures of glucose, xylose, and arabinose. Furthermore, activating xylose pathway or maintaining a low concentration of glucose can also lead to the co-utilization of glucose and pentoses from the lignocellulosic hydrolysates [28].

3. Glucose and Galactose from Marine Plant Biomass

In addition to lignocellulosic biomass, marine plant biomass has become into another renewable and cost-effective feedstock for microbial production [29]. Marine plant biomass is basically marine macroalgae, which is a kind of potential feedstock with many advantages. Compared with lignocellulosic biomass, marine macroalgae have higher photon conversion efficiency and carbon dioxide fixation rate. Moreover, marine macroalgae lack the recalcitrant cell wall structure so that the enzymatic hydrolysis proceeds much easier in marine macroalgae than lignocellulosic biomass [29,30]. Especially, the red algae are one of the different types of marine macroalgae. They have high fermentable carbohydrate content, including glucose and galactose, which are released from cellulose and agarose in red algae, respectively [17,31]. CCR is still the obstacle in the co-utilization of glucose and galactose. To bypass the repression of glucose, cellobiose, and galactose were co-fermented by engineered S. cerevisiae, in which genes encoding a cellodextrin transporter and an intracellular β-glucosidase were over-expressed [30]. Finally, the cell growth, titer, and yield of ethanol were increased, compared with sequential utilization of glucose and galactose [30]. Since the consumption rate of cellobiose was slow, glucose was co-utilized with galactose in S. cerevisiae to produce enantiopure (2R,3R)-butanediol with 70% of the theoretical maximum yield. The repression of glucose was alleviated after introducing the internal truncated MTH1 that encodes a transcription factor related to hexose transport [17]. Further improvement on the glucose consumption rate was realized by adaptive evolution for practical use [17]. In addition to S. cerevisiae, the galactose metabolic pathway was rebuilt with synthetic components, such as promoters and terminators in E. coli, for the co-fermentation of glucose and galactose [32]. The uptake of galactose could increase by 53.1% in the final strain compared with the wild-type strain.

4. Glucose and Non-Carbohydrates

Simultaneous utilization of glucose and non-carbohydrates such as glycerol and acetate is another way to achieve highly-efficient biological production. Glycerol is a renewable and abundant feedstock with great availability, low price, and high degree of reduction [33]. The co-utilization of glucose and glycerol can be realized by deletion of the ptsHIcrr operon of PTS in E.coli [34]. The strain PB12 which could grow fast on glucose was isolated to produce aromatic compounds from glucose and other carbon sources. In the mixture of glucose and glycerol, the yield of aromatic compounds was higher and the titer of byproduct acetate was much lower [34]. Shiue et al. constructed an E. coli Δpgi Δzwf mutant strain for D-glucaric acid production. The deletion of genes pgi and zwf could not only eliminate catabolite repression but also improve the strain’s ability to produce glucose-derived products. As the results, the titer of D-glucaric acid was higher in the E. coli Δpgi Δzwf mutant strain and the yield of D-glucaric acid on glucose increased by nine-fold in the medium with glucose and glycerol [35]. For the co-utilization of glucose and acetate, acetate could play different roles. To increase the intracellular PEP for aromatics production, the genes pykA, pykF, and ppsA were deleted in an E. coli ptsHIcrr− strain. As a consequence, the mutant strain could not grow on either glucose or acetate but could grow in the presence of both carbon sources. Herein, acetate played a role as a gluconeogenic substrate. At last, the yield of total aromatics could increase by four-fold, compared with the control strain [36]. To increase the carbon yield and balance the co-factor, acetate was used as the substrate to
generate acetyl-CoA for isobutyl acetate production. The deletion of native genes for conversion of pyruvate to acetyl-CoA, and the introduction of an acetate-assimilating pathway, led to the theoretical maximum carbon yield of isobutyl acetate increased from 67% to 75% and the practical yield improved from 35% to 59% after 120 h of production [2].

5. Conclusions and Future Perspectives

The preferential utilization of glucose is one of the major limitations for the co-utilization of carbon sources in microbes. The current metabolic engineering strategies to address this limitation mainly focus on the following aspects: (1) eliminating the CCR caused by glucose transportation; (2) enhancing the efficiency of other carbohydrate transportation; and (3) introducing exogenous genes or pathways to improve the efficiency of co-utilization of carbon sources. Although the mechanisms of CCR in different microbes are various, the principles used to relieve CCR are similar. Overall, co-utilization of carbon sources is essential for low-cost microbial production. The existing strategies can be further combined and optimized to achieve target production based upon the carbon sources to be used.

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