Reconstruction and analysis of carbon metabolic pathway of *Ketogulonicigenium vulgare* SPU B805 by genome and transcriptome

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*Ketogulonicigenium vulgare* has been widely used in vitamin C two-step fermentation. Four *K. vulgare* strains (WSH-001, Y25, Hbe602 and SKV) have been completely genome-sequenced, however, less attention was paid to elucidate the reason for the differences in 2-KGA yield on genetic level. Here, a novel *K. vulgare* SPU B805 with higher 2-keto-L-gulonic acid (2-KGA) yield, was genome-sequenced to confirm harboring one circular chromosome with plasmid free. Comparative genome analyses showed that the absence of plasmid 2 was an important factor for its high 2-KGA productivity. The amino acid biosynthetic pathways in strain SPU B805 are much more complete than those in other *K. vulgare* strains. Meanwhile, strain SPU B805 harbored a complete PPP and TCA route, as well as a disabled EMP and ED pathway, same as to strain SKV, whereas strain WSH-001, Y25 and Hbe602 harbored complete PPP, ED, TCA pathway and a nonfunctional EMP pathway. The transcriptome of strain SPU B805 validated the carbon metabolism in cytoplasm mainly through the PPP pathway due to its higher transcriptional levels. This is the first time to elucidate the underlying mechanism for the difference in 2-KGA yield, and it is of great significance for strain improvement in the industrial fermentation.

*Ketogulonicigenium vulgare* has been widely used for the bioconversion of L-sorbose to 2-keto-L-gulonic acid (2-KGA), the key precursor of vitamin C in the industrial production. In this process, *K. vulgare* is usually co-cultured with companion strain, *Bacillus megatherium*, which is generally considered to provide nutrients or metabolic substances for the growth of *K. vulgare*.¹ The known 2-KGA-producing strains, *K. vulgare* Y25, *K. vulgare* WSH-001, *K. vulgare* SKV and *K. vulgare* Hbe602 have been completely genome-sequenced and reported²-⁵. The genome-scale metabolic model of *K. vulgare* WSH-001 was reconstructed and analyzed by flux balance analysis (FBA).⁶ Total 116 genes, located on the chromosome, were identified as the essential genes for the growth of *K. vulgare* WSH-001⁶. Among them, approximately half of the essential genes participated in the metabolism of nucleotides and amino acids. Some specific amino acids significantly affected the growth of cells and the accumulation of 2-KGA⁷,⁸; moreover, the uptake rates of amino acids had a linear positive correlation with cell growth rate⁹. Additional, *K. vulgare* WSH-001 had a powerful transporting system to absorb nutrients and various amino acids³. Although many researches on metabolic pathways have been carried out, the understanding of the *K. vulgare* is still a partial picture due to its unique growth characters. Therefore, it is necessary to discover the panorama of whole metabolism features of different 2-KGA-producing strains.

In prokaryotes, sugar oxidation pathways, the most common of which are the Embden-Meyerhoff-Parnass (EMP) pathway, the Entner-Doudoroff (ED) pathway and the pentose phosphate pathway (PPP)¹⁰. However, these routes are different in the ATP yield, cofactors and chemical intermediates. The EMP pathway is a process by which glucose is not completely oxidized to pyruvate under anaerobic conditions¹⁰. In the classical EMP pathway, glucose is phosphorylated twice and cleaved into glyceraldehyde 3-phosphate (G3P) and dihydroxyacetone phosphate (DHAP), and then two G3P are formed by the reversible interconversion between DHAP and G3P. G3P is metabolized to pyruvate in the subsequent glycolysis and coupled with ATP

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production through the substrate-level phosphorylation. So, one mole of glucose is oxidized to produce a net of two moles of ATP and two moles of NADH through the EMP pathway.

The ED pathway is a wildly distributed glucose catabolic route in nature\(^\text{11,12}\). In the ED pathway, glucose is phosphorylated to form glucose 6-phosphate (G6P), and then oxidized to 6-phosphogluconate (6PG) by glucose-6-phosphate dehydrogenase (Zwf)\(^\text{13}\). 6PG is then oxidized to 2-keto-3-deoxy-6-phosphogluconic acid (KDPG) by KDPG aldolase (Eda), which is then cleaved to one pyruvate and one G3P. G3P is used to produce ATP through substrate-level phosphorylation in latter glycolysis. KDPG is exclusively generated in the ED pathway and Eda is the key enzyme unique to the ED pathway\(^\text{15}\). The ED pathway can achieve the same conversion rate as the EMP pathway do, but fewer enzymes are required\(^\text{16}\). During the process, only one mole of ATP as well as one mole of NADH and one mole of NADP are produced when one mole of glucoses is decomposed.

The PPP pathway, as a fundamental component in the carbon metabolism, can be divided into the oxidative branch and non-oxidative branch\(^\text{15}\). In the oxidative branch, G6P is metabolized into ribulose 5-phosphate (Ru5P), NADPH and carbon dioxide by the enzyme Zwf and 6-phosphate glucuronate dehydrogenase (Gnd). In the non-oxidative branch, the glycolytic intermediates fructose 6-phosphate (F6P), G3P and sedoheptulose 7-phosphate (STP) are rearranged to generate ribose 5-phosphate (R5P) and other sugar phosphate for the subsequent biosynthesis of nucleic acids and amino acids\(^\text{17}\). Zwf plays an important role in the fine-tuning of PPP pathway, but it is not the unique to the PPP pathway because of its product 6PG can be channeled into ED pathway for further metabolism\(^\text{15}\). Unlike the EMP and ED pathway, one mole of glucose is decomposed to produce a net of two moles of NADPH through the PPP pathway.

Although several strains of _K. vulgaris_ have been genome-sequenced, annotated and published\(^\text{2–5}\), but their attentions were mainly focused on the elucidation of the metabolic pathway of 2-KGA biosynthesis and symbiosis mechanism with companion strains\(^\text{14}\). Less attempt of different 2-KGA yield on the genetic level in different strains. Therefore, in this work we studied the differences in genome and carbon metabolism pathway between *K. vulgaris* SPU B805 and the four reported *K. vulgaris* strains to understand the variations in the growth and 2-KGA production, as well as the advantages of *K. vulgaris* SPU B805. *K. vulgaris* SPU B805, a new 2-KGA-producing strain, contains only a circular chromosome without any plasmid. The COG and RAST annotation revealed that the genes encoding glucuronate 2-dehydrogenase (GA2DH), one of L-sorbose dehydrogenase (SNDH) and the key enzymes (Eda and Edd) unique to the ED pathway were all absent from the genome of strain SPU B805, which may be the underling mechanism that contributes to its high 2-KGA productivity. The complete pathways of glycine, serine, threonine and proline in SPU B805, whereas absent from other *K. vulgaris* strains, may be another reason for high 2-KGA yield of SPU B805. Meanwhile, the comparative genome analysis showed that strain SPU B805 and SKV contained a complete PPP and TCA cycle, as well as a disabled EMP and ED pathway. Whereas, strain WSH-001, Y25 and Hbe602 harbor complete PPP, ED, TCA pathways and a nonfunctional EMP pathway. The transcriptome of strain SPU B805 validated that the PPP pathway was of major importance for cytoplasmic carbon oxidation. Our findings provide a complementary platform for understanding the underlying mechanism for the different central carbon metabolic features and 2-KGA yield of different *K. vulgaris* strains. It also provides a useful framework for further physiology and gene engineering research. This is of great significance for _K. vulgaris_ strain improvement and genetic engineering in the 2-KGA industrial fermentation.

**Results**

**2-KGA yield of K. vulgaris SPU B805 in the co-culture system**. *K. vulgaris* SPU B805 was fermented with _B. megatherrium_ SPU B806 in 5-L bioreactor for 2-KGA production. The 2-KGA production was 82.74 ± 0.77 g/L with a 95.97% conversion rate at the end of the fermentation, and the absorbance at 660 nm was 5.63 ± 0.17, which means the cell concentration is 52 ± 1.4 × 10⁶ CFU/mL (Table 1; Supplementary Fig. S1). Because no fermentation data of Y25 and SKV, the 2-KGA production abilities were compared between Hbe602, WSH-001 and SPU B805. As shown in Table 1, strain SPU B805 showed significantly differences (p < 0.001) compared with *K. vulgaris* Hbe602 in the 2-KGA yield (increased by 25.55%, increment of 16.84 mg/ml), productivity (increased by 241.82%), 2-KGA conversion rate (increased by 25.55%) and fermentation cycle (decreased by 172.73%); similarly, strain SPU B805 showed significantly differences (p < 0.001) compared with *K. vulgaris* WSH-001 in the 2-KGA yield (increased by 23.81%, increment of 15.91 mg/ml), productivity (increased by 102.15%), 2-KGA conversion rate (increased by 8.48%) and fermentation cycle (decreased by 59.09%). The cell growth (OD₆₀₀) of SPU B805 was also significantly higher than that of WSH-001 (p < 0.05). These superior qualities aroused our great interest to understand the mechanism. So, the genome of *K. vulgaris* SPU B805 was sequenced, annotated and compared with all of the reported 2-KGA-producing strains (SKV, Hbe602, WSH-001 and Y25).

**Genomic features of K. vulgaris SPU B805.** The complete genome of *K. vulgaris* SPU B805 was sequenced and deposited in the GenBank database with the accession number CP017622. The genome schema was drawn as Supplementary Fig. S2. It is 3,032,608 bp in length with a G + C content of 61.7%. Based on 16s rRNA gene phylogenetic analysis, *K. vulgaris* SPU B805 is highly homologous to a reported 2-KGA-producing strain of _K. vulgaris_ SKV\(^\text{3}\) (see Supplementary Fig. S3). The comparison of *K. vulgaris* SPU B805 and _K. vulgaris_ Y25\(^\text{5}\), _K. vulgaris_ WSH-001\(^\text{1}\), _K. vulgaris_ SKV\(^\text{3}\) and _K. vulgaris_ Hbe602\(^\text{2}\) in genome characteristics was summarized in Table 2. _K. vulgaris_ SPU B805 has the largest genome among all the strains, but it doesn't encode the most genes and proteins. No plasmid is found in _K. vulgaris_ SPU B805, while other four strains have 1–2 plasmids. The number of rRNA (15) and tRNA (58) are the same as those in other strains except for tRNA (59) in WSH-001. The number of sorbose dehydrogenase (sdh) and sorbosone dehydrogenase (sndh), key genes responsible for the bioconversion of L-sorbose to 2-KGA, is different among all the strains, shown in Table 2. There are 5 _sdh_ genes (KVC_2764, KVC_2744, KVC_0718, KVC_1927, KVC_0337) and 1 _sndh_ gene (KVC_0605) in strain SPU B805.
Furthermore, the genome-scale sequence comparison conducted by LAST software showed that the genome of *K. vulgare* SPU B805 is more similar with *K. vulgare* SKV (see Supplementary Fig. S4). Strain SPU B805, showing a 99% identity to strain SKV, harbors one circular chromosome without any plasmid. However, strain SKV contains one circle chromosome and one circle plasmid, while the other strains (Y25, WSH-001, Hbe602) consist of one circle chromosome and two circle plasmids, respectively. Therefore, it is the first time to find a 2-KGA-producing strain without any plasmid. The frame diagram of genome comparison between *K. vulgare* SPU B805, SKV, Hbe602, WSH-001 and Y25 was shown in Fig. 1. Most of the genome sequences are highly consistent except for some insertion, dislocation and rearrangement. Further analysis revealed that the DNA fragment of plasmid 1 (for example, namely as plasmid pKvSKV1 in SKV) was completely inserted into the circular chromosome (from 125,709 bp to 393,697 bp) of SPU B805, while the DNA fragment of plasmid 2 (belonging to strain Hbe602, WSH-001 and Y25), was completely lost in SPU B805.

**Genome annotation of *K. vulgare* SPU B805.** To facilitate the gene function analysis, the genes on chromosome and plasmid 1 of the four published *K. vulgare* strains (SKV, Hbe602, WSH-001 and Y25) were selected to compare the distribution of COG classification with SPU B805 (see Supplementary Fig. S5). Consequently, in SPU B805, the number of genes related to amino acid transport and metabolism (E) is similar to that of Hbe602 and Y25 and higher than that in SKV and WSH-001. These functional genes encode many transporters, which absorb nutrients from the environment to compensate for its metabolic defects. Besides, the number of genes related to inorganic ion transport and metabolism (P) and translation, ribosomal structure and biogenesis (J) in SPU B805 is more than those of the reported *K. vulgare* strains. Inorganic ions are a kind of important cofactor involved in various physiological reactions in cell. The strong translation ability may be helpful to biosynthesize protein in *K. vulgare* SPU B805.

On the basis of the COG and RAST annotation, the genes on plasmid 2 of strain Hbe602, WSH-001 and Y25 account for about 7% of the genome. A majority of them are related to amino acid transport and metabolism (E), and assigned as "an ABC transporter" (Supplementary Fig. S5 and Table S1). Besides, some of the genes for carbohydrate metabolism are annotated as gluconate and ketogluconate metabolism. For example, the enzyme

**Table 1.** The fermentation parameters of different *K. vulgare* strains. Note: *"*** indicate statistically significant difference when compared with *K. vulgare* SPU B805 (*p* < 0.5, 0.001).

| Strain     | *K. vulgare* SPU B805 | *K. vulgare* SKV | *K. vulgare* Hbe602 | *K. vulgare* WSH-001 | *K. vulgare* Y25 |
|------------|-----------------------|------------------|---------------------|----------------------|------------------|
| Cell growth (CFU/mL) | 52.1 ± 1.4 × 10^7 | Null             | Null                | Null                 | Null             |
| Cell growth (OD660)  | 5.63 ± 0.17          | Null             | Null                | 5.13 ± 0.12*        |                  |
| Volume (L)         | 5                     | 5                | 7                   |
| Total L-sorbose (g/L) | 80                   | 80               | 70                  |
| Fermentation period (h) | 44                  | 120***           | 70***               |
| 2-KGA production (g/L) | 82.74 ± 0.77         | 65.9 ± 0.4***    | 66.83 ± 1.1***      |
| 2-KGA productivity (g/L/h) | 1.88 ± 0.02          | 0.55 ± 0.00***   | 0.93 ± 0.01***      |
| Conversion rate (mol/mol) | 95.97%              | 76.44%***        | 88.47%***          |
| References        | This study           |                  |                     |

**Table 2.** Genome features of different 2-KGA-producing strains of *K. vulgare*.

| Strain     | *K. vulgare* SPU B805 | *K. vulgare* SKV | *K. vulgare* Hbe602 | *K. vulgare* WSH-001 | *K. vulgare* Y25 |
|------------|-----------------------|------------------|---------------------|----------------------|------------------|
| chromosome (Mb)      | 3.03                  | 2.76             | 2.77                | 2.77                 | 2.78             |
| G + C (%)          | 61.7                  | 61.7             | 61.7                | 61.7                 | 61.7             |
| Plasmid 1 (bp)     | Null                  | 267,949          | 267,988             | 267,986              | 268,675          |
| Plasmid 2 (bp)     | Null                  | Null             | 242,716             | 242,715              | 243,645          |
| Protein            | 2947                  | 2851             | 3178                | 3054                 | 2905             |
| Gene               | 3062                  | 3067             | 3281                | 3198                 | 3286             |
| rRNA               | 15                    | 15               | 15                  | 15                   | 15               |
| tRNA               | 58                    | 58               | 58                  | 59                   | 58               |
| SDH                | 5                     | 6                | 6                   | 5                    | 5                |
| SNDH               | 1                     | 1                | 2                   | 2                    | 2                |
| GA2DH              | Null                  | Null             | 1                   | 1                    | 1                |
| Reference         | This work             |                  |                     |                      |                  |
| Accession number   | CP017622              | CP016592         | CP012908            | CP002018             | CP002224         |
|                   |                      |                  | CP012909            | CP002019             | CP002225         |
|                   |                      |                  | CP016593            | CP012910             | CP002020         |
|                   |                      |                  |                     |                      | CP002226         |
gluconate 2-dehydrogenase (GA2DH) can convert 2-KGA into idonate, followed flowing to 6-phosphogluconate (6PG) and subsequently entering the PPP for energy and biomass production6, finally decrease the carbon flow to 2-KGA and lower the 2-KGA production. Additionally, one of the L-sorbosone dehydrogenase (SNDH) genes is also found to be located on plasmid 2. The over-expression of SNDH (encoded by plasmid 2) in \( K. \) \( vulgare \) \( Hbe602 \) produced an obvious byproduct (not identified its chemical structure in the published paper), so as to decrease the 2-KGA yield15. Moreover, the key genes unique to ED pathway, 2-keto-3-deoxy-6-phosphogluconate aldolase (Eda) and phosphogluconate dehydratase (Edd), are annotated to be located on plasmid 2. The deletion of edd and eda genes from \( Gluconobacter \) \( oxydans \) 621H resulted in a lower overall sugar uptake by cells in the cytoplasm, and led to more sugar being left in the periplasm and converted to the end-product (2-keto-gluconate) with high yield16. The loss of ED pathway maybe has a similar effect on 2-KGA production in \( K. \) \( vulgare \). Fortunately, these genes (GA2DH, SNDH, Eda and Edd) located on the indigenous plasmid 2, are all absent from the genome of \( K. \) \( vulgare \) SPU B805, which may be a reason for its high 2-KGA production. Therefore, presumably, elimination of the above genes or indigenous plasmid 2 is an effective method to enhance 2-KGA production by \( K. \) \( vulgare \).

Moreover, a total of 2933 coding sequences were obtained by RAST annotation in strain SPU B805, which belonged to 26 subsystems, including cofactor, membrane transport metabolism, nucleotides metabolism, protein metabolism, regulation and cell signaling, amino acids and carbohydrate metabolism, and so on. Among them, genes regarding to amino acid metabolism were accounted for the highest proportion (15%) (Fig. 2 and Supplementary Table S2). However, the genes coding histidinol-phosphate (EC 3.1.3.15), leucine-alanine transaminase (EC 2.6.1.12), alanine transaminase (EC 2.6.1.2) and asparagine synthetase (EC 6.3.1.1) were found to be absent in strain SPU B805, which resulted in the incomplete biosynthesis pathway of histidine, alanine and asparagine (Fig. 3). While in strain WSH-001, the biosynthesis pathways of more amino acids, such as histidine, glycine, lysine, proline, threonine, methionine, leucine and isoleucine, were deficient due to absence of one or more key enzymes7. So the biosynthetic pathways of amino acid in \( K. \) \( vulgare \) SPU B805 are more complete than

Figure 1. The genome comparison of \( K. \) \( vulgare \) SPU B805 with the published genomes. The frames under the main body represent the inverted fragments. The sequence of plasmid pKvSKV1 (belong to SKV), plasmid 1 (belong to Hbe602), pKVU_100 (belong to WSH-001) and pYP1 (belong to Y25) with highly similarity is inserted into the genome of \( K. \) \( vulgare \) SPU B805 (dark grey parts), while plasmid 2 (belong to SKV) or pKVU_200 (belong to WSH-001) or pYP12 (belong to Y25) (white fragments) is missing in the \( K. \) \( vulgare \) SPU B805 genome.
those in the reported \textit{K. vulgare} strain WSH-001. Previous study showed that glycine, serine, threonine and proline were important factors for 2-KGA biosynthesis, and threonine affected cell growth significantly\cite{8}. Glycine and serine are donors of one-carbon units in one-carbon metabolism, and threonine can be converted into glycine and acetyl-CoA. The impairment of these amino acids biosynthesis pathways will weaken the biosynthesis of nucleic acid and protein, and the biogenesis methyl group. Moreover, proline can decrease the destruction of microbial cells resulted by the osmotic stress at high 2-KGA concentrations\cite{18}. Therefore, presumably, the more complete biosynthesis pathways in strain SPU B805 may contribute to the growth of \textit{K. vulgare} and the accumulation of 2-KGA. Besides, the membrane transport genes were abundant in strain SPU B805, which might help to transport and absorb the nutrients and metabolites released by \textit{B. megatherium}\cite{1}.

\textbf{Construction of carbohydrate metabolic network.} Generally, the prokaryote carbohydrate metabolic pathways are diverse, including the EMP pathway, ED pathway and PPP pathway. However, the genomic annotation revealed that the genes coding 6-phosphofructokinase (EC 2.7.1.11, Pfk), 2-keto-3-deoxy-6-phosphogluconate aldolase (EC 4.1.2.14, Eda) and phosphogluconate dehydratase (EC 4.2.1.12, Edd) were missing in strain SPU B805. Because Pfk is an enzyme specific to the EMP pathway, the absence of \textit{pfk} leads to the incompleteness of the EMP route even though all the other enzymes are present. Similarly, Eda and Edd are the key enzymes unique to the ED pathway, therefore the absence of \textit{eda} and \textit{edd} imply the ED pathway is disabled in strain SPU B805. Thus, the PPP pathway was speculated as the main carbohydrate metabolic pathway in strain SPU B805.

The EMP pathways of strain WSH-001, Y25 and Hbe602 were also analyzed based on their published genome, and found to be incomplete because of the absence of gene \textit{pfk} (Table 3), therefore, we speculated that the ED and PPP pathways were the main carbohydrate metabolic routes. However, for strain SKV, the gene \textit{pfk}, \textit{eda} and \textit{edd} were all absent from the genome, so presumably the carbohydrate metabolism was mainly through the PPP pathway. To sum up, there are two types of carbohydrate metabolic pathways in \textit{K. vulgare}. One is simultaneously through the ED and PPP pathways in strain WSH-001, Y25 and Hbe602, the other one is only through the PPP pathway in strain SKV and SPU B805. The loss of the ED pathway had no negative effective on 2-KGA production (Table 1) further confirmed that the ED pathway might be negligible in the cell growth and 2-KGA accumulation, thus the PPP pathway played an important role in the cytoplasmic carbohydrate metabolism in \textit{K. vulgare}. 

\begin{figure}
\centering
\includegraphics[width=\textwidth]{image.png}
\caption{Gene distributions of metabolic subsystems of \textit{K. vulgare} SPU B805.}
\end{figure}
Carbohydrate metabolism analysis based on transcriptome. The abundance of FPKM (fragments per kilo bases of million fragments) was employed to represent the gene transcriptional level. As for strain SPU B805, the whole FPKM was divided into 11 grades (from Rank $^{-4}$ to Rank $^{+6}$), and the median point was 195.564 (Fig. 4). The FPKM of the key genes related to PPP pathway and TCA cycle (Zwf, EC 1.1.1.49; Gnd, EC 1.1.1.44; Rpe, EC 5.1.3.1; RpiA, EC 5.3.1.6; TktA, EC 2.2.1.1; Tal, EC 2.2.1.2; GltA, EC 2.3.3.1; AceE, EC 1.2.4.1; Acn, EC 4.2.1.3; Icd, EC 1.1.1.42; SucA, EC 1.2.4.2; SucDC, EC 6.2.1.5; SdhA, EC 1.3.5.4; FumC, EC 4.2.1.2; Mdh, EC 1.1.1.37) were marked out in Fig. 4, and it was noteworthy that all of the genes related to carbohydrate metabolism (PPP, TCA) were transcribed at higher levels than the median point. However, no transcription signals of pfk, ga2dh, edd and eda could be detected, which confirmed that these genes did not exist in the genome of strain SPU B805, and indicated that the EMP and ED pathway were disabled. Because the PPP pathway, ED pathway and EMP pathway were three parallel carbon metabolic routes before TCA cycle, therefore, the only existing complete PPP pathway was assigned as the major and important cytoplasmic carbohydrate decomposition pathway linked with TCA cycle in strain SPU B805.

In fermentation, the FPKM of the five sdh (KVC_2764, KVC_2744, KVC_0718, KVC_1927, KVC_0337) (the average FPKM value 24691.32, 10501.89, 2078.73, 750.78, 220.99) were obviously higher than those of the central carbon degradation related genes, which partially explained that the majority of L-sorbose was converted to 2-KGA in strain SPU B805. Although the gene sndh (KVC_0605, the average FPKM value 479.86) had a relative lower transcriptional level, it did not affect the 2-KGA production rate because SDH was a dual-function enzyme, which could oxidize L-sorbose to L-sorbosone and further oxidize to 2-KGA. Therefore, the central carbon metabolism, as well as the transcription level (the average FPKM value), were put together in Fig. 5. Strain SPU B805 possesses two separated modes for L-sorbose catabolism. A major part of L-sorbose is oxidized to L-sorbosone in the periplasm by the membrane-bound SDH, and further to 2-KGA. A minor part of L-sorbose as carbon source is catabolized to F6P and subsequently to 6PG in cytoplasm, and then enters the PPP pathway for energy and biomass production. In the oxidative branch, 6PG is converted into ribulose 5-phosphate (Ru5P), carbon dioxide and NADPH. NADPH is the major reducing power used to maintain the redox balance under stress situation. The formed Ru5P is converted to R5P by ribose 5-phosphate isomerase (RpiA, EC 5.3.1.6) or X5P by ribulose 5-phosphate epimerase (Rpe, EC 5.1.3.1) in the non-oxidative branch. R5P and X5P undergo carbon rearrangement to generate sedoheptulose 7-phosphate (S7P), glyceraldehyde
3-phosphate (G3P) and erythrose 4-phosphate (E4P). R5P and E4P, as the precursors, were used for the biosynthesis of RNA, DNA, as well as aromatic amino acids. 

**RT-qPCR validation of the genes related to carbohydrate metabolism.** The key genes in PPP pathway, zwf (KVC_1674), gnd (KVC_0890), transketolase (KVC_2247; tktA) and transaldolase B (KVC_1789; tal) were conducted by RT-qPCR. Compared with internal standard (polA), all of the four genes showed a relative high transcription level, and the changed trends in different time-points were in accordance with the transcriptome results (see Supplementary Fig. S6). The gene transcription level assayed by RNA-seq and RT-qPCR showed high correlation, indicating the reliability of the RNA-seq analysis.

**Discussion**

For a long time, researchers have mainly concentrated on elucidating the symbiosis mechanism of *K. vulgaris* and *B. megaterium*, however, few studies have been explored on the underlying mechanism for the differences in central carbon metabolic pathways and 2-KGA yield of *K. vulgaris*. In this study, the genome of *K. vulgaris* SPU B805 was completely sequenced and the metabolic network was reconstructed, which provided a new insight into the carbohydrate metabolic specificity. *K. vulgaris* SPU B805 was the first 2-KGA-producing strain without any plasmid. Its genome was similar to *K. vulgaris* WSH-001 except for some insertion, dislocation and rearrangement. The genome annotation showed that except for histidine, alanine and asparagine, the biosynthetic pathway of other amino acids was complete in strain SPU B805, which had significantly integral function than that in WSH-001 (eight amino acid biosynthesis pathways incomplete)6. Liu, et al.7 reported that glycine, proline, threonine and...
isoleucine play vital roles in *K. vulgare* growth and 2-KGA production, and the addition of these amino acids increased the 2-KGA productivity by 20.4%, 17.2%, 17.2% and 11.8%, respectively. Zhang, et al.\(^8\) also determined the key factors affected 2-KGA fermentation by orthogonal design experiments, and the results indicated that glycine and threonine were the key components affecting the mixed cell growth; serine, glycine, and proline were the key components that affected the 2-KGA production. Therefore, the more complete amino acid biosynthetic pathway, the more contribution to the high 2-KGA production of *K. vulgare* SPU B805. Moreover, the absence of the SNDH on plasmid 2 can avoid the unknown byproduct in the formation of 2-KGA. Besides, the gene ga2dh,

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**Figure 4.** Transcriptome information of *K. vulgare* SPU B805. X axial represents the FPKM of *K. vulgare* SPU B805, Y axial represents the counts of expressed genes. The blue and pink panels indicate the FPKM is below and above the median point, respectively. The FPKM between 150 and 300, 300 and 600, 600 and 1200, 1200 and 2400, 2400 and 4800, 4800 and 9600 was defined as Rank 0, +1, +2, +3, +4, +5. The FPKM between 75 and 150, 36 and 75, 18 and 36 was defined as Rank −1, −2, and −3. The FPKM above 9600 and below 18 was defined as Rank +6 and Rank −4.

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**Figure 5.** Overview of the average FPKM of carbohydrate metabolism related genes in *K. vulgare* SPU B805. Blue lines and orange lines indicate the average FPKM is lower and higher than the median point, respectively. Dotted lines indicate no transcription signal is detected. The thickness of lines represents the FPKM level. The figures above or below lines represent the EC number of the enzyme and the average FPKM.
The genome and transcriptome analysis showed that the central carbon metabolism of *K. vulgare* was versatile. Strain WSH-001, Y25 and Hbe602 contained all the genes involved in the ED, the PPP and the TCA cycle, meaning that these pathways were complete. While the *pfk* gene was absent from strain WSH-001, Y25 and Hbe602, indicating that the EMP pathway was non-functional in these strains. Whereas, in strain SPU B805 and SKV, all the genes referred to the PPP and the TCA cycle were found, meaning that the PPP and the TCA pathway were complete. The *pfk*, as well as *eda* and *edd* genes were simultaneously missing from the genome of strain SPU B805 and SKV, which suggested that neither of the EMP and the ED pathway were functional. Therefore, the PPP pathway is the exclusively pathway for cytoplasmic carbohydrate decomposition in strain SPU B805 and SKV. In strain WSH-001, Y25 and Hbe602, the ED and the PPP pathways are paralleled for cytoplasmic carbohydrate catabolism. Although the ED pathway was complete in strain WSH-001, Y25 and Hbe602 genome, the carbon flux through this pathway is presumably negligible because of the lower ATP productivity ability of the ED pathway. Moreover, in *G. oxydans* 621H, the deletion of *gnd* to inactive the PPP pathway resulted in a reduced final biomass and a decreased end-product 2-keto-glucuronate, whereas, the inactivation of the ED pathway by deleting the *edd-eda* genes caused a lower overall sugar uptake by cells for the formation of biomass and energy in the cytoplasm, and led to a larger fraction of the sugar being converted to the final product of 2-keto-glucuronate in the periplasm. This research indicated that the PPP pathway was of the major importance for cytoplasmic carbohy-
dride catabolism, whereas the ED pathway pathway is the exclusively pathway for cytoplasmic carbohydrate catabolism, whereas the ED pathway is dispensable. The loss of ED pathway maybe has a similar effect on cell growth, the bacterium usually does not synthesize additional protein only for one more energy.

The PPP pathway, as a fundamental component in cellular metabolism, plays an important role in maintain carbon homoeostasis. The PPP pathway can provide various intermediate metabolites for the biosynthesis of nucleotide and amino acids, such as G3P, E4P, R5P, and S7P, and generate NADPH for the maintenance of redox balance under the stress situations. *K. vulgare* as an aerobic bacterium, its growth and metabolism are inevitably subject to reactive oxygen species (ROS), therefore a reducing environment is an essential requirement for cell metabolism. It is well known that NADPH is the major reducing power in cell, and plays an important role in the GSH and thioredoxin metabolism. Besides, a large number of anti-oxidant enzymes are coupled with NADPH as cofactor, thus the PPP pathway may help *K. vulgare* SPU B805 to defeat intracellular ROS. Moreover, the biomass of *K. vulgare* was composed of protein, DNA, RNA, lipids, peptidoglycans, lipopolysaccharides, glycogen and soluble pool. It is conceivable that the PPP pathway may provide a variety of intermediates as precursors to satisfy the demands for the biosynthesis of cellular components and facilitate to curb different types of environmental hardship in the process of growth.

### Materials and Methods

#### Bacterial strains.

Two strains, *K. vulgare* SPU B805 (2-KGA-producing strain) and *B. megatherium* SPU B806 (the helper strain), were used in Microbial Resource Center of Shenyang Pharmaceutical University, were used for 2-KGA fermentation in this study.

#### Culture medium and fermentation.

The plate medium (P medium) contained 20 g/L L-sorbose, 1 g/L KH₂PO₄, 3 g/L yeast powder, 0.2 g/L MgSO₄, 3 g/L beef extract, 1 g/L urea, 10 g/L peptone, 3 g/L corn steep liquor and 20 g/L agar, pH 6.8. The plates were cultured at 28–30°C for 4 days. The seed medium (S medium) contained 20 g/L L-sorbose, 1 g/L KH₂PO₄, 3 g/L yeast powder, 0.2 g/L MgSO₄, 12 g/L urea, 10 g/L corn steep liquor, pH 6.8. Seeds were cultured at 28–30°C, 250 rpm for 18 h. The fermentation medium (F medium) contained 80 g/L L-sorbose, 20 g/L corn steep liquor, 1 g/L KH₂PO₄, 0.2 g/L MgSO₄, 12 g/L urea. The fermentation was conducted in 5-L fermenter containing 3 L F medium and 300 mL co-inoculated seeds of *K. vulgare* SPU B805 and *B. megatherium* SPU B806. The fermentation pH was maintained automatically at 6.8–7.0 by addition of 30% (m/v) Na₂CO₃ solution. The temperature was controlled at 30°C, and the agitation speed was 400 rpm with the air flow rate of 1.5L/min.

To calculate the Colony Forming Units (CFUs) of *K. vulgare*, the fermentation broth was diluted and spread on P plate, and the growth curve was drawn according to the CFU/mL. Meanwhile, the cell density was measured spectrophotometrically at 660 nm.

#### Genome sequencing and assembly.

*K. vulgare* SPU B805 was cultured in 250 mL flasks with 50 mL S medium at 30°C for 18 h at 220 rpm. The genome of strain SPU B805 was extracted with the TIANamp Genomic DNA Kit (Tiangen Biotech (Beijing) Co., Ltd.) according to the instruction. The concentration and purity of DNA sample were qualified by Agilent 2100 BioAnalyzer (Agilent Technologies, USA). The integrity of genomic DNA was detected by agarose gel electrophoresis. The DNA samples with a 260/280 nm absorbance ratio of 1.8–2.0 and a 260/230 nm absorbance ratio of 2.0–2.2 were considered pure and then used for the library construction and sequencing.

The complete genome sequence was determined by the Chinese National Human Genome Center (Shanghai, China) using 454 single-end sequencing technology. It yielded a total of 116,450 reads with a 17.29-fold coverage of the genome. The reads were assembled into 26 large contigs (>500 nucleotides) and 3 small contigs (100–500 nucleotides) by using the 454 Newbler assembler (454 Life Sciences, Branford, CT). The relationships between the contigs were determined by ContigScape, and the gaps between the contigs were closed by PCR amplification,
followed by DNA sequencing. Finally, the contigs were assembled into a circular chromosome, and no plasmid was found. The complete genome of SPU B805 had an error rate of less than 0.2 in a 10-kb sequence as determined by sequence assembly and quality assessment using the Phred/Phrap/Consed software package.

**Genome annotation.** Protein-coding genes were predicted by combining the results of Glimmer 3.02 and ZCURVE, followed by manual inspection. Each gene was functionally classified by assigning a Cluster of Orthologous groups (COG) number. Both tRNA and rRNA genes were identified by tRNA-scan-SE and tRNAscan2, respectively. Functional annotations of ORFs were performed by searching against public databases, including NCBI (http://www.ncbi.nlm.nih.gov/), KEGG and SWISS-Prot. Basic metabolic pathways were reconstructed by KEGG and MetaCyc. The Rapid Annotation using Subsystem Technology (RAST) server was used to re-annotate the genome of *K. vulgaris* SPU B805. Then, RAST annotation was put through the Model SEED pipeline to obtain the initial network reconstruction.

**Transcriptome sequencing.** *K. vulgaris* SPU B805 and *B. megaterium* SPU B806 were co-cultured in 5-L fermenter. The fermentation processes of 0 h (seed), 8 h, 16 h, 24 h, 32 h and 40 h were chosen to represent different growth phase. The co-culture samples were subjected to centrifugation of 4,000 rpm for 5 min at 4 °C to eliminate the cells of *B. megaterium* SPU B806. The RNA of *K. vulgaris* SPU B805 was extracted with the RNAPrep Pure Bacteria Kit (Tiangen, Beijing, China). The concentration and purity of RNA sample were qualified by a Agilent 2100 BioAnalyzer (Agilent Technologies, USA). The integrity was monitored on 1% agarose gels. The RNA samples with a 260/280 nm absorbance ration of 1.8–2.1 were considered pure, and a RNA ration (28S/18S) ≥ 1.5 and RN ≥ 7 were considered integrity: The NEBNext Ultra RNA Library Prep Kit (NEB, USA) for Illumina was used to construct sequencing libraries. The RNA was removed from total RNA using a Ribo-Zero Magnedia Kit (Epipcentre, USA) to enrich the mRNA and then the cDNA was synthesized. The library preparation was sequenced on Illumina HiSeq 2500 platform. The genome of *K. vulgaris* SPU B805 was used as a reference.

**Analyses of 2-KGA and L-sorbose.** The concentration of 2-KGA and L-sorbose in the fermentation broth were analyzed by the High-Performance Liquid Chromatography (HPLC) (Waters Corp., Massachusetts, USA) using an amino column (Agilent Zorbax NH2) with an ultraviolet detector at 210 nm. Cetonitrile-KH2PO4 (5%/95%, v/v) was used as mobile phase with a flow rate of 0.6 mL/min.

**Real time quantitative PCR.** The primers used for real time quantitative PCR (RT-qPCR) were listed in Supplementary Table S3. RT-qPCR was conducted with the GoTaq qPCR Master Mix (Promega (Beijing) Biotech Co., Ltd.) on a Stratagene Mx3000P (Agilent Technologies Inc, USA). The DNA polymerase I (KVC_2082, polA) gene was used as an internal standard. The fold change of gene expression was calculated by the 2−ΔΔCT method.

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Author Contributions
C.Y.W, Y.L. designed the project and experiments; C.Y.W and Z.W.G. performed the experiments; Y.L. and L.C.L. annotated and uploaded the genome; Y.C.W assisted in the analysis of final data. M.Y.Z and T.Y.Z. assisted in the revision the manuscript; Y.X.Z. revised the manuscript. All the authors read and approved the final version of the manuscript.

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