Evaluation of Carbon and Electron Flow in *Lactobacillus brevis* as a Potential Host for Heterologous 1-Butanol Biosynthesis

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ABSTRACT

Heterofermentative lactic acid bacterium *Lactobacillus brevis* may be considered as a promising host for heterologous butanol synthesis because of tolerance to butanol and ability to ferment pentose and hexose sugars from wood hydrolysates that are cheap and renewable carbohydrate sources. Carbon and electron flow was evaluated in two *L. brevis* strains in order to assess metabolic potential of these bacteria for heterologous butanol synthesis. Conditions required for generation of acetyl-CoA and NADH which are necessary for butanol biosynthesis have been determined. Key enzymes controlling direction of metabolic fluxes in *L. brevis* in various redox conditions were defined. In anaerobic glucose fermentation, the carbon flow through acetyl-CoA is regulated by aldehyde dehydrogenase ALDH possessing low affinity to NADH and activity (\(K_m^{\text{NADH}} = 200 \mu M, V_{\text{max}} = 0.03 \text{ U/mg of total cell protein}\)). Aerobically, the NADH-oxidase NOX (\(K_m^{\text{NADH}} = 25 \mu M, V_{\text{max}} = 1.7 \text{ U/mg}\)) efficiently competes with ALDH for NADH that results in formation of acetate instead of acetyl-CoA. In general, external electron acceptors (oxygen, fructose) and pentoses decrease NADH availability for native ethanol and recombinant butanol enzymes and therefore reduce carbon flux through acetyl-CoA. Pyruvate metabolism was studied in order to reveal redirection possibilities of competitive carbon fluxes towards butanol synthesis. The study provides a basis for the rational development of *L. brevis* strains producing butanol from wood hydrolysate.

Keywords: *Lactobacillus brevis*; Metabolism; Carbon and Electron Flow; Heterologous Butanol Synthesis; SO₂-Ethanol-Water Hydrolysate

1. Introduction

A number of studies have been carried out on the production of liquid fuels from renewable plant sources by microbial synthesis. A special attention is paid to alcohols, mainly ethanol and 1-butanol as an effective substitute for gasoline and diesel in combustion engines [1,2]. 1-butanol has an advantage over ethanol due to its higher energy content, lower volatility, less ignition problems, better miscibility with gasoline and the possibility to use it without modification of engines and infrastructure for supplying and distribution [3]. Anaerobic bacteria of the genus *Clostridium* are the natural producers of 1-butanol [4]. However, the application of existing *Clostridium* strains for large scale industry is not feasible in the current economic conditions because of low 1-butanol titer (<15 g/l), two-phase metabolism (formation of acids precedes the formation of solvents) hampering organization of a continuous fermentation process, synthesis of a significant amount of by-products (acetone, ethanol, acetate, butyrate) and expensive fermentation substrates.

Several attempts have been made to construct 1-butanol-producing strains based on other microbial platforms such as *Escherichia coli*, *Saccharomyces cerevisiae*, *Pseudomonas putida*, *Lactobacillus brevis*, *Bacillus subtilis* [5-10]. A high-titer (30 g/liter) and high-yield (70% to 88% of the theoretical) production of 1-butanol was achieved in *E. coli* [11]. This is comparable to or exceeding the levels demonstrated by native producers.
Economically viable production of biofuels should be based on an inexpensive, renewable raw material like plant biomass. Recently developed SEW (SO₂-ethanol-water) pulping is a promising fractionation process for lignocellulosic biomass. Its advantages over conventional pulping methods include simplified chemical recovery, lower capital costs and rapid impregnation of the feedstocks [12,13]. SEW-hydrolyzate from spruce chips contains mannose, glucose, galactose, xylose, and arabinose which can be used for microbial fermentation [14]. However, hydrolysis by-products such as formic acid, acetic acid, levulinic acid, furfural and hydroxymethyl furfural inhibit fermentation of many industrial microorganisms, including E. coli [15,16].

Heterofermentative lactic acid bacterium Lactobacillus brevis is able to ferment pentose as well as hexose sugars from various plant sources [17] including acid hydrolyzate of hemicellulose [18]. In addition L. brevis is tolerant to 3% butanol and may be easily adapted to increased butanol concentration [9,19]. Thus it can be considered as a potential platform for heterologous butanol synthesis from cheap renewable plant sources. The first objective of the present work is to investigate the ability of wild-type L. brevis strains to ferment sugars of SEW-hydrolyzate from spruce chips that are abundant waste of woodworking and pulp and paper industry.

In the previous study we constructed a L. brevis butanol-producing strain by cloning thl, hbd, crt, bcd, etfB, and etfA genes from Clostridium acetobutylicum [9]. However, the butanol titer in recombinant strain did not exceed ~300 mg·L⁻¹. Expression of the butanol pathway genes did not change the level and the ratio of native L. brevis end-products: the spectra of metabolites other than butanol were similar in recombinant and wild-type L. brevis genes did not change the level and the ratio of native L. brevis strains to ferment sugars of SEW-hydrolyzate from spruce chips that are abundant waste of woodworking and pulp and paper industry.

In the previous study we constructed a L. brevis butanol-producing strain by cloning thl, hbd, crt, bcd, etfB, and etfA genes from Clostridium acetobutylicum [9]. However, the butanol titer in recombinant strain did not exceed ~300 mg·L⁻¹. Expression of the butanol pathway genes did not change the level and the ratio of native L. brevis end-products: the spectra of metabolites other than butanol were similar in recombinant and wild-type L. brevis strains that indicated the inability of the recombinant pathway to compete with native ones. The present study is focused on evaluation of carbon and electron flow in wild-type L. brevis strains in order to assess metabolic potentialities of these bacteria for heterologous butanol synthesis and reveal bottlenecks preventing efficient butanol synthesis by recombinant L. brevis strains.

2. Materials and Methods

2.1. Bacterial Strains and Growth Conditions

Lactobacillus brevis ATCC 367, Lactobacillus brevis ATCC 8287, and Clostridium acetobutylicum ATCC 824 were used for metabolic studies.

C. acetobutylicum was cultivated anaerobically at 37°C in MSS medium as described earlier [20]. L. brevis strains were grown at 30°C aerobically (with agitation at 250 rpm) or semi-aerobically (without agitation in closed vials) in MRS or HM media supplemented with kanamycin (12.5 µg·ml⁻¹). The non-autoclaved MRS medium was composed of (w/v) 1% casein peptone, 1% meat extract, 0.5% yeast extract, 2% glucose, 0.1% Tween 80, 0.2% K₂HPO₄, 0.5% Na-acetate, 0.2% ammonium citrate, 0.02% MgSO₄·7H₂O, and 0.005% MnSO₄·H₂O. In the autoclaved MRS medium glucose was partly converted to fructose. The HM medium had the same content as the MRS medium except the glucose was substituted with 25% SO₂-ethanol-water hydrolyzate of spruce chips. The SEW-hydrolyzate was kindly provided by E. Sklavounos, Aalto University. Sugar composition of the autoclaved MRS medium and the HM medium was determined by HPLC (Table 1).

2.2. Analytical Methods

2.2.1. Chromatograph

Metabolic end-products of L. brevis were identified and quantified by high-performance liquid chromatography (HPLC). Waters 2695 Separations module was equipped with Waters 2414 Refractive Index Detector, Bio-Rad Aminex HPX-87H column (300 mm × 7.8 mm × 9 µm), and Micro-Guard Cation H⁺-Cartridge (Bio-Rad, Hercules, CA, US). The column was heated at 65°C; the eluent (5 mM H₂SO₄) was circulated at a flow rate of 0.60 mL·min⁻¹. The sugars were determined by Waters 2690 Separations module with Waters 2414 Refractive Index Detector, equipped with a Bio-Rad HPX-87P column (300 mm × 7.8 mm × 9 µm) and two Micro-Guard Deashing Cartridges (Bio-Rad, Hercules, CA, US) at 70°C with a flow rate of 0.60 mL·min⁻¹ using de-ionized water as eluent. Cellobiose (Rotth, Karlsruhe, Germany) was added to the samples as an internal standard. Sugar and metabolite concentrations were calculated from at least three experiments, the accuracy was within ±5%.

Table 1. Sugar and uronic acid composition (mM) of MRS and HM media.

<table>
<thead>
<tr>
<th>Sugars</th>
<th>MRS</th>
<th>HM</th>
</tr>
</thead>
<tbody>
<tr>
<td>Glucose</td>
<td>88</td>
<td>22</td>
</tr>
<tr>
<td>Fructose</td>
<td>23</td>
<td>-</td>
</tr>
<tr>
<td>Xylose</td>
<td>-</td>
<td>36</td>
</tr>
<tr>
<td>Mannose</td>
<td>-</td>
<td>59</td>
</tr>
<tr>
<td>Arabinose</td>
<td>-</td>
<td>7</td>
</tr>
<tr>
<td>Galactose</td>
<td>-</td>
<td>12</td>
</tr>
<tr>
<td>Uronic acids: glucuronate, galacturonate and 4-O-Me-glucuronate</td>
<td>-</td>
<td>20</td>
</tr>
</tbody>
</table>
2.2.2. Enzyme Assays
Cell-free extracts were prepared as described previously [9]. The protein content was determined with bicinchoninic acid [21] or by Bradford [22] after removal of the cell debris. Bovine serum albumine (Sigma) was used as a standard.

Lactate dehydrogenase, NADH:H2O oxidase, alcohol dehydrogenase, aldehyde dehydrogenase, and 3-hydroxybutyryl-CoA dehydrogenase activities were measured by NADH oxidation at 340 nm (ε_{NADH} = 6.22) in accordance with the published procedures [9,23,24]. ADH activity was measured by NAD+ reduction as described by Berezina et al. [9]. All activities were assayed at 30°C and pH 6.7 that corresponded to intracellular conditions in logarithmic growth phase [25].

Enzyme activity of pyruvate dehydrogenase was assayed as described by Yahui et al. [26] from the extracts of aerobically or semi-anaerobically grown cells collected at logarithmic growth phase, at pH 7.1 (enzyme optimum) and 6.7 (intracellular pH). Pyruvate formate lyase activity was measured as described by Asanuma et al. [27], in an anaerobic chamber. The cell extracts for PFL assay were prepared anaerobically as described by Berezina et al. [9]. Pyruvate oxidase, and pyruvate decarboxylase activities were assayed as described previously [28,29]. One unit of activity was defined as 1 µmol of substrate utilized or product formed in a minute per mg of total cell protein (U/mg).

2.2.3. Measurement of Intracellular NAD⁺ and NADH Levels
A freshly grown cell culture was concentrated by centrifugation up to 30 mg·ml⁻¹ of total cell protein. For NAD⁺ extraction, 0.1 ml of perchloric acid was mixed with 0.4 ml of cell suspension, incubated for 5 min at 60°C and immediately chilled on ice. The extract was neutralized to pH 7.0 with 0.5 ml of 2 M KOH containing 1.2 M Tris-HCl (pH 9.0) and 1.36 M semicarbazide. The precipitate was removed by centrifugation, and the supernatant was used for NAD⁺ measurements. For NADH extraction, 0.1 ml of 2 M KOH was mixed with 0.3 ml of cell suspension, incubated for 5 min at 60°C and immediately chilled on ice. The extract was neutralized on ice to pH 8.0 with 0.4 ml of 1M HEPES, pH 5.0. The precipitate was removed by centrifugation and the supernatant was used for NADH measurements. NAD⁺ and NADH concentrations in the extract were determined by enzyme assay with commercial alcohol and lactate dehydrogenases (Sigma-Aldrich) according to Bergmeyer [30]. Absorbance at 340 nm was measured by a Cintra 404 spectrophotometer. Total intracellular [NAD⁺ + NADH] and [NADH] concentrations were calculated assuming that 1 mg of total cell protein binds 3.7 µl of cytosole [31].

3. Results
3.1. Effect of Aeration and Sugar Composition on Growth Characteristics of Two L. brevis Strains

Although L. brevis lacks cytochromes, porphyrins, and respiratory enzymes and is generally recognized as an anaerobic bacteria [32], the aeration positively influenced growth rate of both strains (Figure 1).

In aerobic incubation in MRS medium the ATCC 8287 had a higher growth rate than the ATCC 367, while in semi-anaerobic conditions the difference between the strains was very small (Figure 1(a)).

In HM medium the ATCC 8287 strain had a higher growth rate than the ATCC 367 strain both in aerobic and semi-anaerobic conditions (Figure 1(b)).

Both strains were able to ferment glucose, xylene, arabinose and galactose of SEW-hydrolysate, but not mannose (Figure 2). The 8287 strain fermented xylose better than the 367 strain that correlates with faster growth of the 8287 strain in semi-anaerobic conditions (Figure 1(b)).

In semi-anaerobic fermentation of MRS medium lactate, ethanol and mannitol were produced from glucose and fructose. The level of acetate at first went down, and

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Figure 1. Growth curves of L. brevis ATCC 367 and ATCC 8287 strains on MRS medium (a) and HM medium (b). (●): ATCC 367, aerobic growth; (○): ATCC 8287, aerobic growth; (✦): ATCC 367, semi-anaerobic growth; (◇): ATCC 8287, semi-anaerobic growth. The mean values calculated from at least three experiments; the accuracy is within ±5%.

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then came back to initial value (Figure 3(a)). In aerobic MRS fermentation by \textit{L. brevis} 8287, lactate and acetate were produced in equimolar amounts during exponential growth phase. In stationary growth phase the decrease in lactate concentration was equal to increase in acetate concentration. Ethanol was not produced aerobically and mannitol only in minute amounts (Figure 3(b)). Aerobic and semi-anaerobic fermentations patterns of \textit{L. brevis} 367 were similar to \textit{L. brevis} 8287.

The time-dependent profiles of sugar utilization and metabolite formation during semi-anaerobic and aerobic fermentation of HM medium containing hexoses, pentoses, and uronic acids (Table 1) by \textit{L. brevis} 8287 are shown in Figures 3(c) and (d). Lactate, ethanol and acetate were produced semi-anaerobically whereas only lactate and acetate were produced aerobically. In stationary growth phase of aerobic incubation, the concentration of lactate decreased while the acetate concentration increased correspondingly (Figure 3(d)). For \textit{L. brevis} ATCC 367 the pattern of metabolic end-products was similar. Both strains accumulated biomass in HM medium better than in MRS medium, and the largest difference was observed in semi-anaerobic ATCC 8287 culture (Figure 1).

Fermentation balance of the ATCC 8287 strain cultivated in semi-anaerobic and aerobic conditions on MRS and HM media is shown in Table 2.

### 3.2. NADH-Dependent Enzymes of \textit{L. brevis} Metabolism

Lactate dehydrogenase (LDH), alcohol dehydrogenase (ADH), aldehyde dehydrogenase (ALDH), and NADH oxidase (NOX) activities were measured in cell extracts of both \textit{L. brevis} strains. The \textit{L. brevis} ATCC 367 genome sequence was screened for the corresponding genes (Table 3).
Table 2. Fermentation balance of *L. bevis* ATCC 8287 cultivated in various redox conditions. The mean values calculated from at least three experiments. The accuracy of measurements is within ±5%.

<table>
<thead>
<tr>
<th>Fermentable carbohydrate, mM</th>
<th>MRS</th>
<th>HM</th>
</tr>
</thead>
<tbody>
<tr>
<td>Metabolites, mM</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Lactate</td>
<td>Acetate</td>
</tr>
<tr>
<td>Cultivation time, h</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Semi-anaerobic cultivation</td>
<td></td>
<td></td>
</tr>
<tr>
<td>30</td>
<td>87</td>
<td>-</td>
</tr>
<tr>
<td>48</td>
<td>93</td>
<td>3</td>
</tr>
<tr>
<td>Aerobic cultivation</td>
<td></td>
<td></td>
</tr>
<tr>
<td>24</td>
<td>106</td>
<td>103</td>
</tr>
<tr>
<td>48</td>
<td>68</td>
<td>133</td>
</tr>
</tbody>
</table>

Table 3. *L. brevis* genes presumably encoding enzymes involved in pyruvate and acetyl-CoA metabolism.

<table>
<thead>
<tr>
<th>Gene</th>
<th>Predicted enzyme*</th>
<th>EC number</th>
</tr>
</thead>
<tbody>
<tr>
<td>LVIS 0514</td>
<td>NAD⁺-dependent L-lactate dehydrogenase L-LDH</td>
<td>1.1.1.27</td>
</tr>
<tr>
<td>LVIS 1352</td>
<td>NAD⁺-dependent D-lactate dehydrogenase D-LDH</td>
<td>1.1.1.28</td>
</tr>
<tr>
<td>LVIS 0119</td>
<td>Bifunctional acetaldehyde-CoA/Alcohol dehydrogenase ALDH, ADH</td>
<td>1.2.1.10, 1.1.1.1</td>
</tr>
<tr>
<td>LVIS 0254, LVIS 1019</td>
<td>Zn-dependent alcohol dehydrogenases ADH</td>
<td>1.1.1.1</td>
</tr>
<tr>
<td>LVIS 1558</td>
<td>NADH-oxidase NOX</td>
<td>1.6.99.3</td>
</tr>
<tr>
<td>LVIS 0313</td>
<td>Pyruvate oxidase POX</td>
<td>1.2.3.3</td>
</tr>
<tr>
<td>LVIS 1407, LVIS 1408, LVIS 1409, LVIS 1410</td>
<td>Pyruvate dehydrogenase enzyme complex PDH: Dihydrolipoamide dehydrogenase,</td>
<td>1.8.1.4</td>
</tr>
<tr>
<td>LVIS 1408, LVIS 1409</td>
<td>Dihydrolipoamide succinyltransferase (the E2 component of PDH complex)</td>
<td>2.3.1.12</td>
</tr>
<tr>
<td>LVIS 1410</td>
<td>Beta and alpha subunits for E1 component of PDH complex, correspondingly</td>
<td>1.2.4.1</td>
</tr>
<tr>
<td>LVIS 0491</td>
<td>Acetolactate synthase ALS</td>
<td>2.2.1.6</td>
</tr>
<tr>
<td>LVIS 0492</td>
<td>Acetolactate decarboxylase ALDB</td>
<td>4.1.1.5</td>
</tr>
<tr>
<td>LVIS 0187</td>
<td>Acetoain reductase BUTB</td>
<td>1.1.1.4</td>
</tr>
<tr>
<td>LVIS 2218</td>
<td>Acetyl-CoA C-acetyltransferase (thiolase) THL</td>
<td>2.3.1.9</td>
</tr>
</tbody>
</table>

*By analysis of *L. brevis* ATCC 367 genome, GenBank (CP000416), [33,34].

Overall lactate dehydrogenase activity was higher than the other detected enzyme activities: $V_{\text{max}}$ 37.0 and 26.5 U/mg of total cell protein in the 367 and 8287 strains correspondingly. ALDH activity was 0.02 and 0.03 U/mg in 367 and 8287 strains correspondingly (Table 4). ADH activity following the ALDH in the ethanol-forming pathway was 10.8 U/mg in both strains. No reverse ADH activity was detected in cell extracts at pH 6.7 or pH 6.3. This indicates that the process of ethanol formation is irreversible in physiological conditions. $K_m^{\text{NADH}}$ values for LDH and ALDH were comparable and about 2 - 2.5 times higher than $K_m^{\text{NADH}}$ of ADH (Table 4). NOX has higher activity compared to ALDH and the highest affinity to NADH among all the tested enzymes: $V_{\text{max}}$ (NOX) = 1.7 U/mg, $K_m$ (NOX) = 25 µM versus $V_{\text{max}}$ (ALDH) = 0.03 U/mg, $K_m$ (ALDH) = 200 µM for ATCC 8287 strain (Table 4).

In the recombinant butanol-producing *L. brevis* strains [7], the 3-hydroxybutyryl-CoA dehydrogenase (HBD, EC 1.1.1.35) is the first NADH-dependent enzyme of the heterologous butanol pathway originated from acetyl-CoA. The $K_m^{\text{NADH}}$ of HBD is 178 µM, which is slightly lower than $K_m^{\text{NADH}}$ of ALDH but sufficiently higher than $K_m^{\text{NADH}}$ of NOX (Table 4).

Intracellular NAD⁺/NADH ratio and total [NAD⁺ + NADH] concentration as a function of cell growth were determined for batch cultures (Figure 4).

3.3. Pyruvate-Converting Pathways in *L. brevis*

In *L. brevis*, pyruvate is converted to D- and L-lactate by corresponding lactate-dehydrogenases [9] (Table 3). But under certain conditions LAB may use alternative ways of utilizing pyruvate than reduction to lactic acid.
Table 4. $K_{m}^{NADH}$ (µM) and $V_{max}$ (U·mg$^{-1}$ of total cell protein) for NADH-dependent enzymes of *L. brevis* ATCC367, *L. brevis* ATCC8287 and *C. acetobutylicum* ATCC824. The mean values calculated from at least three experiments, the accuracy is within ±10%.

<table>
<thead>
<tr>
<th>Enzyme</th>
<th>$K_{m}^{NADH}$, 367/8287</th>
<th>$V_{max}$, 367/8287</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>L. brevis</em> ATCC 367/8287</td>
<td></td>
<td></td>
</tr>
<tr>
<td>LDH</td>
<td>201/220</td>
<td>37/26.5</td>
</tr>
<tr>
<td>ALDH</td>
<td>204/200</td>
<td>0.02/0.03</td>
</tr>
<tr>
<td>ADH</td>
<td>82/108</td>
<td>10.8/10.8</td>
</tr>
<tr>
<td>NOX</td>
<td>23/25</td>
<td>0.2/1.7</td>
</tr>
<tr>
<td><em>C. acetobutylicum</em> ATCC 824</td>
<td></td>
<td></td>
</tr>
<tr>
<td>HBD</td>
<td>178</td>
<td>5.9</td>
</tr>
</tbody>
</table>

Figure 4. Intra cellular NAD$^+$/NADH ratio (A) and total [NAD$^+$ + NADH] concentrations (B) in *L. brevis* ATCC 8287 (*) and *L. brevis* ATCC 367 (m) strains grown on MRS medium. The mean values calculated from at least three experiments. The accuracy of measurements is within ±10%.

Figure 5. Metabolism of pyruvate in *L. brevis*. The solid arrows indicate active pathways. The dotted arrows—the enzymes are not active, but the putative genes are predicted in *L. brevis* genome. The hollow arrows—neither gene nor enzyme activity or metabolic product were found. L-LDH: L-lactate dehydrogenase, D-LDH: D-lactate dehydrogenase, ADH: alcohol dehydrogenase, ALDH: aldehyde dehydrogenase, PTA: phosphotransacetylase, ACK: acetate kinase, PDH: pyruvate dehydrogenase, PFL: pyruvate formate lyase, PDC: pyruvate decarboxylase, POX: pyruvate oxidase, FDH: formate dehydrogenase, ALS: acetolactate synthase, ALDB: alpha-acetolactate decarboxylase, BUTB: diacetyl acetoin reductase (butanediol dehydrogenase).

Pyruvate oxidase activity was detectable only at the late stationary phase in aerobic conditions being 0.02 and 0.01 U/mg of total cell protein for ATCC 8287 and ATCC 367 strains respectively. The putative pyruvate oxidase-encoding gene was identified in the *L. brevis* genome (Table 3).

Although PDH activity was not detected, the genes predictably encoding enzymes of PDH complex were identified in *L. brevis* ATCC 367 genome (Table 3). No PFL (EC 2.3.1.54) or PDC (EC 4.1.1.1) activities were detected under conditions applied, and the corresponding genes were not identified during genome analysis. Formate produced in the PFL reaction was not found among metabolic end-products.

The genes presumably encoding acetolactate synthase (ALS), alpha-acetolactate decarboxylase (ALDB) and diacetyl acetoin reductase also named butanediol dehydrogenase (BUTB) participating in synthesis of acetolactate, acetoin and 2,3-butenediol from pyruvate, were identified in *L. brevis* ATCC 367 genome (Table 3). However, these metabolites were not detected in cultural liquid during *L. brevis* semi-anaerobic or aerobic fermentation in MRS or HM media.

Thus, besides conversion into lactate, pyruvate may be converted into acetyl-phosphate by pyruvate oxidase in strictly aerobic conditions and under glucose limitation. No other overlaps between lactate and ethanol metabolic branches have been detected in *L. brevis*.

Data of metabolic stoichiometry and enzyme kinetics allowed developing the scheme of carbon and electron flow in *L. brevis* in anaerobic and aerobic glucose fermentation (Figure 6).
Anaerobically: 1 Glucose → 1 Lactate + 1 Ethanol + 1 CO₂ + 1 ATP
Aerobically: 1 Glucose + O₂ → 1 Lactate + 1 Acetate + 1 CO₂ + 2 ATP + H₂O

Figure 6. The scheme of carbon and electron flow in _L. brevis_ in anaerobic and aerobic glucose fermentation. The solid arrows indicate native permanently acting reactions. The dotted arrows indicate reactions acting preferably in anaerobic condition. The hollow arrows indicate reactions active aerobically. NOX: NADH oxidase, LDH: lactate dehydrogenase, ADH: alcohol dehydrogenase, ALDH: aldehyde dehydrogenase, PK: phosphoketolase, PTA: phosphotransacetylase, ACK: acetate kinase, POX: pyruvate oxidase.

4. Discussion


_L. brevis_ metabolism is based on carbohydrate fermentation coupled with substrate level phosphorylation. Oxidation of a substrate leads to formation of NADH from NAD⁺, which has to be regenerated continuously. Stochiometry analysis of fermentation in various redox conditions (Table 2) proved that _L. brevis_ utilizes sugars through the 6-phosphogluconate (6-PG) pathway. The crucial enzyme of the 6-PG pathway is phosphoketolase, which converts xylulose-5-phosphate (C₅) into glyceraldehyde-3-phosphate (C₃) and acetyl phosphate (C₂) (Figure 6). In semi-anaerobic glucose fermentation this bifurcation leads to formation of lactate (C₃) and ethanol (C₂) (Table 2, Figure 3(a)) that have a key role in NAD⁺ regeneration.

In comparison with the lactate branch, the NADH-oxidizing capacity of the ethanol branch is very low: ALDH activity was about three orders lower than LDH activity (Table 4). It makes ALDH the bottleneck of the _L. brevis_ anaerobic metabolism, despite the highly active ADH following the ALDH in the ethanol-forming pathway. The ethanol branch does not bring additional ATP to the cells and it participates only in the maintenance of red-ox balance that determines the low level of biomass accumulation by _L. brevis_ grown semi-anaerobically in MRS medium (Figure 1(a)).

In aerobic conversion of hexoses, the oxygen acts as the external electron acceptor in a reaction catalyzed by NADH:H₂O oxidase (NOX) (Figure 6). Due to significantly higher activity and affinity to NADH compared to ALDH (Table 4), the NOX has the strong benefit of NAD⁺ regeneration and thereby prevents ethanol formation. The acetyl-phosphate is completely redirected from acetyl-CoA formation towards the acetate formation (Figure 3(b)) accompanied by synthesis additional ATP in acetate kinase reaction (Figure 6), and hence faster aerobic growth is observed in comparison to semi-anaerobic conditions (Figure 1(a)).

The rapid growth of ATCC 8287 versus ATCC 367 in MRS medium in aerobic conditions (Figure 1(a)) was obviously dependent on higher NOX activity of the ATCC 8287 strain: 1.7 U/mg in ATCC 8287 vs 0.20 U/mg in ATCC 367 (Table 4).

In the recombinant _L. brevis_ strains the native ethanol and recombinant butanol pathways originate from acetyl-CoA [9]. 3-Hydroxybutyryl-CoA dehydrogenase (HBD) is a first NADH-dependent enzyme of the butanol pathway. In aerobic conditions, the NOX, because of its comparatively high activity and affinity to NADH, efficiently competed for reducing equivalents not only with the ALDH, but with the HBD: the affinity towards NADH of the HBD was about 7 times lower than the NOX affinity (Table 4). For this reason, the redirection of the electron and carbon flow towards butanol synthesis was not achieved aerobically, and the recombinant _L. brevis_ strains produced butanol only in semi-anaerobic fermentation during exponential growth phase [9].

An active role of oxygen and NOX in growth and energy metabolism of various LAB species has been demonstrated in previous studies [35-37]. For example it was shown that the growth rate and the Yₕₑₗ of _Leuconostoc_ sp in aerated cultures were higher compared to non-aerated. An NOX-deficient mutant did not shift from ethanol to acetate production, and Yₕₑₗ was the same aerobically and anaerobically [37].

The ratio of NAD⁺/NADH and total intracellular [NAD⁺ + NADH] concentration correlated with the sugar consumption and influenced the pathway fluxes at the level of various dehydrogenase reactions. In the loga-
rhythmic growth phase of a batch culture, the NAD\(^+\)/NADH ratio increased (Figure 4(a)) while total [NAD\(^+\) + NADH] concentration decreased (Figure 4(b)). The intracellular NADH concentration dropped from 700 to 50 \(\mu\)M consequently passing through the \(K_m\) values of LDH, ALDH, ADH, and hence the activity of the corresponding enzymes decreased. In this way the cell automatically regulates carbon fluxes in response to environmental conditions.

4.2. Fermentation of Complex Sugar Substrates

Anaerobic heterolactic fermentation of 1 mole glucose through 6-PG pathway gives theoretically 1 mole of lactic acid, ethanol, CO\(_2\) and ATP. Semi-aerobic incubation of \(L.\ brevis\) in MRS media supplemented with glucose resulted in almost theoretical distribution of lactate and ethanol [9].

A complex fermentation medium changes the balance and the composition of end-products. Lactate, ethanol, acetate and mannitol were produced during semi-aerobic fermentation of MRS medium containing glucose and fructose (Figure 3(a), Table 2). Mannitol formation indicates that fructose is not only used as a carbon source, but as an additional electron acceptor, and metabolic intensity prevails over efficiency of substrate utilization. Reduction of fructose to mannitol may be catalyzed by NAD\(^+\)-mannitol dehydrogenase that is presumably encoded in \(L.\ brevis\) ATCC 367 by LVIS 2162 gene. This reaction allows overcoming the low capacity of the ethanol branch for NAD\(^+\) regeneration arising at the level of aldehyde dehydrogenase [38]. Using the mannitol pathway in fructose fermentation was earlier described with \(Oenococcus\ oeni\) [39].

In aerobic conditions NOX prevented not only the ethanol formation, but also the use of fructose as an electron acceptor (Figure 3(b), Table 2). \(L.\ brevis\) strains can utilize hexoses (glucose, galactose), pentoses (xylose, arabinose) and uronic acids of SEW-hydrolyzate (Figure 2, Table 2). No CO\(_2\) is formed during pentose fermentation in 6-PG pathway and since no dehydrogenation steps are necessary to reach the xylose-5-phosphate. Consequently the ethanol formation becomes redundant. Instead, acetyl phosphate is used by the acetate kinase yielding acetyl-CoA and ATP. Fermentation of pentoses thus leads to production of equimolar amounts of lactic and acetic acids [32]. Therefore fermentation of HM medium resulted in lower ethanol and higher acetate ratio as well as higher biomass accumulation level compared to MRS medium containing glucose and fructose (Table 2). The uronic acids present in SEW-hydrolyzate may be utilized to form lactate and acetate (http://www.genome.jp/kegg-bin/show_pathway?lbz00040).

4.3. Role of Pyruvate Oxidase in \(L.\ brevis\) Metabolism

Under aerobic conditions and sugar limitation acetate is produced at the expense of lactate as glucose becomes depleted and \(L.\ brevis\) enters the stationary phase of growth (Figures 3(b) and (d)). Various LAB species can use lactate as a carbon source [40,41]. The pathway of lactate to acetate conversion could be performed via three enzymatic steps: oxidation of lactate to pyruvate by the NADH-dependent D- and L-LDHs, oxidative decarboxylation of pyruvate to acetyl-phosphate by pyruvate oxidase, and dephosphorylation of acetyl-phosphate to acetate by acetate kinase (Figures 5, 6). ATP formed in ACK reaction provides the cells with energy in the stationary phase. The LDH-POX-ACK pathway is regulated at the level of pyruvate-oxidase activity, which is induced by O\(_2\) and repressed by glucose [41-45]. In the \(L.\ brevis\) ATCC 367 genome, the putative pyruvate oxidase-encoding gene was identified. The pyruvate oxidase activity was detected at the late stationary phase of aerobic growth. Lactate to acetate conversion could also be performed aerobically by oxygen-dependent lactate oxidase (lactate 2-mono-oxidase). However, no similarity with the corresponding genes was found in \(L.\ brevis\) genome analysis.

Thereby in aerobic conditions and under glucose limitation \(L.\ brevis\) cells may use pyruvate oxidase as a link between pyruvate and acetyl-phosphate in a pathway from lactate to acetate. Since anaerobic conditions are required for heterologous butanol synthesis, the aerobically active pyruvate oxidase cannot be used for redirecting carbon flux from lactate to butanol synthesis.

4.4. Proposed Strategy for Redirection \(L.\ brevis\) Carbon and Electron Flow towards Butanol Synthesis

In recombinant butanol-producing \(L.\ brevis\) strains, acetyl-CoA is the butanol precursor; the reducing equivalents are necessary for enzymes of heterologous butanol pathway. Metabolic study suggests that in \(L.\ brevis\) the acetyl-CoA is formed from acetyl-phosphate in anaerobic glucose fermentation, \(i.e.\) in conditions where NADH-oxidase is inactive and NADH equivalents are available for the enzymes of ethanol biosynthesis. Thus anaerobic conditions are required for heterologous butanol synthesis. Semi-aerobic cultivation is possible too because oxygen availability in the beginning of fermentation meets the energetic needs of cells allowing intensive growth. Presence of pentoses, uronic acids and external electron acceptors other to oxygen (e.g. fructose) in fermentation medium enhances cells growth but unfavorable for butanol synthesis.

Inactivation of the LDHs could not only shut off the
competitive carbon flux, but it could partly compensate NADH-deficiency of butanol pathway by saving 1 mole NADH per 1 mol of butanol. The surplus of pyruvate should be redirected toward butanol synthesis. However, enzymes converting pyruvate into acetyl-CoA in anaerobic conditions have not been detected in \textit{L. brevis}. Therefore pyruvate redirection may be organized by cloning the pyruvate formate lyase genes (Figure 7).

The anaerobically active PFL enzyme system has been shown to be operational in several LAB species [46-48]. Cloning the formate dehydrogenase (FDH) might help to utilize formate produced in PFL reaction and provide an additional mole of NADH per mol of 1-butanol produced (Figure 7).

Despite the principal possibility to support the butanol synthesis [9], the native \textit{L. brevis} ALDH, ADH and THL, and recombinant clostridial BCD/EtfAB cannot provide a high capacity pathway for butanol because of 1) low activity of ALDH and THL, 2) low specificity towards C4 substrate of ALDH and ADH [9] and 3) cofactor (FAD and ferredoxin) deficiency for BCD/EtfAB [49]. These enzymes should be enhanced or substituted with efficient analogues.

5. Conclusions

Carbon and electron flow was investigated in two \textit{L. brevis} strains in order to assess metabolic potential of these bacteria for heterologous butanol synthesis. The study approach was based on analysis of fermentation stoichiometry in various redox conditions together with catalytic properties \( K_{\text{NADH}} \) and \( V_{\text{max}} \) of the NADH-dependent enzymes from the main metabolic pathways.

Conditions necessary for generation of acetyl-CoA and NADH required for butanol biosynthesis have been determined. Key enzymes controlling direction of carbon and electron flow in \textit{L. brevis} were defined.

In anaerobic glucose fermentation, the carbon flow through acetyl-CoA and the slow NAD\(^+\) regeneration rate are controlled by aldehyde dehydrogenase ALDH \( K_{\text{NADH}} = 200 \mu M, V_{\text{max}} = 0.03 \text{ U/mg} \). Aerobically, the NOX, due to its comparatively high affinity to NADH and activity \( K_{\text{NADH}} = 25 \mu M, V_{\text{max}} = 1.7 \text{ U/mg} \), efficiently competes with ALDH for NADH that results in redirecting carbon flow from acetyl-CoA towards acetate formation.

In recombinant butanol-producing strains the enzymes of the native ethanol and heterologous butanol pathways compete for acetyl-CoA and NADH. Aerobically, the NOX, because of low \( K_{\text{NADH}} \), prevents not only the ethanol but also the butanol formation.

In general, availability of external electron acceptors (oxygen, fructose) enhances biomass accumulation associated with additional ATP synthesis in acetate kinase reaction, but reduces NADH availability for enzymes of ethanol pathway and, therefore, decreases carbon flux through acetyl-CoA.

Pyruvate metabolism was investigated to find redirection possibilities of competitive carbon fluxes towards butanol synthesis. Pyruvate may be converted into acetyl-phosphate by pyruvate oxidase in strictly aerobic conditions and under glucose limitation. No other overlaps between lactate and ethanol metabolic branches have been detected in \textit{L. brevis}.

\textit{L. brevis} strains ferment glucose, galactose, arabinose, and xylose from SEW-hydrolysate of spruce chips, but not utilize mannose. Increased acetate formation accompanied by enhanced cell growth and decreased ethanol synthesis indicated reduced carbon flow though the acetyl-CoA. Thus, SEW-hydrolysate could be used for initial biomass accumulation of butanol-producing \textit{L. brevis} strains. Using SEW-hydrolysate for butanol synthesis requires optimization of fermentation media in order to increase NADH availability for enzymes of butanol pathway.

The study provides a basis for rational development of \textit{L. brevis} strains producing butanol from SEW-hydrolysate and proposes possible engineering ways for rerouting carbon and electron flow towards butanol synthesis.

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