Communication

Production of Ethylene Glycol from Glycerol Using an In Vitro Enzymatic Cascade

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Abstract: Glycerol is a readily available and inexpensive substance that is mostly generated during biofuel production processes. In order to ensure the viability of the biofuel industry, it is essential to develop complementing technologies for the resource utilization of glycerol. Ethylene glycol is a two-carbon organic chemical with multiple applications and a huge market. In this study, an artificial enzymatic cascade comprised alditol oxidase, catalase, glyoxylate/hydroxypyruvate reductase, pyruvate decarboxylase and lactaldehyde:propanediol oxidoreductase was developed for the production of ethylene glycol from glycerol. The reduced nicotinamide adenine dinucleotide (NADH) generated during the dehydrogenation of the glycerol oxidation product D-glycerate can be as the reductant to support the ethylene glycol production. Using this in vitro synthetic system with self-sufficient NADH recycling, 7.64 ± 0.15 mM ethylene glycol was produced from 10 mM glycerol in 10 h, with a high yield of 0.515 ± 0.1 g/g. The in vitro enzymatic cascade is not only a promising alternative for the generation of ethylene glycol but also a successful example of the value-added utilization of glycerol.

Keywords: glycerol; ethylene glycol; biocatalysis; in vitro enzymatic cascade

1. Introduction

The manufacturing of biofuels from reproducible feedstocks is increasing throughout the world [1–3]. The development of the biofuel industry has generated a large amount of glycerol as an inevitable byproduct [4]. An excess of glycerol production in the biofuel industry leads to a dramatic decrease of the price of glycerol, which has a negative impact on the development of the biofuels industry [5]. Thus, the conversion of glycerol into value-added compounds through chemical or biotechnological routes is now urgently needed for the improvement of the viability of the biofuel industry.

Ethylene glycol is a large-volume commodity that is widely used in many industrial processes, such as glycolic acid production and polyethylene terephthalate synthesis [6]. It is predominantly produced from ethylene oxide through carbonation and subsequent hydrolyzation, or by the hydration of ethylene oxide [7,8]. Direct ethylene glycol production from biomass-derived carbohydrates through chemical catalysis has also been reported [9]. Importantly, ethylene glycol can be produced from glycerol through liquid-phase catalytic hydrogenolysis [10,11]. However, this process usually needs high pressures (4.0–8.0 MPa), high temperatures (453–473 K) and expensive noble metal catalysts (Pt, Ni, Ru, etc.). Several metabolic pathways have been constructed for the biotechnological production of ethylene glycol from renewable substrates such as D-glucose, D-xylose and L-arabinose [12–14]. Nevertheless, the production of ethylene glycol from glycerol via a biotechnological route has never been reported, until now.
An in vitro synthetic system is emerging as a promising technique for the synthesis of high-value products from cheap and renewable substrates [15,16]. This biomanufacturing platform leaves out the use of cells, and can thus exclude cell-associated process barriers, such as the transmembrane transport of substances, the toxicity of products, and undesired metabolism pathways. In addition, an in vitro synthetic system exclusively employs purified enzymes, and makes the purification of the target products rather simple to conduct. Some in vitro synthetic systems have been constructed for the generation of valuable chemicals from renewable substrates, such as D-glucose, D-xylose and glycerol [17–19]. In this study, we have designed an artificial enzymic cascade for the conversion of glycerol to ethylene glycol that only requires five enzymes. The artificial pathway is also completely redox balanced through self-sufficient reduced nicotinamide adenine dinucleotide (NADH) recycling.

2. Results and Discussion

2.1. Design of the In Vitro Biosystem for Ethylene Glycol Production

Here, an artificial enzymatic reaction cascade was designed for the conversion of glycerol into ethylene glycol (Figure 1). Glycerol is first oxidized to D-glycerate via D-glyceraldehyde by alditol oxidase (AldO) catalyzed two-step oxidation. The D-Glyceraldehyde produced from glycerol is then transformed into 3-hydroxypyruvate by oxidized nicotinamide adenine dinucleotide (NAD)+-dependent glyoxylate/hydroxypyruvate reductase (GRHPR). Then, 3-hydroxypyruvrate is decarboxylated to glycolaldehyde by pyruvate decarboxylase (PDC). Finally, the glycolaldehyde is reduced into ethylene glycol by lactaldehyde:propanediol oxidoreductase (FucO), using the NADH produced during the oxidation of the glycerol is decomposed by catalase in order to avoid the spontaneous oxidative of the intermediate products [20]. As a result, one mole of glycerol can generate one mole of ethylene glycol.

![Diagram](https://via.placeholder.com/150)

**Figure 1.** Scheme of ethylene glycol production from glycerol based on the artificial in vitro enzymatic cascade. The enzymes are alditol oxidase (AldO) from *Streptomyces coelicolor* A3(2), catalase from *Aspergillus niger*, glyoxylate/hydroxypyruvate reductase (GRHPR) from *Pyrococcus furiosus*, pyruvate decarboxylase (PDC) from *Zymomonas mobile*, and lactaldehyde:propanediol oxidoreductase (FucO) from *Escherichia coli*.

2.2. Substrate Specificity of AldO<sub>mt</sub>

AldO is a soluble monomeric flavoprotein containing covalently-bound FAD as its cofactor [21]. This enzyme is first isolated from *Streptomyces coelicolor* A3(2) and identified as an oxidase primarily active with alditols, and as producing aldoses as products. Interestingly, AldO exhibits regio- and enantio-selective oxidative activity toward 1,2-diols and α-hydroxy carboxylic acids but not the α-hydroxy aldehydes which will be formed as the end products [22]. AldO<sub>mt</sub> with enhanced oxidative activity towards glycerol was also
constructed through directed evolution, and this enzyme has been successfully applied in D-glycerate production from glycerol [23].

The target product of this study, ethylene glycol, is the simplest 1,2-diol, and is also the possible substrate of AldO\textsubscript{mt}. Thus, the activity of AldO\textsubscript{mt} toward ethylene glycol was assayed using a Clark-type oxygen electrode, as described in a previous study [24]. As shown in Figure 2a, the AldO\textsubscript{mt}-dependent oxygen consumption was rather low with the addition of ethylene glycol, while noticeable oxygen consumption was observed when glycerol was added (Figure 2c). The estimated specific activities of AldO\textsubscript{mt} for glycerol and ethylene glycol were 0.53 ± 0.03 U/mg and 0.018 ± 0.004 U/mg, respectively (Figure 2b). The oxidative activities of AldO\textsubscript{mt} toward glycerol and ethylene glycol were also determined by high-performance liquid chromatography (HPLC) analysis. As shown in Figure 2c,d, the depletion of glycerol could be observed obviously, while very little consumption of ethylene glycol was detected.

Figure 2. Substrate specificity of AldO\textsubscript{mt}. (a) The determination of the oxygen consumption during the AldO\textsubscript{mt}-catalyzed oxidation of the target substrates. (b) The specific activity of AldO\textsubscript{mt} toward glycerol, D-glyceraldehyde, glycolaldehyde, and ethylene glycol, respectively. The error bars indicate the standard deviations from three parallel replicates. (c,d) HPLC analysis of AldO\textsubscript{mt}-catalyzed glycerol (c) or ethylene glycol (d) oxidation. The reactions were carried out in a reaction mixture containing 50 mM 4-(2-hydroxyethyl)-1-piperazineethanesulfonic acid (HEPES)-NaOH buffer (pH 7.4), 10 mM glycerol or 10 mM ethylene glycol and 0.2 mg/mL AldO\textsubscript{mt}, at 30 °C, with a shaking speed of 150 rpm.

AldO oxidizes glycerol into D-glycerate in a two-step reaction via D-glyceraldehyde [23]. Glycolaldehyde, the direct precursor of ethylene glycol in the designed enzymatic reaction cascade, is the analogue of D-glyceraldehyde, and might be oxidized by AldO\textsubscript{mt}. Thus, the activities of AldO\textsubscript{mt} toward glyceraldehyde and D-glyceraldehyde were also assayed. As shown in Figure 2a,b, a very much lower oxygen consumption rate was observed when glyceraldehyde, but not D-glyceraldehyde, was added to the reaction mixture, indicating that AldO\textsubscript{mt} exhibits rather low oxidative activity toward glyceraldehyde. The result of the HPLC analysis further confirmed that AldO\textsubscript{mt} can hardly catalyze the oxidation of glyceraldehyde (Figure S1).
Based on the data mentioned above, AldO\textsubscript{mt} can efficiently oxidize glycerol into D-glycerate, but will barely consume the glycolaldehyde and ethylene glycol in the reaction system. Thus, AldO\textsubscript{mt} is a proper catalyst in the designed enzymatic reaction cascade, and it exhibits little promiscuous activity affecting the production of ethylene glycol.

2.3. 3-Hydroxyxypyruvate Decarboxylation Catalyzed by PDC

The GRHPR of \textit{Pyrococcus furiosus} can catalyze the NAD\textsuperscript{+}-dependent dehydrogenation of D-glycerate to produce 3-hydroxyxypyruvate \cite{27,28}. However, the standard Gibbs energy change (\(\Delta r\text{G}^\theta\)) of D-glycerate dehydrogenation is estimated to be 20.26 KJ/mol, indicating that this reaction is not thermodynamically favorable. In order to support the GRHPR-catalyzed, thermodynamically unfavorable D-glycerate dehydrogenation, the integration of thermodynamically favorable reactions with 3-hydroxyxypyruvate or NADH as the substrates is needed.

The PDC of \textit{Zymomonas mobilis} can catalyze the decarboxylation of pyruvate to produce carbon dioxide and acetaldehyde \cite{26}. In this study, the activity of PDC toward 3-hydroxyxypyruvate decarboxylation, a thermodynamically favorable reaction (\(\Delta r\text{G}^\theta = -55.23\) KJ/mol), was also assayed. As shown in Figure S2, PDC also exhibited decarboxylation activity to catalyze 3-hydroxyxypyruvate into glycolaldehyde. About 8 mM glycolaldehyde was produced from 10 mM 3-hydroxyxypyruvate in 4 h by 10 \(\mu\)g/mL PDC.

2.4. Substrate Specificity of FucO

The NADH-dependent reduction of glycolaldehyde can regenerate the NAD\textsuperscript{+} required for D-glycerate dehydrogenation, and can support the production of ethylene glycol. The lactaldehyde:propanediol oxidoreductase of \textit{Escherichia coli} (FucO) was reported to reduce glycolaldehyde \cite{27,28}. This enzyme has been overexpressed in \textit{E. coli} for in vivo ethylene glycol production via a synthetic pathway \cite{29}. In this study, the activity of FucO using glycolaldehyde and NADH as the substrates was also assayed (Figure 3a,b). The specific activity of FucO toward glycolaldehyde reduction was calculated to be 30.9 \(\pm\) 1.2 U/mg (Figure 3b). The HPLC analysis results showed that FucO rapidly reduces glycolaldehyde into ethylene glycol (Figure 3c).

![Figure 3](https://example.com/figure3.png)

**Figure 3.** Substrate specificity of FucO. (a) Time course of reactions of FucO with glycerol, D-glycerate, glycolaldehyde, or ethylene glycol as the substrate and NADH as the cofactor. (b) Specific activity of FucO toward glycerol, D-glycerate, glycolaldehyde, and ethylene glycol, respectively. The error bars indicate the standard deviations from three parallel replicates. (c) HPLC analysis of the FucO-catalyzed glycolaldehyde reduction. The reaction was carried out in a reaction mixture containing 50 mM HEPES-NaOH buffer (pH 7.4), 10 mM glycolaldehyde, 10 mM NADH and 0.5 mg/mL FucO, at 30 °C, with a shaking speed of 150 rpm.

The activities of FucO toward other reactants in the in vitro system including ethylene glycol, glycerol and D-glyceraldehyde were also assayed. As shown in Figure 3a,b, all of these reactant especially D-glyceraldehyde, the analogue glycolaldehyde, cannot be reduced by FucO. Thus, the NADH generated during D-glycerate oxidation can only be consumed for glycolaldehyde reduction. This property of FucO ensured the self-sufficient NADH
recycling of the in vitro system, and avoided the requirement for exogenous reducing agent addition.

2.5. Production of Ethylene Glycol from Glycerol In Vitro

AldO\textsubscript{mt}, GRHPR, PDC and FucO were overexpressed, purified (Figure S3), and coupled with commercial catalase from \textit{A. niger} (Sigma-Aldrich, St. Louis, MO, USA) for the production of ethylene glycol from glycerol. As shown in Figure 4, 7.64 ± 0.15 mM ethylene glycol was generated from 10 mM glycerol in 10 h. Some chemists have also attempted to produce ethylene glycol from glycerol by hydrolysis. For example, Feng and his colleagues studied the conversion of glycerol to ethylene glycol over Ru–Co/ZrO\textsubscript{2} catalysts [30]. However, this catalyst exhibited higher selectivity toward 1,2-propanediol generation than ethylene glycol production. The yield of ethylene glycol was only 0.30 g/g glycerol. Ueda and his colleagues reported the conversion of glycerol to ethylene glycol over Pt-modified Ni catalysts under the conditions of 453 K and 8.0 MPa [10]. The yield of ethylene glycol was 0.479 g/g, which was lower than that of a biocatalytic process utilizing the artificially designed enzymatic cascade (0.515 ± 0.1 g/g) under modest reaction conditions (303 K, 0.1 MPa).

Some biotechnological routes have been developed for the production of ethylene glycol from biomass-derived carbohydrates [12–14,19,29,31–35] (Table 1). For example, Feng and his colleagues disrupted \textit{arcA} [29]. Recently, a cell-free bioreaction scheme was also reported to generate ethylene glycol from D-xylene, with a yield of 0.227 g/g [19]. In this study, we designed an artificial enzymatic cascade to produce ethylene glycol from glycerol with only one carbon atom loss. A high yield of 0.515 ± 0.1 g/g ethylene glycol from glycerol was acquired using the in vitro synthetic system.

![Figure 4. Conversion of glycerol into ethylene glycol by the artificial enzymatic reaction cascade. The reaction was carried out in a reaction mixture containing 50 mM HEPES-NaOH buffer (pH 7.4), 10 mM glycerol, 1 mM NAD\textsuperscript{+}, 2 mM MgSO\textsubscript{4}, 0.25 mM thiamine pyrophosphate (TPP), 1 mg/mL AldO\textsubscript{mt}, 10 mg/mL GRHPR, 5.4 mg/mL PDC, 13 mg/mL FucO and 500 U/mL catalase. The reaction was conducted at 30 °C and a shaking speed of 150 rpm. The error bars indicate the standard deviations from three parallel replicates.](image-url)
Table 1. Yield of ethylene glycol production from various substrates by metabolically-engineered strains or an in vitro biosystem.

| Strain or Enzymes                          | Substrate   | Ethylene Glycol Yield (g/g) | Reference |
|-------------------------------------------|-------------|----------------------------|-----------|
| E. coli BW25113(DE3)                      | D-Xylose    | 0.29                       | [12]      |
| E. coli MG1655 ΔxylB/ pET20-ΔldC          | D-Xylose    | 0.19                       | [13]      |
| E. coli BL21(DE3) ΔarcAΔldC/pETDuet1-ΔldC | D-Xylose    | 0.40                       | [29]      |
| E. coli MG1655(DE3) ΔendAΔrecA ΔxylBΔldC/pT5-Pc.dte-fucO-fucK | D-Xylose | 0.35                       | [33]      |
| E. coli W3110 ΔlacI/pTacxylBC-ΔldC/D1-anti-ΔldC | D-Xylose | 0.36                       | [34]      |
| E. coli W3110 ΔxylAΔldC/pKMX & pTrcHis2A-ΔldC | D-Xylose | 0.39                       | [35]      |
| Enterobacter cloacae S1                   | Xylose acid | 0.288                      | [32]      |
| E. coli MG1655(DE3) ΔendAΔrecAΔxylBΔldC/p5_T7-Pc.dte-rbaB-rhaD-fucO | l-Arabinose | 0.38                       | [33]      |
| E. coli MG1655(DE3) ΔldDΔeutBC/p10_T7-satC-ΔldC-satC-fucO & pCDFDuet1-ΔldD-ΔldD-ΔldD-ΔldD | D-Glucose | 0.14                       | [31]      |
| C. glutamicum PABS1/pec-P0-GldP-Pmdc-ΔldC | D-Glucose | 0.09                       | [14]      |
| AldOmt, catalase, GRHPR, PDC, FucO         | Glycerol    | 0.515                      | This study |

The in vitro synthetic system has appealing advantages, such as a high product yield, easy product separation, and simple process control [16]. It has made great progress in recent years. For example, the production of inositol from starch using the in vitro synthetic system has been implemented on an industrial scale [36]. In this study, we designed an enzymatic cascade for the production of ethylene glycol from glycerol. Although a higher yield of ethylene glycol was obtained using the artificially designed enzymatic cascade, the final concentration and productivity of ethylene glycol cannot meet the requirements for industrial application. The low activity of GRHPR may be the key factor restricting the efficiency of the in vitro enzymatic cascade. Systematic screening or the directed evolution of a new biocatalyst with higher activity toward the dehydrogenation of D-glycerate to 3-hydroxypyruvate will be necessary to further improve the production of ethylene glycol. The optimization of the reaction system may also improve the efficiency of the enzymatic cascade, and this deserves an intensive study [37].

3. Materials and Methods

Materials: reduced/oxidized nicotinamide adenine dinucleotide (NADH/NAD\(^+\)), isopropyl-β-D-1-thiogalactopyranoside (IPTG), thiamine pyrophosphate (TPP), 3-hydroxypyruvate, D-glycerate and glycolaldehyde were purchased from Sigma-Aldrich (St. Louis, MO, USA). Restriction endonuclease and T4 DNA ligase were purchased from Thermo Scientific (Waltham, MA, USA). FastPfu DNA polymerase was purchased from TransGen Biotech Ltd. (Beijing, China). The other chemicals used in this study were analytical pure grade and commercially available.

The cloning and expression of fucO: the strains, plasmids and primers used in this study are listed in Supplementary Materials Table S1. E. coli DH5α was used for the plasmid construction, and E. coli BL21(DE3) was used for the overexpression of the recombinant proteins. The encoding gene of FucO was amplified from the genomic DNA of E. coli K12 with the primers fucO-t/fucO-r (Supplementary Table S1), digested with BamHI/HindIII, and cloned into pET28a in order to construct pET28a-FucO. The plasmid pET28a-FucO was then transferred into E. coli BL21(DE3) to obtain E. coli BL21(DE3)/pET28a-FucO. All of the E. coli strains were cultivated in Luria-Bertani (LB) medium at 37 °C and 180 rpm or 16 °C.
and 160 rpm. If necessary, ampicillin (at a final concentration of 100 µg/mL) or kanamycin (at a final concentration of 50 µg/mL) was added into the medium.

Purification of the enzymes: the catalase from *A. niger* used in this study was purchased from Sigma-Aldrich (St. Louis, MO, USA). The AldO<sub>mt</sub>, GRHPR and PDC were purified as described by Li and his colleagues [20]. *E. coli* BL21(DE3)/pET28a-FucO was grown at 37 °C and 180 rpm in LB medium with 100 µg/mL kanamycin to an optical density of 0.5 at 620 nm. Then, IPTG was added at a final concentration of 1.0 mM in order to induce the expression of FucO. After cultivation for another 4 h at 37 °C, the cells were harvested and washed twice with 20 mM phosphate solution buffer (PBS, pH 7.4) by centrifugation at 6000 rpm for 10 min; the cells were subsequently suspended in binding buffer (20 mM phosphate buffer, 500 mM NaCl, and 20 mM imidazole, pH 7.4) and lysed using a continuous high-pressure cell disrupter working at 4 °C and 900 bar. The cell lysate was centrifugated at 12,000 rpm for 1 h, and the supernatant then was loaded onto a 5 mL HisTrap HP column (GE Healthcare, Sweden). The target proteins were eluted with a gradient ratio of elution buffer (20 mM phosphate buffer, 500 mM NaCl, and 500 mM imidazole, pH 7.4). The purified proteins including AldO<sub>mt</sub>, GRHPR, PDC and FucO were subjected to sodium dodecyl sulphate-polyacrylamide gel electrophoresis (SDS-PAGE), and the results are shown in Supplementary Figure S3.

Ethylene glycol production: the production of ethylene glycol from glycerol was performed in a 5 mL reaction system containing 1 mg/mL AldO<sub>mt</sub>, 500 U/mL catalase, 10 mg/mL GRHPR, 5.4 mg/mL PDC, 10 mg/mL FucO, 10 mM glycerol, 1 mM NAD<sup>+</sup>, 0.25 mM TPP, 2 mM MgSO<sub>4</sub>, and 50 mM HEPES-NaOH (pH 7.4) at 30 °C and 150 rpm. Samples (0.2 mL) were withdrawn every 2 h and subjected to analysis by HPLC.

Analytical methods: the concentrations of the purified proteins were measured by the use of the Bradford protein assay (Bio-Rad, Hercules, CA, USA). The activity of the FucO was assayed by recording the absorption of NADH at 340 nm using a UV–visible spectrophotometer (Ultrospec 2100 pro; Amersham Biosciences, Little Chalfont, UK) [20]. All of the reactions were carried out in a 1.0 mL quartz cuvette at 30 °C, and HEPES-NaOH buffer (50 mM, pH 7.4) with 0.2 mM NADH and 2.5 mM substrate (glycerol, D-glyceraldehyde, glycolaldehyde or ethylene glycol). One unit of FucO activity was defined as the amount of protein catalyzing the consumption of one µmol of NADH per minute.

The oxidase activities of AldO<sub>mt</sub> toward glycerol, D-glyceraldehyde, glycolaldehyde and ethylene glycol were assayed by measuring the rate of oxygen consumption. The reaction mixture contained 10 mM substrate, 50 mM HEPES-NaOH buffer (pH 7.4), and 0.2 mg/mL AldO<sub>mt</sub>. The oxygen consumption in the reaction system was monitored using a Clark-type oxygen electrode (Oxytherm System; Hansatech Instruments Ltd., Pentney, UK). One unit of AldO<sub>mt</sub> activity was defined as the amount of protein catalyzing the consumption of one µmol O<sub>2</sub> per minute.

The concentrations of glycerol, glyceraldehyde, D-glycerate and 3-hydroxypyruvate were analyzed using an HPLC system (Agilent 1100 series, Hewlett-Packard, Palo Alto, CA, USA) equipped with an anion exchange column (Aminex HPX-87H, 300 × 7.8 mm; Bio-Rad, USA) and a refractive index detector [20]. The mobile phase was 5.0 mM H<sub>2</sub>SO<sub>4</sub> at a flow rate of 0.4 mL/min and at a temperature of 55 °C. The concentrations of the analytes were calculated by converting the peak areas detected by HPLC using external calibration curves.

4. Conclusions

In conclusion, we designed an artificial in vitro biosystem involving five enzymes for the production of ethylene glycol from glycerol. Ethylene glycol was produced from glycerol with a yield of 0.515 ± 0.1 g/g, which is higher than any reported chemical or biotechnological routes to date. The method presented in this work would not only provide a promising process for ethylene glycol production, but would also expand the utilization of the glycerol produced by the biofuel industry.
Supplementary Materials: The following are available online at https://www.mdpi.com/2073-4344/11/2/214/s1. Figure S1: HPLC analysis of AldOmt-catalyzed d-glyceraldehyde (a) and glycolaldehyde (b) oxidation. The reactions were carried out in a reaction mixture containing 50 mM HEPES-NaOH buffer (pH 7.4), 0.2 mg/mL AldOmt, 10 mM d-glyceraldehyde or 10 mM glyceraldehyde, at 30 °C and 150 rpm. Figure S2: HPLC analysis of PDC catalyzed 3-hydroxypropionate decarboxylation. The reaction was carried out in 50 mM HEPES-NaOH buffer (pH 7.4) containing 2 mM MgSO4, 0.25 mM TPP, 10 mM 3-hydroxypropionate and 10 µg/mL PDC, at 30 °C and 150 rpm. Figure S3: SDS-PAGE results of the purified enzymes. M: Marker; 1: AldOmt; 2: GRHPR; 3: PDC; 4: FucO. Table S1: Strains, plasmids and primers used in this study.

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Conflicts of Interest: The authors declare no conflict of interest.

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