



Supplementary Materials

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

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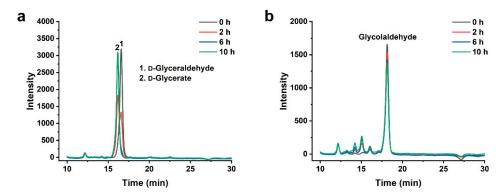


Figure S1. HPLC analysis of AldO_{mt} catalyzed D-glyceraldehyde (a) and glycolaldehyde (b) oxidation. The reactions were carried out in a reaction mixture containing 50 mM HEPES-NaOH buffer (pH 7.4), 0.2 mg/mL AldO_{mt}, 10 mM D-glyceraldehyde or 10 mM glycolaldehyde, at 30 °C and 150 rpm.

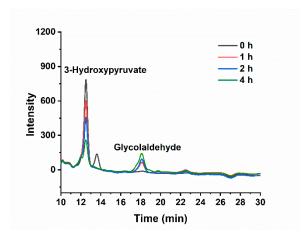


Figure S2. HPLC analysis of PDC catalyzed 3-hydroxypyruvate decarboxylation. The reaction was carried out in 50 mM HEPES-NaOH buffer (pH 7.4) containing 2 mM MgSO₄, 0.25 mM TPP, 10 mM 3-hydroxypyruvate and 10 μ g/mL PDC, at 30 °C and 150 rpm.

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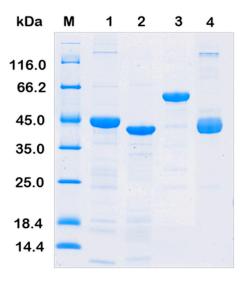


Figure S3. SDS-PAGE results of purified enzymes. M: Marker; 1: AldOmt; 2: GRHPR; 3: PDC; 4: FucO.

Table S1. Strains, plasmids and primers used in this study.

Strain, plasmid and primer	Description	Source
Strain		
E. coli DH5α	supE44 ΔlacU169 (φ80 lacZΔM15) hsdR17 recA1 endA1 gyrA96 thi-1 relA1	Novagen
E. coli BL21(DE3)	F - ompT hsdSB (rB-mB-) gal (λ c I 857 ind1 Sam7 nin5 lacUV5 T7gene1) dcm (DE3)	Novagen
E. coli K12	Wild-type	Lab stock
E. coli BL21(DE3)/pETDuet- AldO _{mt}	E. coli BL21(DE3) expressing AldO from Streptomyces coelicolor A3(2) with double mutant (V125M/A244T)	Lab stock
E. coli BL21(DE3)/pETDuet- GRHPR	E. coli BL21(DE3) expressing GRHPR from Pyrococcus furiosus	Lab stock
E. coli BL21(DE3)/pETDuet- PDC	E. coli BL21(DE3) expressing PDC from Zymomonas mobile	Lab stock
E. coli BL21(DE3)/p28a-FucO	E. coli BL21(DE3) expressing FucO from Escherichia coli K12	This study
Plasmid		
pET28a	Expression vector, kanamycin ^r	Novagen
pET28a-FucO	pET28a harboring gene fucO from Escherichia coli K12	This study
Primer	Sequence (5'→3')	
fucO-f	CG <u>GGATCC</u> ATGGCTAACAGAATGATTCTGAAC	This study
fucO-r	CCC <u>AAGCTT</u> TTACCAGGCGGTATGGTAAAGCTC	This study



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