

Table S1. Strains used in this study.

Strains	Genotype	Source
<i>E. coli</i> DH5a	F80d,lacZΔ(lacZYA_argF),U169,deoR,recA1,endA1,p hoA, hsdR17 (r -k , mk+) supE44λ_thi -1, gyrA96, relA1	Transgen
<i>E. coli</i> BL21(DE3)	F-, ompT, hsdS _B (r _B -, m _B -), gal, dcm (DE3)	Transgen
<i>E. coli</i> TOP10	F-, mcrA Δ(mrr-hsd RMS-mcr BC), φ80 lacZ ΔM15ΔlacX74, recA1, araΔ139Δ(ara-leu)7697, galU, galK, rpsL(Str ^R), endA1, nupG	This study
ECPLD1	<i>E. coli</i> BL21(DE3) carrying pET28a-SaPLD	This study
ECPLD2	<i>E. coli</i> BL21(DE3) carrying pET22b-SaPLD	This study
ECPLD3	<i>E. coli</i> BL21(DE3) carrying pET22b-OmpA-SaPLD	This study
ECPLD4	<i>E. coli</i> TOP10 carrying pBAD-gIIIC-OmpA-SaPLD	This study

Table S2. Plasmids used in this study.

Plasmids	Genotype	Source
pET28a	Kana, T7 promoter	GENEWIZ
pET22b- PelB	Amp, T7 promoter, pelB signal sequence	GENEWIZ
pBAD-gIIIC	Amp, pBAD promoter	GENEWIZ
pET28a-SaPLD	pET28a carrying PLD gene from <i>Streptomyces antibioticus</i>	This study
pET22b-SaPLD	pET22b carrying PLD gene from <i>Streptomyces antibioticus</i>	This study
pET22b-DsbA-SaPLD	pET22b carrying PLD gene from <i>Streptomyces antibioticus</i> , pelB signal sequence replaced by DsbA signal sequence	This study
pET22b-FhuD-SaPLD	pET22b carrying PLD gene from <i>Streptomyces antibioticus</i> , pelB signal sequence replaced by FhuD signal sequence	This study
pET22b-TorA-SaPLD	pET22b carrying PLD gene from <i>Streptomyces antibioticus</i> , pelB signal sequence replaced by TorA signal sequence	This study
pET22b-OmpA-SaPLD	pET22b carrying PLD gene from <i>Streptomyces antibioticus</i> , pelB signal sequence replaced by OmpA signal sequence	This study
pET22b-PhoA-SaPLD	pET22b carrying PLD gene from <i>Streptomyces antibioticus</i> , pelB signal sequence replaced by PhoA signal sequence	This study
pBAD-gIIIC-OmpA- SaPLD	pBAD-gIIIC carrying PLD gene from <i>Streptomyces antibioticus</i> and OmpA signal sequence	This study

Table S3. Primers and synthetic oligos used in this study.

No.	Primer	Nucleotide sequence (5' >3')
1	DsbA-SaPLD-1F	GCCTGGTGCTGGCGTTTAGCGCTAGCGCCATGGATATCGGAATTAATTCGGATCCGG
2	DsbA-SaPLD-1R	CTCGAGTGCGGCCGCAAGCTTGCCCGCTTGGCGCGC
3	DsbA-SaPLD-2F	AACTTTAAGAAGGAGATATACATATGAAAAAAATTTGGCTGGC GCTGGCGGGCCTGGTGCTGGCGTTTAGC
4	DsbA-SaPLD-2R	CTCGAGTGCGGCCGC
5	pBAD-IF	GGCTAACAGGAGGAATTAACCATGAAAAAAACCGCGATTGCG ATTGCGG
6	pBAD-IR	TGAGTTTTTGTCTAGACCGCCCGCTTGGCGCGC
7	pBAD-VF	GGTCTAGAACAAAACTCATCTCAGAAG
8	pBAD-VR	GGTTAATTCCTCCTGTTAGCC
9	mrcB-up-F	CAAAGGCAAAGGGCGTAAG
10	mrcB-up-R	GATCACCCACCACCAGCAAATCCGGGAAACC
11	mrcB-down-F	CGGATTTGCTGGTGGTGGGGTGATCTCTCCTCAG
12	mrcB-down-R	GGCACGTTTCATCGAACGGGTC
13	dacB-up-F	CTCTTGAATATTCTGATAGGGCAAGTC
14	dacB-up-R	GCCTGTTTAACTCATTTCAGCGTCGTGGTAAAACG
15	dacB-down-F	CGACGCTTGAAATGAGTTAAACAGGCGGGTATCAC
16	dacB-down-R	CCAGGTTATATACCCCCTGCAAC

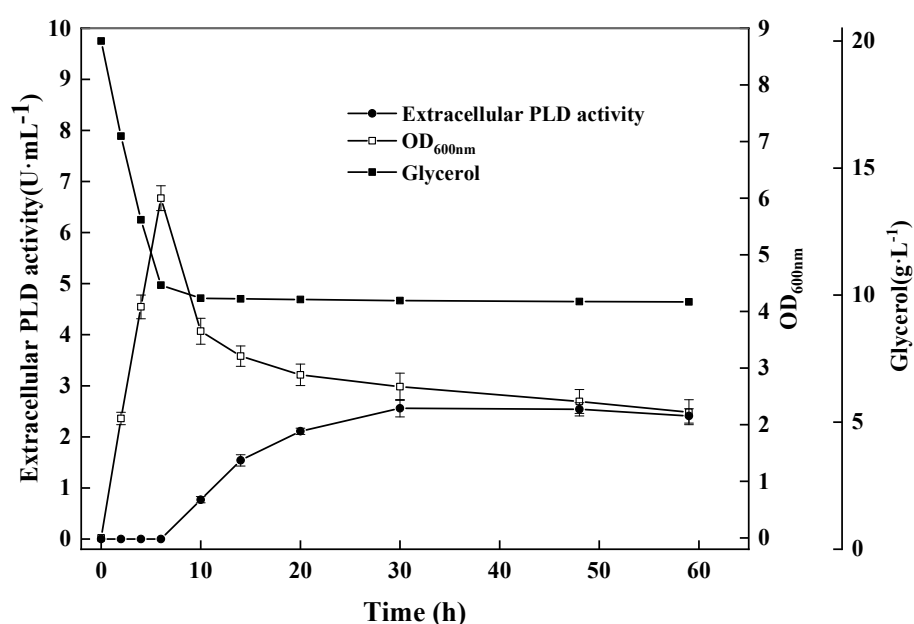


Figure S1. PLD production, cell growth and glycerol consumption by the engineered ECPLD4 after knockout of mrcB and dacB.

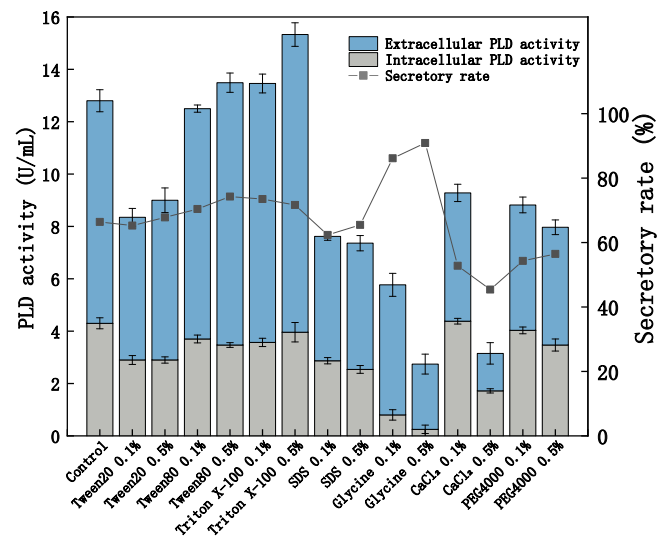


Figure S2. Comparison of the effects of different surfactants with 0.1% and 0.5% addition level on PLD production.