Metabolic Engineering of Escherichia Coli for Ectoine Production with the Fermentation Strategy of Supplementing Amino Donor

Hao Zhang  
Nanjing Tech University

Ming Zhao  
Nanjing Tech University

Yingsheng Dong  
Nanjing Tech University

Yingying Cao  
Nanjing Tech University

Yanqin Ma  
Nanjing Tech University

Zhengshan Luo  
Nanjing Tech University

Sha Li (lisha@njtech.edu.cn)  
Nanjing Tech University  https://orcid.org/0000-0001-8256-1263

Hong Xu  
Nanjing Tech University

Research

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Abstract

Background: Ectoine, a compatible solute, has broad application prospects in food biotechnology, agriculture, medicine, and cosmetics because of its protective action on biological compounds. Industrially, ectoine is produced by halophilic bacteria in a complex process. Recently, various works focus on improving ectoine production by using engineered strains, but there are still problems of low yield and low ectoine production efficiency.

Results: To overcome the drawback, a final metabolic engineered strain *E. coli* ET08 was constructed by eliminating lysine synthesis branch and by-product metabolic pathways, and ectoine production reached 10.2 g/L through culture medium optimization. Compared with nitrate, addition of ammonium salt contributed more to the ectoine synthesis. Furthermore, the ammonium sulphate boosted more ectoine titers than ammonium chloride and sodium glutamate. The analysis of transcriptional levels revealed that the ammonium sulfate enhanced ectoine biosynthesis by enhancing metabolic flux toward ectoine biosynthesis and providing affluent synthetic precursors. Ultimately, the ectoine production and yield of the *E. coli* ET08 reached 36.5 g/L and 0.3 g/g glucose with supplementing amino donor in a 7.5 L bioreactor.

Conclusions: a novel potential metabolic engineered *Escherichia coli* for ectoine production was constructed. optimizing amino donor and analyzing the transcription levels conclude that ammonium sulfate, as the optimal amino donor, has a positive effect on ectoine synthesis. It is the first report about the effect of exogenous amino donor on ectoine fermentation by metabolic engineered strain. The maximum ectoine production and yield from glucose synthesized by *E. coli* were obtained by two-stage feeding fermentation with supplementing amino donor. It provides a novel strategy for the synthesis of ectoine by engineered strain in industry. This research provides the basis for an effective process for ectoine production, together with the further applications of ectoine in food and cosmetics, and could also be used to produce other high value amino acid derivative.

Background

As a compatible solute, ectoine (1,4,5,6-tetrahydro-2-methyl-4-pyrimidinecarboxylicacid) was commonly found in halophilic and halotolerant microorganisms. It was first discovered in *Ectothiorhodospira halochloris* by Galinski et al. [1] and its structure was identified to be a cyclic amino acid derivative. In addition to its primary function of maintaining cell osmotic balance and resisting the impact of high osmotic pressure, an increasing attention was focused on its remarkable ability in protecting biological compounds [2, 3]. Therefore the potential applications of ectoine lies in food biotechnology, agriculture, skin caring, and medical fields [4–8].

As shown in Figure. 1, ectoine is synthesized from the precursor L-aspartate-β-semialdehyde (ASA), a central hub in microbial amino acid production, by three successive enzymatic steps that are catalyzed by the L-2,4-diaminobutyrate transaminase (EctB), the 2,4-diaminobutyrate acetyltransferase (EctA) and the ectoine synthase (EctC) [9–12]. The ectoine biosynthetic genes are normally organized in an operon (*ectABC*) that might also comprise the *ectD* gene [13]. Similar gene clusters involved in ectoine biosynthesis were disclosed in strain *Marinococcus halophilus* [14], *Halobacillus dabanensis* D-8T [15], *Methylmicrobium alcaliphilum* 20Z [16], *Nesterenkonia halobia* DSM20541 [17] and so on.

Commercially, ectoine production is realized through fermentation of halophiles in a special and complex process called “bacterial milking” [18, 19]. Although this method can be used to obtain ectoine in large scale, the use of large amounts of salt will corrode the equipment, which requires higher corrosion resistance of the fermentation equipment. The ability of bacteria to resist osmotic pressure shock needs to be higher in the process. To address the
shortcomings of this process, great efforts have been made, such as optimizing process conditions, improving ectoine production performance by breeding of halophilic bacteria. However, using transgenic nonhalophilic bacteria for ectoine production is more efficient in recent years.

Therefore, Becker et al. successfully integrated the ectABCD gene operon of Pseudomonas stutzeri A1501 into Corynebacterium glutamicum based on systematic metabolic engineering, and mutated aspartate kinase to ensure sufficient supply of ASA, and the final overall spacetime yield achieved 6.7 g/L ectoine per day [20]. In 2015, ectABC gene cluster of Halomonas elongata DSM 2581 was over expressed in E. coli K-12/BW25113, and the titer of ectoine reached 25.1 g/L, with a productive yield of 4048 mg/g DCW by a whole cell biocatalytic process using aspartate and glycerol as substrates [21]. Subsequently, an engineered strain E. coli ECT05 was constructed through a series of metabolic engineering strategies, besides the final titer reached 25.1 g/L, and overall ectoine yield was 0.11 g/g of glucose [22]. Recently, production of ectoine was up to 65 g/L within 56 h by transcriptional balancing of the ectoine pathway in Corynebacterium. glutamicum [23].

Although the ectoine production can be realized by fermentation of engineered strains, there are still problems of low glucose conversion rate and low ectoine production efficiency. The recent reports mainly focus on metabolic modification of engineered strains. There were few studies on nutritional requirements optimization and fermentation regulation of engineered strains. Particularly, nitrogen is a constituent of cellular components such as proteins, nucleic acids and several cofactors. It also regulates primary and secondary metabolism in different bacteria. Xu et al. increased the yield of ε-poly-L-lysine through nitrogen source regulation and optimization [24]. It was also reported that nitrogen source regulation has great effect on ethanol and antibiotic production [25, 26]. Ectoine is synthesized from aspartate, which acts as the direct precursor. The supply of amounts of nitrogen is essential for ectoine synthesis.

In this study, we constructed a metabolically engineered strain E. coli ET08 capable of producing ectoine efficiently. Further studies focus on the effects of complex nitrogen sources and amino donors on ectoine production. Then the transcription levels of the key genes in ectoine and ammonium metabolic pathways were analyzed for description the function of the amino donor. Finally, ectoine production of engineered E. coli ET08 was evaluated by two-stage feeding fermentation with supplementing amino donor. This work provides a novel strategy for the synthesis of ectoine by engineered strain in industry.

**Results And Discussion**

**Construction of engineered E. coli for ectoine production**

For heterologous synthesis of ectoine in E. coli, the basic strains E. coli MG1655 and E. coli BL21 (DE3) were chosen as hosts in light of their clear genetic background. The constructed plasmids pTrc-ectABC and pET- ectABC were transferred into E. coli MG1655 and E. coli BL21 (DE3), to form the strain ET01 and ET02 respectively (Table 1). The extracellular ectoine titer reached 0.52 g/L after 48 h cultivation with ET01 (Table 2). Although the protein expression levels of ectABC in ET02 were more powerful than those of ectABC in ET01 (see Additional file 1: Figure. S1), and the biomass of ET02 was higher than that of ET01, the ectoine titer of the strain ET02 could not be detected (Table 2). The results clearly indicated that ET01 performed best and thus was chosen for further genetic manipulation.
### Table 1

| Strains/plasmid | Relevant characteristic | Source |
|-----------------|-------------------------|--------|
| **Strains**     |                         |        |
| *E. coli* MG1655| Wild type               | This lab |
| *E. coli* BL21 (DE3) | Expression host     | This lab |
| *Halomonas venusta* | Wild type            | This lab |
| ET01            | MG1655 (pTrc-ectABC)   | This study |
| ET02            | BL21 (pET-ectABC)      | This study |
| ET03            | ET01 ΔlysA             | This study |
| ET04            | ET01 ΔthrA             | This study |
| ET05            | ET03 ΔthrA             | This study |
| ET06            | ET03 Δpta              | This study |
| ET07            | ET03 ΔldhA             | This study |
| ET08            | ET03 ΔpykF             | This study |
| **Plasmid**     |                         |        |
| pTrc99a         | trc promoter, cloning vector, Amp\(^r\) | This lab |
| pET-28a         | T7 promoter, cloning vector, Kan\(^r\) | This lab |
| pKD46           | Temperature sensitive vector carrying Red recombinase, Amp\(^r\) | This lab |
| pKD3            | Template vector, Cm\(^r\) | This lab |
| pCP20           | Temperature sensitive vector carrying FLP recombinase, Amp\(^r\) | This lab |
| pTrc-ectABC     | pTrc99a containing *Halomonas venus* ectABC gene | This study |
| pET-ectABC      | pET28a containing *Halomonas venus* ectABC gene | This study |

### Table 2

**Comparison of fermentation parameters of different strains in shake flask cultivations.**

| Strain | Ectoine titer (g/L) | Glucose consumption (g/L) | DCW (g/L) | Ectoine yield on glucose (g/g) |
|--------|---------------------|---------------------------|-----------|-------------------------------|
| ET01   | 0.52 ± 0.01         | 65 ± 1.37                 | 16.2 ± 0.39 | 0.01                          |
| ET02   | 0                   | 65 ± 1.72                 | 18 ± 0.41  | 0                             |
| ET06   | 1.09 ± 0.21         | 65 ± 1.43                 | 14.3 ± 0.29 | 0.02                          |
| ET07   | 0.6 ± 0.07          | 45 ± 0.91                 | 14.3 ± 0.32 | 0.01                          |
| ET08   | 1.87 ± 0.18         | 63 ± 1.8                  | 15.02 ± 0.48 | 0.03                          |
Elimination of branch and by-product metabolic pathways

L-aspartate-β-semialdehyde is the important precursor of L-threonine, L-lysine and ectoine synthesis pathway in *E. coli* [12]. To reduce L-aspartate-β-semialdehyde shunting by the branch metabolism, the genes of *lysA* (encoding diaminopimelate decarboxylase) and *thrA* (encoding aspartate kinase/homoserine dehydrogenase) in *E. coli* ET01 were knocked out individually and together, generating the strains *E. coli* ET03, *E. coli* ET04 and *E. coli* ET05 respectively. As shown in Figure 2, in comparison to that of *E. coli* ET01, the ectoine titer of *E. coli* ET03 (1.08 g/L) increased by 1.08 folds but biomass (13.9 g/L) decreased at the same glucose consumption (65 g/L). When *thrA* was deleted, ectoine titer of *E. coli* ET04 decreased to 0.34 g/L while glucose consumption and cell growth showed the similar trend, declining to 58 g/L and 8.2 g/L respectively. Similarly, as *lysA* and *thrA* were knocked out together, ectoine titer of *E. coli* ET05 decreased to a lower level (0.2 g/L) with 53 g/L glucose consumption and 7.05 g/L DCW. The result indicated that the deletion of *thrA* greatly inhibited the growth of *E. coli* ET01 and had negative impact on the ectoine synthesis. This might be due to the fact that *thrA* is involved in the synthesis of many important amino acids such as homoserine, methionine and threonine which are necessary for the cell growth. The deletion of *thrA* inhibited the synthesis of amino acids, and consequently the biomass decreased notably. Finally, *E. coli* ET03 was chosen for further study.

In addition to ectoine, some quantitative organic acids such as acetate, pyruvate and lactate were thought to be in the substrate. The productions of organic acids distribute the carbon flux and could significantly inhibit the cell growth by reducing the pH level, thereby affecting the titer. To further increase the ectoine production, *E. coli* ET06, *E. coli* ET07, *E. coli* ET08 were constructed by deletion of *pta*, *ldhA*, and *pykF* in *E. coli* ET03 respectively, which are the genes encoding key enzyme in acetate, lactate and pyruvate synthetic pathway respectively (Table 1). When *pta* was deleted, the ectoine production and biomass of *E. coli* ET06 were nearly equal to those of *E. coli* ET03, but the lag phase of *E. coli* ET06 was extended. However, the ectoine titer of *E. coli* ET08 increased by 73.1% (reached to 1.87 g/L) compared with that of *E. coli* ET03. Meanwhile the titer of *E. coli* ET07 was 0.6 g/L with the lowest glucose consumption (45 g/L). These results indicated that the deletion of *pta* and *ldhA* had a negative impact on the production and biomass, while knocking out *pykF* could increase the ectoine titer. This might be due to the positive role that the weakening of pyruvate pathway played in the accumulation of phosphoenolpyruvate, a prerequisite for ectoine synthesis. *E. coli* ET08 was chosen as the potential strain for further study on optimization of fermentation.

Optimization of nitrogen sources for ectoine production

Nitrogen is a constituent of cellular components such as proteins, nucleic acids and several cofactors [27]. It also regulates primary and secondary metabolism in different bacteria. To further improve the ectoine production, the effects of nitrogen sources combined with yeast extract and inorganic nitrogen sources on ectoine titer were investigated. As shown in Table 3, with the increase of yeast extract concentration in the medium, the ectoine production of *E. coli* ET08 presented an increasing trend except for the titer obtained by sodium nitrate addition. The noticeable ectoine yield (7.3 g/L) was achieved at the combination of 20 g/L yeast extract and 152 mM ammonium chloride as inorganic nitrogen source. And the ectoine yield obtained by ammonium chloride addition was 3.9 times higher than that of the control. At the same yeast extract concentration, the ectoine titer gained by ammonium chloride addition was significantly higher than that of the blank control and that obtained by sodium nitrate addition. The combination of yeast extract and nitrate had no positive effect on the production of ectoine. Both the biomass and sugar consumption capacity of *E. coli* ET08 were also reduced to a certain extent (Table 3). This implied that ammonium salt played a more significant role in promoting the production of ectoine during fermentation compared with nitrate salt. The former was more conducive to microbial absorption in comparison.
with the latter and acted as exogenous amino donor to provide \( \text{NH}_4^+ \) involved in the synthesis of glutamate directly (Figure. 1). Besides, glutamate, as a co-substrate, was involved in the catalytic reaction of the key enzyme EctB in the synthesis pathway of ectoine, and it provided an amidogen to L-aspartate-\( \beta \)-semialdehyde [12].

| Yeast extract (g/L) | 10 | 15 | 20 |
|---------------------|----|----|----|
| Inorganic nitrogen   |    |    |    |
| CK                  |    |    |    |
| \( \text{NH}_4\text{Cl} \) | 0.37 ± 0.03 | 1.35 ± 0.07 | 0.47 ± 0.04 |
| \( \text{NaNO}_3 \)   | 1.26 ± 0.11 | 4.19 ± 0.18 | 0.23 ± 0.05 |
| Ectoine titer (g/L)  |    |    |    |
|                     | 0.87 ± 0.18 | 7.3 ± 0.18 | 0.21 ± 0.02 |
| Glucose consumption (g/L) | 59 ± 1.23 | 59 ± 1.30 | 41 ± 0.47 |
|                     | 64 ± 1.63 | 60.6 ± 0.95 | 46 ± 0.94 |
|                     | 63 ± 1.8 | 61.5 ± 1.6 | 47 ± 1.27 |
| DCW (g/L)           |    |    |    |
|                     | 10.97 ± 0.5 | 12.06 ± 0.36 | 9.27 ± 0.39 |
|                     | 12.79 ± 0.43 | 13.06 ± 0.34 | 11.03 ± 0.47 |
|                     | 15.02 ± 0.48 | 14.8 ± 0.34 | 11.60 ± 0.42 |

### Optimization of exogenous amino donor

From the above results, we consider that ammonium chloride as an amino donor plays an important role in improving the ectoine production. Therefore, the types and concentrations of amino donors should be optimized. As shown in Figure. 3, the maximal ectoine output (10.2 g/L) was obtained by ammonium sulfate addition, higher than that obtained by ammonium chloride addition (7.3 g/L) and sodium glutamate addition (6.2 g/L). With the concentration of amino donor increasing, the titer presented a trend that changed from increase to decrease. And the maximum titer was obtained when the concentration of \( \text{NH}_4^+ \) (for ammonium salt)/\( \text{NH}_3^+ \) (for sodium glutamate) was 152 mM, no matter it is under the condition of ammonium sulfate (Figure. 3A), ammonium chloride (Figure. 3B) or sodium glutamate addition (Figure. 3C). The concentration of \( \text{NH}_4^+ / \text{NH}_3^+ \) had little effect on the growth rate of bacteria when the concentration of \( \text{NH}_4^+ / \text{NH}_3^+ \) was less than or equal to 152 mM. However, with the concentration of \( \text{NH}_4^+ / \text{NH}_3^+ \) reaching to 228 mM, the growth of bacteria was significantly inhibited, which was viewed as an important reason for the decrease of total glucose consumption and ectoine production. Although all these three amino donors improved ectoine biosynthesis, ammonium sulfate was the best while sodium glutamate was the worst. This might be attributed to the higher transport efficiency of ammonium sulfate compared with sodium glutamate. Overall, ammonium sulfate (76 mM) was chosen to act as final exogenous amino donor after optimization.

### Effect of \( \text{NH}_4^+ \) on the relevant metabolic pathways

Results above indicated the supplement of \( \text{NH}_4^+ \) was of great importance for improving ectoine production. We anticipated the participation of ammonium sulfate made a difference to the key genes in ammonium metabolic pathways and ectoine synthesis pathway. Thus, the transcription levels of relative genes were investigated under addition of 76 mM ammonium sulfate or not (control).

Ammonium is utilized as the main nitrogen source via two stages of uptake and assimilation. After ammonium is transported into the cells, assimilation proceeds via glutamate dehydrogenase (GDH) encoded by \( \text{gdhA} \) or glutamine
synthetase/glutamate synthase (GS/GOGAT) encoded by glnA and gltB separately, depending on the ammonium availability in the medium [28, 29]. In the study, the transcription levels of gdhA and gltB increased by 3.36 and 2.92 times respectively, whereas the transcription levels of glnA decreased significantly compared with those of the control (Figure 4). The up-regulated gene gltB promoted the synthesis of glutamate in strains over the process of fermentation. The high expression of gdhA could also enhance the accumulation of glutamate [30]. And the down-regulated gene glnA indicated the inhibition of glutamine synthesis but the promotion of glutamate accumulation. Furthermore, the glutamate as an essential substance could be directly involved in ectoine synthesis through transamination (Figure 1). Therefore, the ammonium sulfate addition could act as boost in ectoine synthesis by improving glutamate synthesis.

Aspartokinase (AK), EctA, EctB as well as EctC played significant roles in the pathway of ectoine synthesis. As depicted in Figure 4, the transcription levels of the aspartate kinase gene (lysC) was up regulated by 2.05 folds, which demonstrated that the ability of aspartate metabolism to produce downstream products was improved. Obvious increases in the transcription levels of ectABC acted as a main gene cluster in ectoine synthesis could also be seen. the transcription levels of ectB was up-regulated by 6.84 folds, higher than ectA (3.93 folds) and ectC (1.78 folds). This result indicated that EctB was the most important enzyme among the key enzyme of the ectoine synthesis, which was consistent with the results previously reported [23]. It was speculated that the ammonium sulfate addition enhanced the supplement of the precursor L-aspartate-β-semialdehyde and glutamate, providing affluent co-substrate for the key enzyme EctB. At the same time, the strong expression of ectABC, especially ectB, promoted the flow of substrate to the ectoine synthesis pathway, improving the ectoine production.

**Fermentation performance in a 7.5L bioreactor**

To assess the overall production performance of *E. coli* ET08 under the addition of exogenous ammonium sulfate (76 mM), ectoine fermentation in a 7.5 L bioreactor with feeding batch fermentation using two-stage feed was conducted. As shown in Figure 5, the ectoine concentration reached the maximum (36.5 g/L) at 36 h. At the first phase of fermentation (0 h ~ 18 h), the bacteria grew fast, but just 9.4 g/L extracellular ectoine titer could be seen. At the second phase of fermentation (18 h ~ 36 h), the ectoine titer accumulated rapidly even though there was a slow growth in dry cell weight, reaching 27 g/L at about 30 h and remaining until the end of the fermentation. The specific ectoine production and productivity reached 1.4 g/g DCW and 1.01 g/L/h respectively. Plus, the yield of ectoine was 0.3 g/g glucose. To our best knowledge, our work shows the highest ectoine titer and yield from glucose synthesized by *E. coli*, and still has room for further improvement. Compared with previous studies (Table 4), whole-cell catalysis seems to have certain advantages in yield, but it requires the addition of extra glycerol and aspartate, which results in high cost of extra substrate, cell culture before catalytic reaction. Although *C. glutamicum* ectABC<sup>opt</sup> [23] achieved higher titer, the yield was only 0.19 g/g, which was lower than our yield. Thus, the metabolic engineered strain *E. coli* ET08 and the fermentation strategy of supplementing amino donor shows a promising value for industrial production of ectoine.
Table 4

Microbial production of ectoine using different fermentative strains or biocatalysts.

| Strain                                | Titer (g/L) | Specific production (g/g DCW) | Yield (g/g) | Productivity (g/L/h) | Process strategy            | Reference |
|---------------------------------------|-------------|-------------------------------|-------------|-----------------------|-----------------------------|-----------|
| E. coli ET08                          | 36.5        | 1.4                           | 0.30        | 1.01                  | Fed-batch                   | This work |
| E. coli Ect05                          | 25.1        | 0.8                           | 0.11        | 0.84                  | Fed-batch                   | [22]      |
| E. coli BW25113 (pBAD-ectABC)         | 25.1        | 4.1                           | -           | 1.04 $^c$             | Whole-cell catalysis $^a$    | [21]      |
| E. coli ECT2                           | 12.7        | -                             | 1.27        | 0.53 $^c$             | Whole-cell catalysis $^a$    | [34]      |
| C. glutamicum ectABC$^{opt}$          | 65.3        | -                             | 0.19        | 1.16                  | Fed-batch                   | [23]      |
| Chromohalobacter salexigens DSM3043   | 32.9        | 0.5                           | -           | 1.35                  | continuous reactors with cell $^b$ | [19]      |

$^a$ Whole-cell catalysis using aspartate and glycerol as substrates at a high cell density.

$^b$ A special fermentation process using two continuously operated bioreactors.

$^c$ Achieved by calculating reported data.

Conclusions

In this study, we constructed an efficient strain *E. coli* ET08 by metabolic engineering. By optimizing amino donor and analyzing the transcription levels, we could conclude that ammonium sulfate, as the optimal amino donor, has a positive effect on ectoine synthesis. Batch fermentation of the *E. coli* ET08 with supplementing ammonium sulfate led to an accumulation of 36.5 g/L ectoine, and the specific ectoine production, yield and productivity reached 1.4 g/g DCW, 0.3 g/g glucose and 1.01 g/L/h respectively. To the best of our knowledge, this is the first report about the effect of exogenous amino donor on ectoine fermentation by metabolic engineered strain. It provides a novel strategy for the synthesis of ectoine by engineered strain in industry. This research provides the basis for an effective process for ectoine production and could also be used to produce other high value amino acid derivative.

Methods

Strains, plasmids and culture medium

The strains and plasmids constructed and used in this study are listed in Table 1. All restriction digest enzymes, DNA polymerase and DNA ligase were obtained from Takara Bio Inc. (Dalian, China). Oligonucleotides were synthesized by GENEWIZ Bio Inc. (Suzhou, China). Ectoine was purchased from Sigma-Aldrich (Shanghai, China). The seed culture medium was the Luria–Bertani (LB) medium. The fermentation culture medium in shake flasks contains (per liter) 5 g glucose, 20 g yeast extract, 5 g glycerol, 0.5 g NaCl, 2.5 g KH$_2$PO$_4$, 7.5 g K$_2$HPO$_4$, 1 g MgSO$_4$·7H$_2$O, 15 g magnesium carbonate hydroxide pentahydrate. The fed-batch fermentation culture medium in a
7.5 L bioreactor contained (per liter) 5 g glucose, 20 g yeast extract, 5 g glycerol, 0.5 g NaCl, 2.5 g KH₂PO₄, 7.5 g K₂HPO₄, 1 g MgSO₄·7 H₂O, 10 g ammonium sulfate, 15 g magnesium carbonate hydroxide pentahydrate.

Construction of plasmids and gene knockout

The ECTABC gene (ectABC) was amplified from the genome of *Halomonas venusta* using the primers ET01-F and ET01-R. The PCR products were purified and then ligated into the linearized vector pTrc99a by in-fusion cloning with restriction sites of *Kpn*I and *BamH*I, forming plasmid pTrc-ectABC. For the construction of the plasmid pET-ectABC, the ectABC was amplified by PCR with ET02-F and ET02-R using the genomic DNA of *H. venusta* as a template. The PCR products were purified and then inserted into pET-28a by in-fusion cloning with restriction sites of *BamH*I and *Xho*I. Deletion of the lysA, thrA, pykF, ldhA and pta genes were carried out using the Red recombination method [31]. The primers used for gene cloning and chromosomal manipulation are listed in Additional file 1: Table S1.

Culture conditions in shake flasks

Seed cultures were prepared by transferring an appropriate amount of agar slant cultured cells into a 500 mL Erlenmeyer flask containing 50 mL medium and culturing at 37 °C with shaking at 200 rpm for 8 h. The ectoine production of engineered strains was investigated by fed-batch culture, with a 10% inoculum size in a 500 mL baffled shake flask containing 50 mL medium, at 37 °C with shaking at 200 rpm for 48 h. Both media were supplemented with 100 mg/L ampicillin. As the inducer, 0.1 mmol/L IPTG was added four hours after the fermentation began. 50% glucose was supplied at a final concentration of 20 g/L every 12 h.

Nitrogen sources were optimized by mixing various concentrations of yeast extract (10 g/L, 15 g/L, 20 g/L) up with ammonium chloride (152 mM) and sodium nitrate (152 mM) respectively. And the medium without inorganic nitrogen addition was used as control check (CK). To optimize the exogenous amino donor, the effects of various concentrations of ammonium chloride (0, 38, 76, 152, 228 mM), ammonium sulfate (0, 19, 38, 76 114 mM) and sodium glutamate (0, 38, 76, 152, 228 mM) were investigated in shake-flask tests.

Two-stage fed-batch fermentation in a 7.5 L bioreactor

Primary seed cultures were prepared by transferring an appropriate amount of agar slant cultured cells into a 30 mL Screw glass sample bottle containing 5 mL medium and culturing at 37 °C with orbital shaking at 200 rpm for about 10 h. The bacterial solution was inoculated into a 1 L shake flask containing 200 mL of seed medium, inoculated in an amount of 1%, and cultured at 37 °C for 10 h at 200 rpm to prepare a secondary seed culture. The ectoine production of engineered strains was investigated by fed-batch culture, with a 15% (v/v) inoculum size in a 7.5 L bioreactor (New Brunswick BioFlo/CelliGen 115, New Brunswick, Germany) containing 3.5 L medium. The pH was kept constant at 7.0 by magnesium carbonate hydroxide pentahydrate, and the temperature was maintained at 37 °C. The aeration rate and agitation speed were 1.5vvm and 600 rpm, respectively. The glucose concentration constantly fluctuated during 0 h ~ 24 h, and then it reached almost zero after 24 h. As the inducer, 0.1 mmol/L IPTG was added when the optical density at 600 nm (OD₆₀₀) reached 0.4.

RNA sample preparation and RTqPCR analysis

Total RNA was isolated using RNAiso Plus (9108Q, TaKaRa Biotechnology Company, China). The synthesis of cDNA was then performed with PrimeScript™ II 1st Strand cDNA Synthesis Kit (6210A, TaKaRa Biotechnology Company, China), using the total RNA as template. Real-time PCR (StepOnePlus™ Real-Time PCR System, Applied Biosystems, USA) was carried out with SYBR® Premix Ex TaqTM (RR420Q, TaKaRa Biotechnology Company, China) as fluorochrome, with 16S rDNA as an endogenous control gene. The primers used for qPCR were designed.
according to the *E. coli* genome sequence and were summarized in Additional file 1: Table S2. The amplification program consisted of one cycle at 95°C for 30 s, followed by 40 cycles at 95°C for 5 s, 60°C for 30 s. All reactions were repeated three folds. Data from RT-qPCR were treated with the $2^{-\Delta\Delta C_{t}}$ method for relative quantification [32]. To present the results in a better way, the formula $2^{-\Delta\Delta C_{t}}$ was multiplied by 1. Therefore, the comparative expression level of each gene under the control was always 1 [33].

**Analytic methods**

Cell growth was monitored by measuring the absorbance at 600 nm (OD$_{600}$) that was then converted to dry cell weight (DCW) by a calibration curve. The concentration of extracellular ectoine was determined via high-performance liquid chromatography (HPLC) using TSKgel ODS-80Ts column (4.6 × 250 mm, Tosoh, Tokyo) with an acetonitrile/water mixture (2:98 v/v) at a flow rate of 0.5 mL/min as the mobile phase. Ectoine was monitored by a UV detector at a wavelength of 210 nm. The residual sugar in the fermentation broth was measured using an SBA-40C biological analyzer (Shandong Academy of Sciences, China).

**Declarations**

**Ethics approval and consent to participate**

Not applicable

**Consent for publication**

Not applicable

**Availability of data and materials**

All data generated or analyzed during this study are included in this published article (and its Additional file 1).

**Competing interests**

The authors declare that they have no competing interests

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**Authors' contributions**

ZH performed the experiments and drafted the manuscript. LS, ZH and LZS designed this study. ZM, DYS, CYY and MYQ supervised the experiments and helped to draft the manuscript. LS and XH conceived the study and reviewed the final manuscript. All authors read and approved the final manuscript.

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35. Declarations.
Related metabolic pathways for the synthesis of ectoine in engineered E. coli. The genes marked in green indicate the key genes of biosynthetic ectoine introduced into the engineered E. coli. The genes marked in red indicate deletion of the corresponding gene. Abbreviations: PEP, phosphoenolpyruvate.
Figure 2

The effects of lysA and thrA deletion on ectoine fermentation.
Figure 3

Effect of different amino donors on ectoine fermentation. A: Ectoine titer, DCW and glucose consumption of ET08 with ammonium sulfate (0, 19, 38, 76 and 114 mM) addition; B: Ectoine titer, DCW and glucose consumption of ET08 with ammonium chloride (0, 38, 76, 152, 228 mM) addition; C: Ectoine titer, DCW and glucose consumption of ET08 with sodium glutamate (0, 38, 76, 152, 228 mM) addition.
Figure 4

Transcription levels of the key genes in ammonium metabolic pathways and ectoine synthesis. The level of transcription was calculated relative to transcription of the control (0 mM ammonium sulfate), which were defined as 1. The error bars indicate the standard deviation of three samples taken from the same RNA sample.
Figure 5
Fed-batch fermentation of ET08 in a 7.5 L bioreactor with the supplement of ammonium sulfate.

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