

**Distinct physiological roles for the two L-asparaginase isozymes of  
*Escherichia coli***

Author

Srikhanta, Yogitha N, Atack, John M, Beacham, Ifor R, Jennings, Michael P

Published

2013

Journal Title

Biochemical and Biophysical Research Communications

DOI

[10.1016/j.bbrc.2013.05.066](https://doi.org/10.1016/j.bbrc.2013.05.066)

Rights statement

© 2013 Elsevier. This is the author-manuscript version of this paper. Reproduced in accordance with the copyright policy of the publisher. Please refer to the journal's website for access to the definitive, published version.

Downloaded from

<http://hdl.handle.net/10072/55540>

Griffith Research Online

<https://research-repository.griffith.edu.au>

**Distinct physiological roles for the two L-asparaginase isozymes of *Escherichia coli*.**

Yogitha N. Srikhanta<sup>b</sup>, John M. Atack<sup>a</sup>, Ifor R. Beacham<sup>a</sup>, and Michael P. Jennings<sup>a\*</sup>,

<sup>a</sup>Institute for Glycomics, Griffith University, Gold Coast, QLD 4222, Australia.

<sup>b</sup>Department of Microbiology and Immunology, The University of Melbourne, Victoria 3010, Australia.

\*Corresponding author. *E-mail address*: m.jennings@griffith.edu.au

## **Abstract**

*Escherichia coli* expresses two L-asparaginase (EC 3.5.1.1) isozymes: L-asparaginase I, which is a low affinity, cytoplasmic enzyme that is expressed constitutively, and L-asparaginase II, a high affinity periplasmic enzyme that is under complex co-transcriptional regulation by both Fnr and Crp. The distinct localisation and regulation of these enzymes suggest different roles. To define these roles, a set of isogenic mutants was constructed that lacked either or both enzymes. Evidence is provided that L-asparaginase II, in contrast to L-asparaginase I, can be used in the provision of an anaerobic electron acceptor when using a non-fermentable carbon source in the presence of excess nitrogen.

## 1. Introduction

L-asparaginases are widely distributed in the three domains of life and play a central role in amino acid metabolism and utilization. L-asparaginase enzymes catalyse the hydrolysis of L-asparagine (EC 3.5.1.1) generating L-aspartate and ammonia ( $\text{NH}_3$ ). Based on studies with *Escherichia coli* and *Salmonella enterica*, bacterial L-asparaginases are of two types: a high affinity enzyme located in the periplasm (L-asparaginase II encoded by *ansB*) and a cytoplasmic low affinity enzyme (L-asparaginase I encoded by *ansA*). *E. coli* and a few other bacteria have been shown to possess both type I and type II isozymes [1,2,3,4]. The amino acid sequences are substantially diverged but retain sequence similarity around the active site residues [4]. The high affinity enzymes have attracted a great deal of interest due to their utility as anti-tumour agents in the treatment of childhood acute lymphocytic leukemia [5]. This is due to the efficient depletion of exogenous L-asparagine, which leukemic cells depend on for growth. More recently it has been reported in *Salmonella enterica* Typhimurium that L-asparaginase II (AnsB) has a role in infection and immunity by inhibiting T-cell responses [6] and it also contributes to virulence in the human pathogens *Campylobacter jejuni*, *Helicobacter pylori* and *S. enterica* Typhimurium [6,7,8] via a role in colonisation.

As well as two L-asparaginase isozymes, *E. coli* also possess two distinct systems for L-asparagine uptake distinguishable on the basis of specificity and regulation [9,10]: a low-affinity system ( $K_m = 80\mu\text{M}$ ); and a high affinity system ( $K_m = 3.5\mu\text{M}$ ) which is repressed by the presence of L-asparagine in the growth medium at concentrations greater than 1mM. These systems have not been studied at the molecular level; however, an asparagine permease, AnsP, has also been reported in *S. enterica* and *E. coli* [11,12].

There is evidence that L-asparaginase I (AnsA) plays a role in *E. coli* in the utilisation of L-asparagine as a nitrogen source [13] and it presumably functions to degrade L-asparagine when it has accumulated to an appropriate intracellular concentration. However, the role of the high affinity

L-asparaginase II has never been determined. While L-asparaginase I is expressed constitutively, L-asparaginase II expression requires co-dependent activation by both anaerobiosis, *via* the Fnr transcriptional activator, and by the cyclic-AMP receptor protein (Crp) [3,14,15]. Several possible roles are consistent with the activity, localization, and regulation of the L-asparaginase II. Firstly, L-asparaginase II may serve to utilize L-asparagine as a carbon source under unfavorable conditions, as in the case of, for example, exported phosphatases, proteases and lipases. Such a role is consistent with regulation by Crp. Secondly, this enzyme may be required for growth on low concentrations of L-asparagine as a nitrogen source. Thirdly, it has been suggested that during anaerobic growth on a non-fermentable carbon source, hydrolysis of L-asparagine could lead to the provision of fumarate as a terminal electron acceptor for anaerobic respiration since the resulting L-aspartate could be taken up by an L-aspartate transporter and catabolised to fumarate by L-aspartase (See Fig. 2) [3]. This function is consistent with the anaerobic regulation of L-asparaginase II in both *E. coli*, *via* Fnr, and *S. enterica* by an unknown mechanism, [15] and with the anaerobic regulation of aspartase and dicarboxylate transporters [16,17,18].

## 2. Materials and Methods

### 2.1 Media and Growth Conditions

Strains and plasmids used in this study are detailed in Table 1. HB94 (wild type) and its three mutant derivatives RC614 (*ansA*<sup>-</sup>); HB94K (*ansB*<sup>-</sup>); RC614K (*ansA*<sup>-</sup>, *ansB*<sup>-</sup>) were grown as follows: for Fig. 1A, HB94 was grown in 1X M9 medium and supplemented with 10 mM Ca, 100 mM Mg, 20% glycerol, 100 mM tryptophan and 18 mM NH<sub>4</sub>Cl. Cultures were grown anaerobically at 37<sup>0</sup>C. Cells were harvested by centrifugation (45,000 x g, 15 min, RT) after overnight growth, washed twice with 1X M9 medium and resuspended in 2 ml of the same medium. This was used as a starter culture (initial OD<sub>600</sub> of 0.03) to inoculate tubes containing 15ml of 1X M9, and supplemented with 10 mM Ca, 100 mM Mg and 100 mM tryptophan in the presence or absence of 0.05mM L-asparagine and 20% glycerol. For Figs. 1B and 1C, the growth rate of HB94 and the isogenic

mutants RC614, HB94K and RC614K was observed with 0.05 mM L-asparagine in the absence (1B) or presence (1C) of NH<sub>4</sub>Cl. Strains were grown as above without or with 18 mM NH<sub>4</sub>Cl. All cultures were grown anaerobically at 37<sup>0</sup>C. Where required, kanamycin was used at a concentration of 10 µg/ml. Cells were harvested by centrifugation (45,000 x g, 15 min, RT) after overnight growth, washed twice with 1X M9 medium and resuspended in 2 ml of the same medium. These were used as starter cultures (initial OD<sub>600</sub> of 0.03) to inoculate tubes containing 15 ml of 1X M9, 0.05 mM L-asparagine and supplemented with 10 mM Ca, 100 mM Mg, 20% glycerol and 100 mM tryptophan. Readings were taken in triplicate for each strain and averaged.

## 2.2 Construction of the *ansB* mutant.

An *ansB* mutant was constructed by insertion of a kanamycin resistance cassette, excised from pKIX (Pharmacia) by digestion with *Sma*I, into a unique *Hpa*I site present in the coding region of *ansB* on plasmid pMJ13. The resulting plasmid containing the *ansB::kan* allele was linearised and used to transform *E. coli* strain V355 competent cells (*recD*; [19]). Kanamycin resistant colonies were isolated and the presence of the kanamycin cassette confirmed by Southern blot. The *ansB* phenotype was confirmed by Western blot using anti-AnsB rabbit polyclonal sera [20] and absence of high affinity L-asparaginase activity in anaerobically grown cultures. One of the characterised mutants, MPJ10 was used to make a P1 lysogen, and the *ansB::kan* mutation transduced to recipient strains HB94 and RC614 [13], resulting in strains HB94kan and RC614kan.

## 3. Results and Discussion

### 3.1. *L-asparaginase II* allows anaerobic respiration in the presence of excess nitrogen.

In order to test these putative functions of the high affinity, exported L-asparaginase II, and in particular the possibility of a role in anaerobic respiration, we grew strain HB94 anaerobically with glycerol as the non-fermentable carbon source and with L-asparagine. Slow growth was observed which was dependent on both glycerol and L-asparagine (Fig.1A). To investigate the role of L-

asparaginases in the provision of L-asparagine as electron acceptor, we used isogenic mutants of *E. coli* strain HB94 which lack either L-asparaginase I (RC614; see [13]), L-asparaginase II (HB94K) or both isozymes (RC614K; see Fig. 1B, C). In medium lacking NH<sub>4</sub>Cl, all strains were able to grow slowly, indicating that either L-asparaginase I or II can be used as a source of fumarate (Fig. 1B). However, in medium containing NH<sub>4</sub>Cl, growth was dependent on the presence of L-asparaginase II (Fig. 1C), indicating that L-asparaginase I was unable to deaminate internal L-asparagine under these conditions. These results indicate that both L-asparaginase isozymes are capable of allowing use of L-asparagine as a source of fumarate as an anaerobic electron acceptor, but that in the presence of an excess of nitrogen, in the form of NH<sub>4</sub>Cl, L-asparaginase II is obligatory. This may be due to a critical L-asparagine transporter being inoperative in the presence of excess nitrogen, due either to inhibition of its synthesis or transport function: it is known that the GlnLG system (encoded by *glnLG*; also known as NtrBC) is not active under conditions of high ammonia [21], and hence expression of an L-asparagine transporter would not occur if it was part of the GlnLG/NtrBC regulon. Alternatively, the global regulator Fis has been shown to repress expression of the asparagine transporter AnsP in *E. coli* under nutrient-rich conditions, which may include directly or indirectly, excess nitrogen [12]. The AnsP protein has been shown to be critical for asparagine transport in *S. enterica* [11], but whether AnsP is the L-asparagine transporter down-regulated in *E. coli* in response to ammonia remains to be elucidated. Nonetheless, if L-asparagine is unable to be transported, the action of L-asparaginase II in the periplasm becomes essential in the generation of aspartate, and hence fumarate as an electron acceptor, from exogenous L-asparagine.

The utilisation of L-asparagine, via pathway II, in anaerobic respiration (Fig. 2), is consistently positively regulated by anaerobiosis *via* Fnr: transcription of the C4-dicarboxylate transporter and L-aspartase genes (*dcuB* and *aspA*) are both regulated by Fnr and Crp [16,17,18]. Fumarate reductase, responsible for the terminal step of anaerobic electron transfer to fumarate, is likewise Fnr-regulated [22]. In addition, pathway II in its entirety, from L-asparagine, is repressed by nitrate

via the NarXL two-component system [23,24]. Since nitrate is a preferred electron acceptor, this is consistent with the role of asparaginase II, and pathway II, in anaerobic respiration.

In addition to anaerobic regulation, it is notable that in both *S. enterica* Typhimurium and *E. coli*, production of L-asparaginase II is also absolutely dependent on Crp binding at the respective *ansB* promoters; in *E. coli* the promoter is co-dependent on Fnr and Crp whilst in *S. enterica* the upstream Fnr site has been modified to form a second Crp binding site but the promoter retains anaerobic regulation by an unknown mechanism [3,25]. The *ansB* gene is also transcriptionally up-regulated by the presence of amino acids [26]. Thus, in both enteric organisms, asparaginase II is highly expressed under stringent nutritional conditions and anaerobiosis allowing the conversion of L-asparagine to L-aspartate and anaerobic respiration. Chemotaxis in response to L-asparagine has been demonstrated in a strain expressing the aspartate receptor (Tar) and asparaginase II, the latter under aerobic conditions from an artificial promoter [27]; since the native *ansB* promoter is very highly expressed under Crp-permissive plus anaerobic conditions [3,28], it can be assumed that chemotaxis towards L-asparagine-derived L-aspartate will be similarly enabled under these native conditions, followed by uptake and conversion to fumarate.

In summary, there are two pathways by which L-asparagine may be utilised (Fig. 2). One involves the use of L-asparagine as a source of nitrogen and/or carbon and depends on active transport of L-asparagine followed by conversion of the accumulated L-asparagine to L-aspartate by the low affinity cytoplasmic L-asparaginase I. The second (Fig. 2, pathway II), in the presence of ammonia as nitrogen source, is the utilisation of low concentrations of exogenous L-asparagine as a terminal electron acceptor following conversion by L-asparaginase II to L-aspartate and thence fumarate by L-aspartase. Initial conversion to L-aspartate by the high affinity L-asparaginase II in the periplasm will allow (a) chemotaxis to L-asparagine via the product, L-aspartate, and Tar signalling; and (b) transport of L-aspartate into the cell followed by conversion to fumarate. This becomes essential when ammonia is present which we suggest leads to abolition of asparagine transport into the cytoplasm through potential down-regulation of a likely sole asparagine



transporter. Significantly, this pathway is subject to anaerobic regulation, involving Fnr, and by nitrate, *via* NarL, at the periplasmic, membrane transport and cytoplasmic levels, consistent with its role in anaerobic respiration.

The role of L-asparaginase II in pathogenesis in *S. enterica* serovar Typhimurium may well follow selection for this enzyme in pathway II in non-host environments, or as a commensal, and hence be regarded as ‘coincidental selection’ [29].

## Acknowledgements

The work was supported by NHMRC Program Grant 565526 to M.P.J.

## References

- [1] H. Cedar, J.H. Schwartz, Localization of the two-L-asparaginases in anaerobically grown *Escherichia coli*, *J Biol Chem* 242 (1967) 3753-3755.
- [2] H. Cedar, J.H. Schwartz, Production of L-asparaginase II by *Escherichia coli*, *J Bacteriol* 96 (1968) 2043-2048.
- [3] M.P. Jennings, I.R. Beacham, Co-dependent positive regulation of the *ansB* promoter of *Escherichia coli* by CRP and the FNR protein: a molecular analysis, *Mol Microbiol* 9 (1993) 155-164.
- [4] P.G. Jerlstrom, D.A. Bezjak, M.P. Jennings, I.R. Beacham, Structure and expression in *Escherichia coli* K-12 of the L-asparaginase I-encoding *ansA* gene and its flanking regions, *Gene* 78 (1989) 37-46.
- [5] J.H. Schwartz, J.Y. Reeves, J.D. Broome, Two L-asparaginases from *E. coli* and their action against tumors, *Proc Natl Acad Sci U S A* 56 (1966) 1516-1519.
- [6] A.L. Kullas, M. McClelland, H.J. Yang, J.W. Tam, A. Torres, S. Porwollik, P. Mena, J.B. McPhee, L. Bogomolnaya, H. Andrews-Polymeris, A.W. van der Velden, L-asparaginase II produced by *Salmonella typhimurium* inhibits T cell responses and mediates virulence, *Cell Host Microbe* 12 (2012) 791-798.
- [7] D. Hofreuter, V. Novik, J.E. Galan, Metabolic diversity in *Campylobacter jejuni* enhances specific tissue colonization, *Cell Host Microbe* 4 (2008) 425-433.
- [8] K. Shibayama, H. Takeuchi, J. Wachino, S. Mori, Y. Arakawa, Biochemical and pathophysiological characterization of *Helicobacter pylori* asparaginase, *Microbiol Immunol* 55 (2011) 408-417.
- [9] R.C. Willis, C.A. Woolfolk, Asparagine utilization in *Escherichia coli*, *J Bacteriol* 118 (1974) 231-241.
- [10] R.C. Willis, C.A. Woolfolk, L-asparagine uptake in *Escherichia coli*, *J Bacteriol* 123 (1975) 937-945.
- [11] M.P. Jennings, J.K. Anderson, I.R. Beacham, Cloning and molecular analysis of the *Salmonella enterica ansP* gene, encoding an L-asparagine permease, *Microbiology* 141 (1995) 141-146.

- [12] M.D. Bradley, M.B. Beach, A.P. de Koning, T.S. Pratt, R. Osuna, Effects of Fis on *Escherichia coli* gene expression during different growth stages, *Microbiology* 153 (2007) 2922-2940.
- [13] T. Del Casale, P. Sollitti, R.H. Chesney, Cytoplasmic L-asparaginase: isolation of a defective strain and mapping of *ansA*, *J Bacteriol* 154 (1983) 513-515.
- [14] R.H. Chesney, *E. coli*-asparaginase II production in the presence and absence of catabolite activating protein, *FEMS Microbiology Letters* 17 (1983) 161-162.
- [15] S. Scott, S. Busby, I. Beacham, Transcriptional co-activation at the *ansB* promoters: involvement of the activating regions of CRP and FNR when bound in tandem, *Mol Microbiol* 18 (1995) 521-531.
- [16] P. Golby, D.J. Kelly, J.R. Guest, S.C. Andrews, Transcriptional regulation and organization of the *dcuA* and *dcuB* genes, encoding homologous anaerobic C4-dicarboxylate transporters in *Escherichia coli*, *J Bacteriol* 180 (1998) 6586-6596.
- [17] P.G. Jerlstrom, Liu, J. and Beacham, I. R., Regulation of *Escherichia coli* L-asparaginase II and L-aspartase by the *fnr* gene product, *FEMS Microbiology Letters* 41 (1987) 127-130.
- [18] S.A. Woods, J.R. Guest, Differential roles of the *Escherichia coli* fumarases and *fnr*-dependent expression of fumarase B and aspartase, *FEMS Microbiology Letters* 48 (1987) 219-224.
- [19] C.B. Russell, D.S. Thaler, F.W. Dahlquist, Chromosomal transformation of *Escherichia coli recD* strains with linearized plasmids, *J Bacteriol* 171 (1989) 2609-2613.
- [20] K.J. Spring, P.G. Jerlstrom, D.M. Burns, I.R. Beacham, L-asparaginase genes in *Escherichia coli*: isolation of mutants and characterization of the *ansA* gene and its protein product, *J Bacteriol* 166 (1986) 135-142.
- [21] T.A. Blauwkamp, A.J. Ninfa, Physiological role of the GlnK signal transduction protein of *Escherichia coli*: survival of nitrogen starvation, *Mol Microbiol* 46 (2002) 203-214.
- [22] H.M. Jones, R.P. Gunsalus, Regulation of *Escherichia coli* fumarate reductase (*frdABCD*) operon expression by respiratory electron acceptors and the *fnr* gene product, *J Bacteriol* 169 (1987) 3340-3349.
- [23] E.B. Goh, P.J. Bledsoe, L.L. Chen, P. Gyaneshwar, V. Stewart, M.M. Igo, Hierarchical control of anaerobic gene expression in *Escherichia coli* K-12: the nitrate-responsive NarX-NarL regulatory system represses synthesis of the fumarate-responsive DcuS-DcuR regulatory system, *J Bacteriol* 187 (2005) 4890-4899.
- [24] C. Constantinidou, J.L. Hobman, L. Griffiths, M.D. Patel, C.W. Penn, J.A. Cole, T.W. Overton, A reassessment of the FNR regulon and transcriptomic analysis of the effects of nitrate, nitrite, NarXL, and NarQP as *Escherichia coli* K12 adapts from aerobic to anaerobic growth, *J Biol Chem* 281 (2006) 4802-4815.
- [25] M.P. Jennings, S.P. Scott, I.R. Beacham, Regulation of the *ansB* gene of *Salmonella enterica*, *Mol Microbiol* 9 (1993) 165-172.
- [26] J. Green, M.F. Anjum, J.R. Guest, Regulation of the *ndh* gene of *Escherichia coli* by integration host factor and a novel regulator, *Arr*, *Microbiology* 143 ( Pt 9) (1997) 2865-2875.
- [27] S.D. Goldberg, P. Derr, W.F. DeGrado, M. Goulian, Engineered single- and multi-cell chemotaxis pathways in *E. coli*, *Mol Syst Biol* 5 (2009) 283.
- [28] M.P. Jennings, I.R. Beacham, Analysis of the *Escherichia coli* gene encoding L-asparaginase II, *ansB*, and its regulation by cyclic AMP receptor and FNR proteins, *J Bacteriol* 172 (1990) 1491-1498.
- [29] S.P. Brown, D.M. Cornforth, N. Mideo, Evolution of virulence in opportunistic pathogens: generalism, plasticity, and control, *Trends Microbiol* 20 (2012) 336-342.

## Figure legends

**Figure 1. The role of L-asparaginase II in anaerobic respiration.** (1A) Growth of HB94 in the presence or absence of asparagine and glycerol. (1B) Growth of HB94 and the isogenic mutants RC614, HB94K and RC614K in the presence of asparagine and absence of  $\text{NH}_4\text{Cl}$ . (1C) Growth of HB94 and the isogenic mutants RC614, HB94K and RC614K in the presence of asparagine and  $\text{NH}_4\text{Cl}$ . Growth conditions are detailed in Materials and Methods.

**Figure 2. Proposed role for L-asparaginase II in asparagine utilisation.** Pathway I indicates utilization of L-asparagine in the absence of L-asparaginase II. Under conditions of both anaerobiosis and high cAMP L-asparaginase II is expressed (Pathway II) and converts L-asparagine to L-aspartate in the periplasm, followed by transport into the cytoplasm and utilisation as a terminal electron acceptor. A consequence of conversion of L-asparagine to L-aspartate in the periplasm is signal conversion with respect to Tar chemotaxis.

**Table 1.** Strains and plasmids used in this study. CGSC: Coli Genetic Stock Centre

Name	Relevant genotype	Source/Reference
<i>E.coli</i> Strains		
HB94 HB94kan	Wild type <i>ansB::kan</i>	Del Casale <i>et al.</i> , 1983 This work
RC614 RC614kan CGSC6355(P1)	<i>ansA</i> <i>ansA, ansB::kan</i> P1 lysogen	Del Casale <i>et al.</i> , 1983 This work CGSC
Plasmids		
pMJ13 pMJ13kan pPJ1	<i>ansB</i> <i>ansB::kan</i> <i>ansA</i>	Jennings <i>et al.</i> 1990 This work Jerlstrom <i>et al.</i> 1989