

The *Escherichia coli* Citrate Carrier CitT: a Member of a Novel Eubacterial Transporter Family Related to the 2-Oxoglutarate/Malate Translocator from Spinach Chloroplasts

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Under anoxic conditions in the presence of an oxidizable cosubstrate such as glucose or glycerol, *Escherichia coli* converts citrate to acetate and succinate. Two enzymes are specifically required for the fermentation of the tricarboxylic acid, i.e., a citrate uptake system and citrate lyase. Here we report that the open reading frame (designated *citT*) located at 13.90 min on the *E. coli* chromosome between *rna* and the citrate lyase genes encodes a citrate carrier. *E. coli* transformed with a plasmid expressing *citT* was capable of aerobic growth on citrate, which provides convincing evidence for a function of CitT as a citrate carrier. Transport studies with cell suspensions of the transformed strain indicated that CitT catalyzes a homologous exchange of citrate or a heterologous exchange against succinate, fumarate, or tartrate. Since succinate is the end product of citrate fermentation in *E. coli*, it is likely that CitT functions in vivo as a citrate/succinate antiporter. Analysis of the primary sequence showed that CitT (487 amino acids, 53.1 kDa) is a highly hydrophobic protein with 12 putative transmembrane helices. Sequence comparisons revealed that CitT is related to the 2-oxoglutarate/malate translocator (SODi1 gene product) from spinach chloroplasts and five bacterial gene products, none of which has yet been functionally characterized. It is suggested that the *E. coli* CitT protein is a member of a novel family of eubacterial transporters involved in the transport of di- and tricarboxylic acids.

Under oxic growth conditions, most *Escherichia coli* strains are not able to utilize citrate due to the lack of a functional transport system. This is a key characteristic of *E. coli* among enterobacteria (15). Some *E. coli* strains capable of aerobic growth on citrate possess plasmid-encoded citrate uptake systems. The citrate carrier genes from two of these plasmids were cloned and sequenced (13, 29). The deduced proteins, which consisted of 431 amino acids and differed in six positions only, exhibited 93% sequence identity to CitA from *Salmonella typhimurium* (30) and 66% to CitH from *Klebsiella pneumoniae* (35), both of which are chromosomally encoded. Citrate transport by CitH was shown to occur in symport with protons (36), and a similar mechanism is likely to apply for the plasmid-encoded CitA carriers from *E. coli*.

Under anoxic conditions, *E. coli* can utilize citrate if an oxidizable cosubstrate is present. The corresponding fermentation pathway is shown in Fig. 1. After uptake into the cell, citrate is split by citrate lyase to acetate and oxaloacetate. The latter is subsequently converted via malate and fumarate to succinate by malate dehydrogenase, fumarase, and fumarate reductase. The reducing equivalents required for this conversion must be provided by the oxidation of the cosubstrate, e.g., glucose or glycerol (17). Two enzymes must be specifically induced for anaerobic citrate dissimilation, i.e., a citrate uptake system and citrate lyase. The latter enzyme has been purified from *E. coli* and was shown to consist of three subunits (α [55.5 kDa], β [35 kDa], and γ [12.5 kDa]), similar to citrate lyase from other bacterial species (25).

After the completion of the *E. coli* genome sequence (2), a gene cluster which showed a high degree of similarity to the *citCDEFG* operon of *K. pneumoniae* was identified between 13.9 and 14.2 min (Fig. 2). These genes encode the three subunits of citrate lyase (CitD, CitE, and CitF), a ligase required for the acetylation of the 2-(5'-phosphoribosyl)-3'-dephosphocoenzyme-A prosthetic group (CitC), and a protein (CitG) which presumably is involved in the biosynthesis or the covalent attachment of the prosthetic group (4). The proteins deduced from the *E. coli* genes *citC*, *citD*, *citE*, *citF*, and *citG* exhibited 50.6, 44.9, 65.1, 70.7, and 48.3% sequence identity to the corresponding *K. pneumoniae* proteins, respectively. A noticeable difference between the two gene clusters was the presence of an additional open reading frame (designated *citX* in Fig. 2) between the *E. coli* *citF* and *citG* genes. The *citAB* genes located upstream and divergent to *E. coli* *citC* encode proteins which are most closely related to the *K. pneumoniae* CitA-CitB two-component signal transduction system (42.0 and 48.7% identity, respectively). The *K. pneumoniae* CitA and CitB proteins are essential for expression of the genes specifically involved in citrate fermentation, including *citCDEFG* (5). The sensor kinase CitA was proposed to function as a citrate sensor (5), and the response regulator CitB was shown to bind to two sites extending from -50 to -96 upstream of the *citC* transcription start site and from -55 to -89 upstream of the *citS* transcription start site (20). Phosphorylation led to 10- to 100-fold increase of the apparent binding affinity (20). The *E. coli* *citB* gene has also been designated *criR*, because the deduced amino acid sequence is identical to the sequence of the CriR protein from *Shigella flexneri*, which has been implicated in the regulation of the *ipa* genes (24). Since *E. coli* does not possess an invasion plasmid carrying *ipa* genes, the primary function of the *citAB* gene products is presumably the regulation of the

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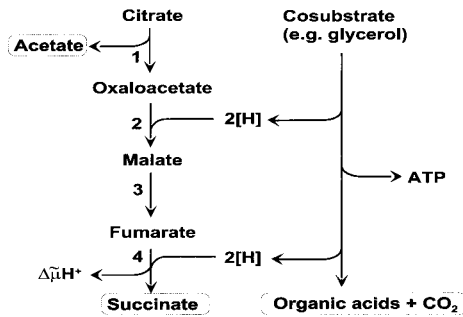


FIG. 1. Cosubstrate-dependent citrate fermentation by *E. coli*. The enzymes involved in the conversion of citrate to succinate are (1) citrate lyase, (2) malate dehydrogenase, (3) fumarase, and (4) fumarate reductase.

citrate lyase genes, as in *K. pneumoniae*. This supposition is supported by the similarity of the DNA-binding helix-turn-helix motifs of the two CitB response regulators and by the similarity of the *citC* upstream regions. With respect to the gene designations, one should be aware that in *E. coli* *citA* denotes both a plasmid-linked gene encoding a citrate carrier and a chromosomally located gene encoding a histidine sensor kinase. Similarly, *citB* denotes a gene associated with the plasmid-linked *citA* and also a chromosomally located gene encoding the cognate response regulator of the sensor kinase CitA.

An important difference between the *K. pneumoniae* and *E. coli* *cit* gene clusters is the lack of genes encoding a Na⁺-dependent citrate carrier (*citS*) and oxaloacetate decarboxylase (*oadGAB*) in the latter species (Fig. 2). In fact, these genes are not present on the whole *E. coli* chromosome. The absence of oxaloacetate decarboxylase provides an explanation for the different fermentation pathways in these organisms (3), and the absence of a CitS-type protein necessitates the use of a different citrate uptake system. Inspection of the *E. coli* DNA region downstream of *citG* revealed a gene (*ybdS*) encoding a highly hydrophobic protein with 34% sequence identity to the 2-oxoglutarate/malate translocator from spinach chloroplasts. The *ybdS* start codon is located only 50 bp downstream of the stop codon of *citG*, indicating that *ybdS* is cotranscribed with *citCDEFXG*. In this report, we present evidence that the protein encoded by *ybdS* functions as a citrate carrier; we therefore renamed the gene *citT*, for citrate transporter.

MATERIALS AND METHODS

Bacterial strains and growth conditions. *E. coli* DH5α (Bethesda Research Laboratories) was routinely used as the host for the cloning procedures. *E. coli* JM83 (38) was used for the preparation of chromosomal DNA by the method of Marmur (18). *E. coli* BL21(DE3), which contains the phage T7 RNA polymerase gene under the control of the *lacUV5* promoter (33), served as the host for the expression of *citT* and *citS* from pET-derived plasmids (Novagen). The strains

were routinely grown at 37°C in Luria-Bertani (LB) medium (21) with shaking at 180 rpm. For testing the ability to grow with citrate as the sole carbon and energy source, we used either Simmons' citrate agar (31) supplemented with 12 μM thiamine or a liquid medium (adjusted to pH 6.8 with Na⁺-poor KOH) that contained 7 mM citric acid, 20 mM KH₂PO₄, 11 mM (NH₄)₂SO₄, 1 mM MgSO₄, 12 μM thiamine, 0.4% (vol/vol) trace elements solution (7), and either 25 mM Na₂SO₄ or 25 mM K₂SO₄. The latter medium contained less than 50 μM Na⁺ as determined by atomic absorption spectroscopy. The antibiotics ampicillin (100 to 200 μg/ml) and kanamycin (50 μg/ml) were used as appropriate. Cells used for growth studies with citrate minimal medium were pregrown in LB medium and washed with Na⁺-poor minimal medium before use as inoculum.

Recombinant DNA work. For routine work with recombinant DNA, established protocols were used (27). For the construction of a *citT* expression plasmid, the *citT* gene was amplified from chromosomal *E. coli* DNA by using the oligonucleotides ec-citT-for (5'-GATTCTGAAGCTTCATATGTCTTTAGCAA AAGATAATATATGG-3') and ec-citT-rev (5'-CCGCGAATTCTTAGTCCCA CATGGCAGAAATCGGCCAG-3'). In ec-citT-for, the ATG start codon of *citT* is part of an *NdeI* restriction site, which is preceded by a *HindIII* site and five additional nucleotides, allowing increased restriction efficiency. In ec-citT-rev, a *BamHI* restriction site is introduced after the *citT* stop codon. The PCR mixture contained 500 ng of genomic DNA of *E. coli* JM83, 0.5 μM each primer, 0.2 mM deoxynucleoside triphosphates, 1× buffer for cloned *Pfu* DNA polymerase, and 2.5 U of *Pfu* DNA polymerase (Stratagene). After an initial denaturation step (2 min at 95°C), 30 cycles consisting of 15 s at 95°C, 15 s at 62°C, and 4 min at 72°C were performed, followed by a terminal elongation step (4 min at 72°C). The complete PCR mix was subsequently separated on a 1% agarose gel, and the expected 1.49-kb fragment was isolated with Qiaex (Qiagen). After restriction with *HindIII* and *BamHI*, the PCR product was purified with a QIA-Quick spin column and ligated with *HindIII/BamHI*-restricted pUC19 (38), resulting in pUC19-citT. Since the *citT* gene in pUC19-citT was not preceded by a well-conserved ribosome binding site, a 1.46-kb *NdeI/BamHI* fragment from pUC19-citT was cloned in pET24b (Novagen) restricted with the same enzymes, resulting in pET24-citT. Plasmid pCitS_{His-3} is a derivative of pET16b (Novagen) and is used for synthesis of the *K. pneumoniae* CitS citrate carrier modified with an N-terminal His₁₀ tag (23).

DNA sequence analysis. The sequence of the *citT* gene present in plasmid pET24-citT was determined by the dideoxynucleotide chain termination method (28), using the protocols and equipment for automated DNA sequencing (Sequencher 310 and PRISM Ready Reaction Dye-Deoxy terminator cycle sequencing kit from Applied Biosystems). For this purpose, pET24-citT was purified with a Qiagen Tip-500 column. A primer-walking strategy involving eight primers derived from *citT* and two primers derived from pET24b was applied. Computer-assisted DNA and protein sequence analysis was performed with the software package of the University of Wisconsin Genetics Computer Group. Prediction of the transmembrane helices indicated in Fig. 3 was performed with the TopPred II software (6) and the TMpred software (12).

Transport experiments. For transport experiments, *E. coli* BL21(DE3) transformed with either pET24b, pET24-citT, or pCitS_{His-3} was grown in LB medium with appropriate antibiotics to an optical density at 600 nm (OD₆₀₀) of ca. 0.7 to 1.0. Subsequently, cells were washed once with 50 mM morpholineethanesulfonic acid-Tris buffer (pH 7.0) and concentrated 10-fold in the same buffer. The protein concentration of the resulting cell suspension was calculated by assuming that an OD₆₀₀ of 1.4 corresponds to 10⁹ cells/ml and those 10⁹ cells contain 150 μg of protein (21). At time zero, 98 μl of the concentrated cell suspension preincubated at 25°C was added to 2 μl of 1.2 mM [1,5-¹⁴C]citrate (145 cpm/pmol). After various times at 25°C, transport was terminated by the addition of 0.9 ml of ice-cold 0.1 M LiCl followed by rapid filtration through 0.45-μm-pore-size cellulose nitrate filters (diameter, 25 mm; Sartorius). The filters were washed once with 1 ml of ice-cold 0.1 M LiCl and then placed into scintillation vials. Immediately afterwards, 4 ml of scintillation fluid (Irga-Safe Plus; Packard) was added, and the entrapped [1,5-¹⁴C]citrate was determined by liquid scintillation counting. The values obtained in this way were corrected for a time zero value obtained as follows: 98 μl of cell suspension was first mixed with 0.9 ml of

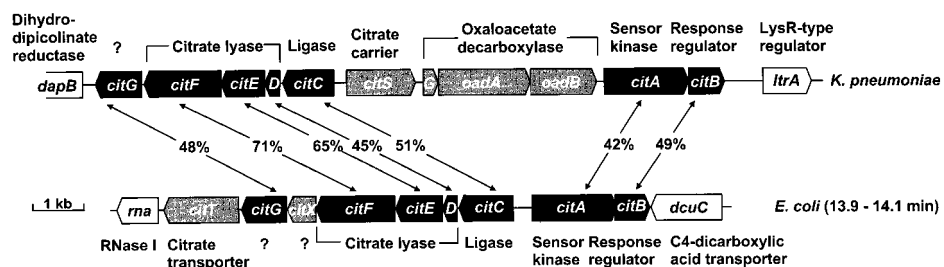


FIG. 2. *E. coli* genes required for citrate fermentation and comparison with the corresponding *K. pneumoniae* genes. Genes shaded in dark gray represent those *K. pneumoniae* genes involved in citrate fermentation which are also present in *E. coli*; genes shaded in light gray are those present only in the *E. coli* or only in the *K. pneumoniae* cluster. All other indicated genes are presumably not directly involved in citrate fermentation.

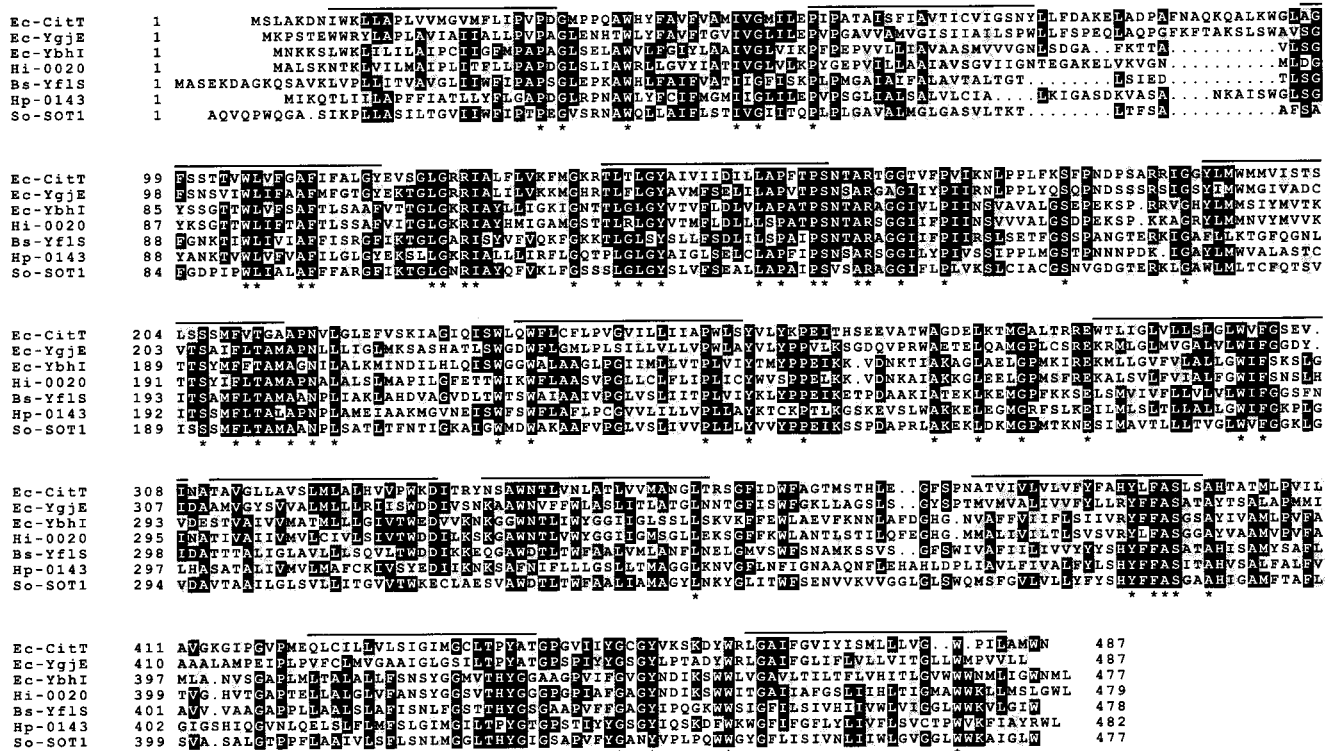


FIG. 3. Amino acid sequence alignment of the *E. coli* (Ec) CitT protein with the 2-oxoglutarate/malate translocator from spinach (*Spinacia oleracea* [So]) chloroplasts and five eubacterial gene products. Identical residues present in at least four of the sequences are framed in black; conservative exchanges are framed in gray. Asterisks indicate residues conserved in all sequences. Putative transmembrane helices within the CitT sequence are overscored. Details on the aligned sequences can be found in the text and in Table 1. Hi, *Haemophilus influenzae*; Bs, *Bacillus subtilis*; Hp, *Helicobacter pylori*.

ice-cold 0.1 M LiCl and then applied to 2 μl of 1.2 mM [1,5-¹⁴C]citrate. This mixture was rapidly filtered as described above.

To determine whether certain di- and tricarboxylic acids are able to trigger the efflux of [1,5-¹⁴C]citrate previously taken up by the cells, 98 μl of cell suspension was incubated with 2 μl of 1.2 mM [1,5-¹⁴C]citrate for 30 s. Subsequently, these preloaded cells (100 μl) were applied to 1 μl of a 1 M solution of either citric acid, succinic acid, fumaric acid, or fumarate (Na⁺-salt). After various times, 0.9 ml of ice-cold 0.1 M LiCl was added, and the mixtures were filtered and treated as described above.

The transport experiments with 0.91 mM DL-[1,4-¹⁴C]tartrate (163 cpm/pmol; custom synthesized by Anawa) were performed in the same way as described above for [1,5-¹⁴C]citrate.

RESULTS AND DISCUSSION

Aerobic growth of *E. coli* harboring a *citT* expression plasmid on citrate as the sole carbon and energy source. The open reading frame located at 13.9 min on the *E. coli* chromosome (designated *citT*), starting 50 bp downstream of the *citG* stop codon, encoded a protein of 487 amino acids (53.1 kDa) that was very hydrophobic and contained 12 putative transmembrane helices (Fig. 3). The physical proximity of *citT* to the citrate lyase genes and the fact that CitT showed 34% amino acid sequence identity to the 2-oxoglutarate/malate translocator from spinach chloroplasts (37) suggested to us that CitT might function as a citrate carrier. To test this assumption, the *citT* gene was amplified by PCR from chromosomal DNA of *E. coli* and ligated as a 1.5-kb *HindIII/BamHI* fragment in pUC19, which allows transcription of *citT* from the vector-encoded *lac* promoter. The ligation mixture was transformed into *E. coli* DH5α and plated on Simmons' citrate agar. After 48 h at 37°C, several citrate-positive colonies were identified by the color change of the agar from green to blue. This color change of the pH indicator bromthymol blue is observed only

with cells able to utilize citrate as the carbon and energy source, which results in alkalization of the medium. Cells unable to utilize citrate, such as *E. coli* DH5α containing only the vector pUC19, form only very small colonies (diameter, <1 mm), presumably by using residual carbon sources present in the agar, and bromthymol blue remains green. Restriction analysis of plasmid DNA isolated from several Cit⁺ clones showed that all contained pUC19 with the 1.5-kb *HindIII/BamHI* insert carrying *citT*. One of the pUC19-*citT* plasmids was transformed again into *E. coli* DH5α and plated on Simmons' citrate agar. In this case, all transformants were able to utilize citrate, confirming that *citT* is responsible for the Cit⁺ phenotype. Besides citrate, isocitrate could also be used as the sole carbon and energy source by *E. coli* DH5α harboring pUC19-*citT*.

To provide a good ribosome binding site (5'-AAGGAG-3') upstream of the *citT* start codon, a 1.46-kb *NdeI/BamHI* fragment obtained from pUC19-*citT* was cloned into pET24b. Plasmid pET24-*citT* isolated from one of the resulting Cit⁺ clones was used for DNA sequence analysis of the region encompassing *citT*. The sequence was 100% identical to the one present in the database, and thus this plasmid was suitable for further studies. *E. coli* BL21(DE3) harboring pET24-*citT* was able to grow aerobically in citrate minimal medium. After a lag phase of about 40 h, the cells grew within 24 h from an OD₆₀₀ of 0.05 to an OD₆₀₀ of about 0.5, whereas the control cells containing the vector pET24b were unable to multiply in this medium (data not shown). To find out whether mutations had occurred within *citT* during the long lag phase, pET24-*citT* was isolated from citrate-grown cells, and the region encompassing *citT* was sequenced again. No mutations were detected, showing that other adaptation processes must be responsible for the

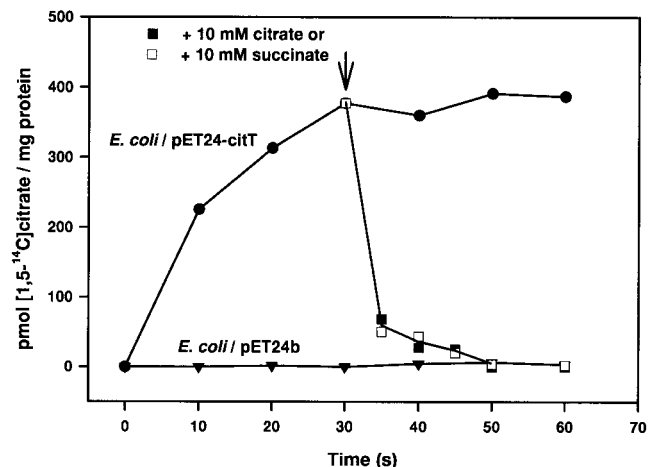


FIG. 4. Uptake and efflux of [1,5-¹⁴C]citrate by cell suspensions of *E. coli* BL21(DE3) containing either pET24b or pET24-citT. The transport experiments were performed as described in Materials and Methods.

40-h lag phase. Growth of *E. coli*/pET24-citT was independent of sodium ions in the concentration range tested (50 μ M to 50 mM).

Transport studies with cell suspensions. The growth experiments described above confirmed our suggestion that CitT catalyzes the uptake of citrate. Further evidence was obtained by transport studies with cell suspensions. As shown in Fig. 4, cell suspensions of *E. coli* BL21(DE3)/pET24-citT catalyzed [1,5-¹⁴C]citrate uptake with an initial rate of 1.4 nmol min⁻¹ (mg of protein)⁻¹, whereas cells containing only the vector pET24b did not show any [1,5-¹⁴C]citrate accumulation. Citrate uptake by cells containing CitT was independent of Na⁺ ions in the concentration range tested (10 μ M to 50 mM; data not shown). The similarity of CitT to the 2-oxoglutarate/malate translocator from spinach chloroplasts and the fact that the final product of citrate fermentation in *E. coli* is succinate led us to assume that CitT may function as a citrate/succinate antiporter. Indeed, the addition of 10 mM succinate to cells that previously had taken up [1,5-¹⁴C]citrate led to a complete efflux of the ¹⁴C label with an initial rate of 3.7 nmol min⁻¹ (mg of protein)⁻¹ (Fig. 4). The same effect was obtained by the addition of 10 mM citrate. Fumarate (10 mM) caused a comparable effect, but the rate was somewhat lower and efflux was not complete (data not shown). In a control experiment, uptake and efflux of [1,5-¹⁴C]citrate were analyzed with *E. coli* BL21(DE3)/pCitS_{His-3}. These cells contain the citrate carrier CitS from *K. pneumoniae* modified by an N-terminal His₁₀ tag (22, 23). Since the CitS carrier was shown to be highly specific for citrate as a substrate (1), we expected that only citrate, not succinate or fumarate, would be able to trigger an efflux of [1,5-¹⁴C]citrate previously taken up. As shown in Fig. 5, cells containing CitS catalyzed the uptake of citrate with an initial rate of 5.6 nmol min⁻¹ (mg of protein)⁻¹. Addition of 10 mM citrate led to a partial efflux of the ¹⁴C label from [1,5-¹⁴C]citrate-loaded cells [initial rate, 7.3 nmol min⁻¹ (mg of protein)⁻¹], whereas succinate or fumarate did not elicit such a response. This result confirms that the succinate- and fumarate-induced efflux observed with the *E. coli*/pET24-citT cells (Fig. 4) is catalyzed by the CitT protein rather than by other carriers present in the cytoplasmic membrane. The fact that only a partial efflux of the ¹⁴C label was observed in the experiment shown in Fig. 5 can be explained by the fact that part of [1,5-¹⁴C]citrate had been converted to other intermediates

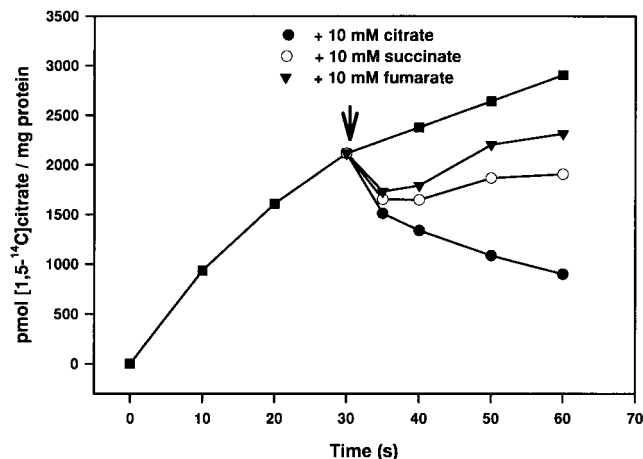


FIG. 5. Uptake and efflux of [1,5-¹⁴C]citrate by cell suspensions of *E. coli* BL21(DE3) containing pCitS_{His-3}. The transport experiments were performed as described in Materials and Methods.

of the tricarboxylic acid cycle within the 30 s before addition of the unlabeled citrate. These other intermediates are transported not by CitS but apparently by CitT, as indicated by the complete efflux shown in Fig. 4.

As outlined below, the protein most similar to CitT is the product of the *E. coli* *yjgE* gene (Fig. 3), which we predict to function as a tartrate carrier. We therefore tested whether CitT also catalyzes the uptake of DL-[1,4-¹⁴C]tartrate. In contrast to citrate, *E. coli* can utilize tartrate as a carbon and energy source under oxic conditions and contains an appropriate transport system (14). This was confirmed by the fact that DL-[1,4-¹⁴C]tartrate uptake was observed with *E. coli* BL21 (DE3) cells harboring the control plasmid pET24b at a rate of 0.25 nmol min⁻¹ (mg of protein)⁻¹ (Fig. 6). Addition of 10 mM citrate did not lead to an efflux of DL-[1,4-¹⁴C]tartrate from these cells. With cells harboring pET24-citT, a significantly higher rate of DL-[1,4-¹⁴C]tartrate uptake [0.62 nmol min⁻¹ (mg of protein)⁻¹] was found, and addition of 10 mM citrate led to a complete efflux of the ¹⁴C label (Fig. 6). As in the case with [1,5-¹⁴C]citrate, efflux was significantly faster [2.5 nmol min⁻¹ (mg of protein)⁻¹] than uptake.

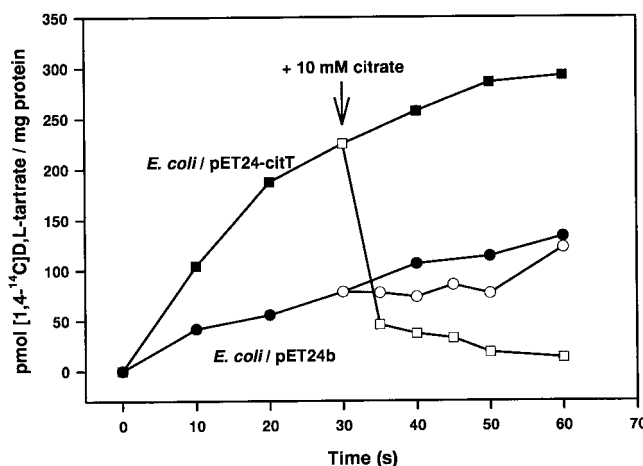


FIG. 6. Uptake and efflux of DL-[1,4-¹⁴C]tartrate by cell suspensions of *E. coli* BL21(DE3) containing either pET24b or pET24-citT. The transport experiments were performed as described in Materials and Methods.

TABLE 1. Bacterial transporters related to the 2-oxoglutarate/malate translocator from spinach chloroplasts

Protein	SwissProt accession no.	Organism	Amino acids	Mol wt (10 ³)	% Apolar amino acids	pI	% Identity with:		Demonstrated (anticipated) function
							CitT	SOT1	
CitT (YbdS)	P77405	<i>Escherichia coli</i>	487	53.09	72	8.9	100	34	Citrate/succinate antiporter
YgjE	P39414	<i>E. coli</i>	487	52.91	75	9.5	44	34	(Tartrate/succinate antiporter)
YbhI	P75763	<i>E. coli</i>	477	51.35	73	10.3	35	36	(Tricarboxylate transporter)
YbhI (HI0020)	Q57048	<i>Haemophilus influenzae</i>	479	51.35	71	10.2	37	35	(Citrate/succinate antiporter)
YfiS	NA ^a	<i>Bacillus subtilis</i>	478	51.43	70	10.2	35	53	(Di- and tricarboxylate transporter)
HP0143 ^b	NA	<i>Helicobacter pylori</i>	482	52.35	73	9.8	42	33	
SOT1	Q41364	<i>Spinacia oleraceae</i>	477	50.69	73	9.9	34	100	2-Oxoglutarate/malate antiporter

^a NA, not available.

^b The gene encoding HP0143 contains a frameshift.

The transport experiments described above show that the *E. coli* CitT protein functions as a Na⁺-independent antiporter with relatively broad substrate specificity for C₄-dicarboxylates and tricarboxylates. Whether the initial uptake is in fact unidirectional uptake or exchange against the internal pool of dicarboxylates and tricarboxylates is not known. Nevertheless, the significantly higher rates of efflux compared to uptake clearly favor exchange as the usual transport mode of CitT. In order to characterize the catalytic properties of CitT in more detail, a proteoliposomal system with purified CitT seems more suitable than studies with whole cells, as shown before for the CitS protein from *K. pneumoniae* (22, 23). Since the *citT* gene is physically linked to the citrate lyase genes and probably coregulated with them, the in vivo function of CitT is likely to be citrate-succinate exchange. The use of an antiporter system for the excretion of succinate formed by fumarate respiration seems to be the general rule in *E. coli*. Three secondary carriers for anaerobic C₄-dicarboxylate transport (DcuA, DcuB, and DcuC), all of which preferentially catalyze exchange but also catalyze unidirectional uptake of C₄-dicarboxylates, have been identified (8, 9, 32, 39). None of these carriers shows significant sequence similarity to CitT (identity of <20%). The *dcuC* gene is located immediately downstream of *citA* in inverse orientation (Fig. 2), but there is no evidence at present for involvement of DcuC in citrate fermentation.

Proteins related to CitT. A search for proteins related to CitT from *E. coli* led to the identification of five bacterial polypeptides, none of which has been functionally characterized hitherto, and the 2-oxoglutarate/malate translocator (SOT1) from *Spinacia oleraceae* (37). The SOT1 protein is located in the inner envelope membrane of spinach chloroplasts, where it catalyzes the import of 2-oxoglutarate in exchange for stromal malate. The protein has recently been purified and functionally reconstituted into liposomes (19). Besides malate, succinate, fumarate, and 2-oxoglutarate can be used as counterions (37). From the sequence alignment shown in Fig. 3, it is obvious that the chloroplast protein is related to CitT and the other bacterial proteins described below. Moreover, preliminary characterization of the catalytic properties of CitT indicates that the transport mechanism of this carrier is similar to that of the SOT1 protein.

The *E. coli* YgjE protein shows 44% sequence identity to CitT (Fig. 3). It is located at 69.08 min on the *E. coli* chromosome immediately downstream of the *tdAB* genes encoding an oxygen-labile L-tartrate dehydratase (26). This enzyme converts L-tartrate to oxaloacetate and is induced by L- and meso-tartrate during anaerobic growth with glycerol as cosubstrate. Oxaloacetate is subsequently converted to succinate, using the reducing equivalents provided by the oxidation of the cosubstrate. Thus, the tartrate fermentation pathway is very similar

to the citrate fermentation pathway depicted in Fig. 1 and leads to the same end product. In view of these facts, it seems plausible that YgjE could function as a tartrate/succinate antiporter.

Besides YgjE, another protein (designated YbhI) with 35% sequence identity to CitT is encoded by the *E. coli* K-12 chromosome at 17.27 min. Interestingly, a spontaneous *E. coli* K-12 mutant (strain D2004) that is able to utilize citrate, *cis*-aconitate, *trans*-aconitate, and isocitrate aerobically has been described (11). Genetic analysis of strain D2004 indicated that the mutations responsible for the Cit⁺ phenotype are located in *cit* genes that are linked to the *gal* operon. Since the *galETKM* genes are located between 16.9 and 17.1 min, *ybhI* is likely to be one of the genes expressed in mutant D2004. Consequently, a function of YbhI as tricarboxylic acid transporter is very likely. Additional support for this assumption is provided by the fact that the open reading frame downstream of *ybhI* encodes a protein of 761 amino acids that exhibits similarity to aconitase from several organisms.

The HI0020 protein of *Haemophilus influenzae* exhibits 38% sequence identity to CitT. It was tentatively identified as a 2-oxoglutarate/malate translocator due to its similarity to the SOT1 protein from spinach chloroplasts (10). However, since the HI0020 gene is located immediately downstream of the citrate lyase genes, as is the *citT* gene of *E. coli*, it seems more likely that the HI0020 protein is functionally related to citrate metabolism.

The YfiS protein from *Bacillus subtilis* (16) exhibits 35% sequence identity to CitT. The corresponding gene is located at 828.9 kb on the chromosome downstream of the *pel* gene encoding pectate lyase. Remarkably, the two genes downstream of *yfiS* (*citS* and *citT*) encode a two-component regulatory system with significant similarity to the CitA/CitB system from *K. pneumoniae* (5). The proteins derived from *citS* and *citT* exhibit 28 and 33% amino acid sequence identity to the sensor kinase CitA and the response regulator CitB, respectively.

The predicted HP0143 protein from *Helicobacter pylori* possesses 42% sequence identity to CitT, if the predicted frameshift within the coding sequence is ignored (34). In the vicinity of the HP0143 gene (located at 156 kb on the chromosome), no genes involved in citrate or tartrate metabolism which might give a clue to the function of the HP0143 protein are present.

In Table 1, some properties of the proteins described above are summarized. It is evident that they all consist of 477 to 487 amino acids, 70 to 75% of which are apolar. Moreover, all of these proteins are basic, with calculated pIs of between 8.9 and 10.3. Together with the overall sequence similarity, the data support the assumption that these proteins form a new family of secondary transporters.

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