INTRODUCTION

Enzymes capable of functioning efficiently at low temperature, termed cold-adapted enzymes, represent one of the strategies for environmental adaptation adopted by psychrophiles, bacteria adapted to cold habitats, because the catalytic rates of enzyme reactions are strongly dependent on temperature. The cold-adapted enzymes display higher specific activities at low temperature and pronounced thermostability compared to their mesophilic and thermostable counterparts; these properties are probably accomplished by the enhanced conformational flexibility of the enzyme proteins, which is regarded as the main adaptive feature to low temperature (Gerday et al., 1997; Fields & Somero, 1998; Russell et al., 1998; Kim et al., 1999; Alvarez et al., 1998; Bentahir et al., 2000).

Directed modification of the enzyme proteins by random PCR mutagenesis is a useful approach to examine their adaptation mechanism. By comparing the amino acid sequences of mutated enzyme proteins that exhibit thermodynamic properties different from those of the wild-type enzyme, the cause of the alteration can be specified. Such studies for thermophilic (Suzuki et al., 2001; Lönn et al., 2002), mesophilic (Taguchi et al., 1999; Wintrode et al., 2000) and psychrophilic enzymes (Miyazaki et al., 2000) have been reported. Even among mutated enzymes with similar characteristics to each other, various patterns of amino acid substitution have been observed (Suzuki et al., 2001). Furthermore, the substituted amino acid residues in a cold-adapted mutant (P3C9) of mesophilic subtilisin SSII were not necessarily identical to the corresponding residues of the naturally cold-adapted subtilisin S41 (Wintrode et al., 2000), supporting the view that nature has not tested all possibilities for the adaptation of proteins to low temperature (Gerday et al., 1997).

Cold adaptation of enzymes in central metabolism should
be essential for the survival of psychrophilic bacteria. Isocitrate lyase (ICL; EC 4.1.3.1) catalyses the cleavage of isocitrate to glyoxylate and succinate, and plays important roles in the metabolism of acetate and fatty acids in micro-organisms and higher plants as a key enzyme of the glyoxylate cycle (Kornberg, 1966; Vanni et al., 1990; Cozzone, 1998). We previously reported that the ICLs of two psychrophilic bacteria, Colwellia maris and Colwellia psychrerythraea, are homotetrameric, as are their counterparts in other organisms, including Escherichia coli, that they are typical cold-adapted enzymes, and that the expression of the genes encoding these enzymes is cold-inducible (Watanabe et al., 2001, 2002a, 2002b). Furthermore, the recently resolved crystal structures of ICLs and their complexes with ligands from several organisms can provide us with useful information to understand their catalytic function (Britton et al., 2000, 2001; Sharma et al., 2000). We previously found that about 20 amino acid residues among those strictly conserved in ICLs from various organisms are substituted in the corresponding enzyme of C. maris (CmICL) (Watanabe et al., 2002a). Furthermore, some of the substitutions were found in amino acid residues essential for the catalytic function of E. coli ICL (Ko & McFadden, 1990; Ko et al., 1991, 1992; Diehl & McFadden, 1993, 1994; Rehman & McFadden, 1996, 1997a, b, c) or the adjacent residues. Similar substitutions of conserved amino acid residues have also been observed in other cold-adapted enzymes (Davail et al., 1994). In this study, to examine whether such substituted residues are related to the cold adaptation of CmICL, they were exchanged for amino acid residues homologous to mesophilic counterparts by site-directed mutagenesis, and the properties of the mutated ICLs were investigated.

**METHODS**

**Construction of expression vectors for C. maris and E. coli ICLs.** The genes encoding the ICLs of C. maris and E. coli (CmICL and EcICL, respectively) were amplified by PCR so as to introduce restriction sites for BamHI and Sall at the 5’ and 3’ terminals of the ORFs, respectively. Therefore, the following primers were synthesized: for CmICL, sense primer 5’-CGATAGAAGATCGTCTACCTATA-CAAGTCGAAATTAGAGCTAAGC-3’ (48-mer) and antisense primer 5’-GATCTTCGAGCTTCTTGAACATGGTATGATGCTTTGCGCCC-3’ (28-mer); for EcICL, sense primer 5’-ATAAGGAT-TCGAAAAACCGTGATACACAAATTTGAAATTTACAGAAAG-3’ (48-mer) and antisense primer 5’-TCTCTGAGCTTCTTGAACATGGTATGATGCTTTGCGCCC-3’ (28-mer). Cleavage sites for restriction enzymes are underlined. The amplification was carried out for 30 cycles in a DNA thermal cycler 2400 (Perkin-Elmer) employing 20 ml of buffer per litre of the original culture medium. Hen-egg lysozyme (2 mg ml⁻¹) was added to the cell suspension, and the mixture was gently shaken overnight at 4°C. The cells were then disrupted by ultrasonic oscillation. After the centrifugation of the cell lysate at 39 120 g for 30 min at 4°C to remove cell debris, the supernatant was centrifuged at 65 600 g for 6 h at 4°C. The resultant supernatant was loaded onto a Ni-NTA column (25 ml; Qiagen) equilibrated with Buffer A (50 mM sodium phosphate, pH 6-85, containing 5 mM MgCl₂, 0-5 M NaCl, 10 mM 2-mercaptoethanol and 10 mM imidazole), employing 20 ml of buffer per litre of the original culture medium. SDS-PAGE was carried out by the method of Laemmli (1970) with a 10% gel at 25 V.

**PAGE.** SDS-PAGE was carried out by the method of Laemmli (1970) with a 10% gel at 25 V.

**Western blot analysis.** After SDS-PAGE of the purified ICLs and the sonicated extracts of the E. coli transformant cells, the expressed proteins, to obtain plasmids pHis-CmWT and pHis-EcWT, respectively. The plasmids were transformed into competent cells of E. coli strain TOP10 by a conventional heat-shock method. Then, the amplified plasmids of pHis-CmWT and pHis-EcWT were purified by the Wizard plus SV minipreps DNA purification system (Promega). To check precise insertion of the CmICL and EcICL genes into pTrcHisB, the insert DNAs in the resulting plasmids were sequenced in both directions by using a DYEnamic ET Terminator Cycling Sequencing Kit (Amersham Pharmacia Biotech) with an ABI 373A DNA sequencer (Applied Biosystems). The sequence was analysed with the Genetyx computer program (Software Development Co.).

**Site-directed mutagenesis.** Site-directed mutagenesis was carried out by the standard PCR method. The following primers were used to introduce the mutations into CmICL: sense primer207 and antisense primer207 for the substitution of His for Gln207 were 5’-GTTGCTTTGGTTCATCTATTGAAAC-3’ (28-mer) and 5’-GCTTTTCAATATGAAGAGCAAGACC-3’ (28-mer), respectively; sense primer307 and antisense primer307 for the substitution of Tyr for Gln207 were 5’-GCTATTGATGCTTTGCGCCC-3’ (31-mer) and 5’-GGTTTCTGTAGTGGCACATTGGAAAC-3’ (31-mer), respectively. The mutated regions are underlined. Except for the use of 100 pmol of each primer, the PCR was performed as described above. Both the PCR products were ligated into pTrcHisB to obtain plasmids pHis-CmWT (the Q207H mutation) and pHis-CmWT (the Q217K mutation). A plasmid carrying both of the mutations (pHis-Cm207/217) was obtained by PCR with the plasmid pHis-Cm207 and a set of sense and antisense primers312. The mutated plasmids were transformed into E. coli TOP10 and purified. To verify the introduced mutations, the relevant regions of the plasmids were sequenced in both directions, as described above.

**Overexpression and purification of His-tagged ICLs.** The E. coli TOP10 cells carrying one of the expression vectors pHis-CmWT, pHis-Cm207, pHis-Cm217, pHis-Cm207/217 or pHis-EcWT were grown at 37°C in Super broth medium (12 g tryptone, 24 g yeast extract, 5 ml glycerol, 3-81 g KH₂PO₄ and 12-5 g K₂HPO₄ per litre, pH 7-0) containing 50 mg ampicillin l⁻¹ until OD₆₀₀ of the culture reached 0-6. The cultures were rapidly cooled on ice and were further incubated for 18-24 h at 15°C after the addition of 0-1 M IPTG. Cells were then collected and resuspended in Buffer A (50 mM sodium phosphate, pH 6-85, containing 2 mM MgCl₂, 0-5 M NaCl, 10 mM 2-mercaptoethanol and 10 mM imidazole), employing 20 ml of buffer per litre of the original culture medium. The cell lysate at 39 120 g for 30 min at 4°C to remove cell debris, the supernatant was centrifuged at 65 600 g for 6 h at 4°C. The resultant supernatant was loaded onto a Ni-NTA column (25 ml; Qiagen) equilibrated with Buffer A. After thorough washing with the same buffer, the column was further washed with 50 ml Buffer B (Buffer A containing 10%, v/v, glycerol and 20 mM imidazole instead of 10 mM imidazole) and next with 50 ml Buffer C (Buffer B containing 50 mM imidazole instead of 20 mM imidazole). In each case the enzyme was then eluted with 50 ml of Buffer D (Buffer C containing 250 mM imidazole instead of 20 mM imidazole), and the eluent was concentrated with polyethylene glycol 8000 (20 ml) and dialysed against Buffer E (20 mM potassium phosphate, pH 6-85, containing 2 mM MgCl₂, 0-5 M NaCl, 1 M DTT and 0-01% NaN₃). All His-tagged recombinant ICLs were stored at 4°C and used in further experiments within a week.
proteins on the gels were transferred onto a nitrocellulose membrane (Hybond-C; Amersham Pharmacia Biotech). Western blot analysis was carried out with the ECL Western blotting detection system (Amersham Pharmacia Biotech) and either rabbit antibody against the C. maris ICL (Watanabe et al., 2002a) or mouse antibody against the additional (His)6-Gly sequence at the N-terminal of the over-expressed ICLs (Invitrogen). Native ICL of C. maris was purified as described previously (Watanabe et al., 2001).

**Enzyme assay.** Unless otherwise stated, the ICL activity was assayed at pH 6.85 as described previously (Watanabe et al., 2001). Optimum pH for activity was determined with the following buffers: potassium phosphate at pH 6.0–7.6 and Tris/HCl at pH 7.5–8.0. Protein was measured by the method of Lowry et al. (1951) with BSA as a standard.

**Measurement of kinetic and thermodynamic parameters.** The kinetic parameters, $K_m$ and $V_{max}$ of ICL were calculated by Lineweaver–Burk analysis. Molecular masses of the wild-type and the mutants of CnICL (Cm-Q207H and Cm-Q217K) and EdICL with the additional N-terminal His-tags were calculated as 246,641, 246,677, 246,641 and 204,124 Da, respectively, from the gene sequences. The activation energies ($E_a$) of the enzymes were calculated by Arrhenius plotting. The thermodynamic activation parameters were calculated according to the following equations.

\[
\Delta G^\circ = RT \times \left(23.76 + \ln T - \ln k_{cat}\right)
\]

\[
\Delta H^\circ = E_a - RT
\]

\[
\Delta S^\circ = (\Delta H^\circ - \Delta G^\circ)/T
\]

**Digestion with trypsin.** The purified recombinant ICLs were dialysed overnight at 4 °C against 0.1 M NaHCO$_3$ (pH 8.1). The ICL enzymes (100 μg) were then digested at 4 °C with 0.1 μg trypsin in the same buffer. At appropriate times after the addition of trypsin into the reaction mixture, the mixture was withdrawn and immediately analysed by SDS-PAGE.

**Circular dichroism (CD) spectra of recombinant ICLs.** All purified recombinant ICLs were dialysed overnight at 4 °C against 20 mM potassium phosphate buffer (pH 6.85) containing 2 mM MgCl$_2$ and 1 mM DTT. UV-CD of the ICLs was measured at 20 °C with a J-725 spectropolarimeter (Jasco) by using a cuvette with a 1 cm path length. The protein concentrations were determined by the absorbance at 280 nm from the extinction coefficients 53056 and 53056 (23,677, 246,641 and 204,124 Da, respectively, from the gene sequences. The activation energies ($E_a$) of the enzymes were calculated by Arrhenius plotting. The thermodynamic activation parameters were calculated according to the following equations.

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\]

\[
\Delta H^\circ = E_a - RT
\]

\[
\Delta S^\circ = (\Delta H^\circ - \Delta G^\circ)/T
\]

**Phylogenetic analysis.** Phylogenetic analysis was done with a CLUSTAL-W program with the unrooted neighbour-joining method. The phylogenetic tree was produced with the program TreeView 1.6.1 distributed by the Bioinformatics Center of Kyoto University, Japan (www.genome.ad.jp/Japanese/).

**Molecular modelling of CnICL.** The three-dimensional structural model of CnICL was built on the basis of its homology to EdICL (PDB No. 1HGW) with the program SWISSPDB VIEWER.

**RESULTS**

**Site-directed mutagenesis of CmICL**

EdICL has been studied more extensively than its counterparts in other organisms with regard to the catalytic mechanism and crystal structure (Britton et al., 2001; Ko & McFadden, 1990; Ko et al., 1991, 1992; Diehl & McFadden, 1993, 1994; Rehman & McFadden, 1996, 1997a, b, c). We previously reported that, in the cold-adapted CmICL, several amino acid residues strictly conserved in the ICLs of many organisms, including E. coli, are substituted, and some unique insertions of 3–36 amino acid residues are present (Watanabe et al., 2002a).

Among these substitutions, we noted Gln207 and Gln217 of CmICL (stars in Fig. 1a) for the following two reasons: (1) from studies on site-directed mutagenesis, His184 and Lys194 of EdICL, corresponding to the two Gln residues of CmICL, were reported to be essential for catalytic function of the enzyme (Diehl & McFadden, 1994; Rehman & McFadden, 1997b), and (2), in spite of low homology between the amino acid sequences of CmICL and EdICL (less than 30%), this region containing the two Gln residues is more highly conserved than other regions of the enzyme proteins, probably due to the existence of amino acid residues constituting the active site (asterisks in Fig. 1a). Thus, the Q207H and Q217K mutations were introduced into the CmICL by site-directed mutagenesis.

**Expression of recombinant ICLs in E. coli**

Wild-types of CmICL and EdICL (Cm-WT and Ec-WT) and three CmICL mutants (Cm-Q207H, Cm-Q217K and Cm-Q207H/Q217K) with the N-terminal His-tags were expressed at 15 °C in E. coli cells because of the marked thermolability of CmICL (Watanabe et al., 2001). In spite of the induction by IPTG, all recombinants of CmICL were expressed at lower levels than Ec-WT (Fig. 2a). The yields of the purified Cm-WT, Cm-Q207H, Cm-Q217K and Ec-WT were estimated to be 3.60, 0.55, 3.64 and 17.0 mg protein per litre of culture, respectively (mean of three independent experiments). All recombinant ICLs, except for Cm-Q207H/Q217K, were purified to homogeneity by single chromatography on a Ni$^{2+}$-chelating affinity column (Fig. 2b). The purification of the Cm-Q207H/Q217K double mutant was unsuccessful since its expression in the E. coli cells was too poor (Fig. 2a). Mobilities on the gel of both wild-type and mutated CmICL recombinants were slightly slower than that of the native CmICL purified from C. maris cells (estimated as 63 and 65 kDa, respectively) due to the addition of 33 amino acid residues including His-tags at their N-terminals (Fig. 2b). Cm-WT, Cm-Q207H and Cm-Q217K cross-reacted with anti-CmICL antibody, whereas Ec-WT did not at all (Fig. 2c). These were consistent with the previous results for the native CmICL and EdICL (Watanabe et al., 2002a). Although Ec-WT as well as Cm-WT had the His-tag attached at the N-terminal, the molecular mass of the Ec-WT subunit (47 kDa) was much smaller than that of Cm-WT (Fig. 2b) because CmICL has several insertions of amino acid residues (Watanabe et al., 2002a). On the other hand, all recombinant ICLs cross-reacted at similar intensities with the anti-His-tag antibody (Fig. 2d).

**Temperature dependence of ICL activity**

The activities of recombinant CmICLs and Ec-WT at various temperatures are shown in Fig. 3(a). The optimum
pH values for the recombinant CmICLs and Ec-WT were 6.85, almost the same as that of the native CmICL (pH 6.8; Watanabe et al., 2001). Specific activities of the ICLs at optimum temperature are summarized in Table 1. The optimum temperature for activity (20–25 °C) and specific activity at the optimum temperature (14±3±0.3 s–1; two independent experiments) of Cm-WT were similar to those of the native CmICL (20 °C and 18±1 s–1) (Watanabe et al., 2001). In addition, rapid reduction of the activity above 30 °C was found in Cm-WT as well as the native CmICL. These results suggest that the additional His-tag at the N-terminal did not affect the activity and thermal characteristics of the enzyme. Cm-Q207H and Cm-Q217K possessed specific activities comparable to that of Cm-WT, but their optimum temperatures increased by about 5 °C and 7±5 °C, respectively, from that of Cm-WT (Fig. 3a and Table 1). Specific activity of the mesophilic Ec-WT increased markedly with the elevation of assay temperature at least up to 40 °C and, above 30 °C, was much higher than the maximum activity of Cm-WT. On the other hand, the specific activity of Cm-WT between 10 °C and 20 °C was 1.3–1.6-fold higher than that of Ec-WT. Interestingly, the activity of both Cm-Q207H and Cm-Q217K between 10 °C and 25 °C was clearly lower than that of Cm-WT. Relative
activities of Cm-Q207H and Cm-Q217K at 10°C were 8·1 ± 1·3 and 9·1 ± 0·4 % (two independent experiments), respectively, and these values were almost the same as that of Ec-WT (8·2 ± 1·1 %; two independent experiments) (Fig. 3b). This result suggests the possibility that Gln207 and Gln217 of CmICL are responsible for cold adaptation.

**Thermostability of the recombinant CmICLs and EcICL**

To estimate whether the mutations of CmICL affect the thermostability of its activity, residual activities were assayed after heat treatment for various times at 30°C for CmICLs and 40°C for Ec-WT (Fig. 3c and Table 1). Similarly to the native CmICL, irreversible inactivation of recombinant CmICLs proceeded during incubation at this temperature. The half time of inactivation of Cm-Q217K was greater than that of Cm-WT. On the other hand, the half time of Cm-Q207H was shorter than that of Cm-WT, in spite of its higher optimum temperature for activity. Although Ec-WT was incubated at 40°C, the half time of inactivation of this enzyme was longer than that of all the CmICLs, and it was hardly inactivated even after incubation for 1 h at 30°C (data not shown).

**Kinetic and thermodynamic parameters of the ICLs**

The effects of temperature on the kinetic parameters of recombinant CmICLs and EcICL were examined (Fig. 4). $K_m$ values for isocitrate ($K_m^{isocitrate}$) of all ICL enzymes tested were increased by elevating temperature. Whereas the $K_m^{isocitrate}$ values of Cm-Q207H were 2·2–3·5-fold larger than those of Cm-WT, the values of Cm-Q217K were
0.6–0.8 times those of Cm-WT, and were almost the same as those of Ec-WT. On the other hand, catalytic efficiencies ($k_{cat}/K_m$) of these ICLs were maximal near the optimum temperature for activity of each enzyme (Fig. 4b). The catalytic efficiency of Cm-WT at 10 °C was 1.2-fold that of Ec-WT and was 41.3% of its maximum value at 20 °C, while the relative efficiency values of Cm-Q207H, Cm-Q217K and Ec-WT at the same temperature were only 13.5, 23.3 and 15%, respectively. These results indicate that the catalytic function of both Cm-Q207H and Cm-Q217K was impaired at low temperature.

**Sensitivities of recombinant ICLs to trypsin**

Partial digestion with proteases has been utilized to detect structural changes in proteins and identify their exposed and flexible regions (Aghajanian et al., 2003). Therefore, the susceptibility of the recombinant CmlICLs and Ec-WT to tryptic digestion was examined (Fig. 5). All CmlICL recombinants were more sensitive to trypsin than Ec-WT. These results indicate that thermolabile CmlICLs are more susceptible to tryptic digestion than Ec-WT.

**CD spectrometric analysis of the ICLs**

CD spectra of the ICLs were measured at 20 °C both to estimate the effect of these mutations on the secondary structure of the enzyme proteins and to compare the secondary structures of recombinant CmlICLs and Ec-WT with each other (Fig. 6). The contents of $\alpha$-helix and $\beta$-sheet calculated from the CD spectra were 42 and 17% in Cm-WT and 61 and 7% in Ec-WT, respectively, and the latter enzyme contained more secondary structure than the former. Little difference in the CD spectra between Cm-WT and Cm-Q217K was observed, while Cm-Q207H resembled Ec-WT rather than Cm-WT. For all ICLs, the CD spectra measured at 10 °C were almost the same as those at 20 °C (data not shown).

**Phylogenetic relationship of ICLs from various organisms**

Serrano & Bonete (2001) have reported that ICLs from various organisms can be classified into two groups: eubacterial and eukaryotic (termed subfamilies 1 and 2, respectively, in Fig. 1b), although some archaeal ICLs belong to neither subfamily. On the other hand, phylogenetic analysis with the CLUSTAL-W program of ICL amino acid sequences from 29 species of eubacteria, 24 eukaryotes, 3 Archaea and several putative eubacterial ICLs revealed that CmlICL is different from these three groups and, together with the ICL of *Hyphomicrobium methylivorum* and five putative eubacterial ICLs from *Pseudomonas*

![Fig. 3. Effect of temperature on specific activity (a) and relative activity (b) of ICL, and thermal inactivation of CmICLs and Ec-WT (c). Symbols: Cm-WT (●), Cm-Q207H (▲), Cm-Q217K (■) and Ec-WT (○). Error bars indicate values from two separate experiments. In (a) and (b), the specific activities of Cm-WT, Cm-Q217K and Ec-WT were measured by a standard assay method containing 1-7 mM isocitrate. Since Cm-Q207H showed a high $K_m$ value (see Fig. 4a), 17 mM isocitrate was used for the assay. Relative activities were expressed as percentages of the maximum activities. (c) The purified enzymes were dialysed at 4 °C against 20 mM potassium phosphate (pH 6.85) containing 2 mM MgCl$_2$ and 1 mM DTT. After incubation for the indicated time periods, at 30 °C for CmlICLs or 40 °C for Ec-WT, the dialysed enzymes were withdrawn and immediately cooled for 10 min on ice. The ICL activity was then measured at each optimum temperature as described in Methods.]()
*aeruginosa, Ralstonia metallidurans, Novosphingobium aromaticivorans, Rhodopseudomonas palustris and Caulobacter crescentus*, can be categorized to a novel group, termed subfamily 3 in Fig. 1(b). Partial sequences of typical ICLs of these subfamilies are shown in Fig. 1(a). Gln207 of *Cm* ICL was conserved in subfamily 3 ICLs, but the corresponding residue in all other ICL subfamilies was His. Furthermore, Gln217 of *Cm* ICL was substituted by different amino acid residues in ICLs other than subfamily 3: Arg in Archaea and Lys in other ICLs, respectively. Although no ICL gene has been cloned from psychrophilic and psychrotrophic bacteria except *Cm* ICL, putative ICL genes belonging to subfamily 1 could be found in the genome sequences of several psychrophilic and psychrotrophic bacteria (organisms in grey type in Fig. 1a).

**DISCUSSION**

The wild-type and mutants of *Cm* ICL together with *Ec* ICL were expressed as recombinant forms in *E. coli* cells, but the expression levels of the *Cm* ICLs were much less than that of *Ec* ICL (Fig. 2a). A similar result has been found when elongation factor 2 of an Antarctic methanogenic archaeon, *Methanococcoides burtonii*, is expressed in *E. coli* cells, and is attributed to the existence of many Arg codons (AGA and AGG) which are used rarely in *E. coli* in the aef2 gene.

<table>
<thead>
<tr>
<th>Property</th>
<th>Cm-WT</th>
<th>Cm-Q207H</th>
<th>Cm-Q217K</th>
<th>Ec-WT</th>
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</thead>
<tbody>
<tr>
<td>Optimum temperature (°C)</td>
<td>20–25</td>
<td>27–30</td>
<td>30–32.5</td>
<td>Above 40</td>
</tr>
<tr>
<td>Specific activity (s⁻¹)*</td>
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<td>14-5</td>
<td>10-5</td>
<td>54-0</td>
</tr>
<tr>
<td>Half time of thermal inactivation (s)†</td>
<td>88-2</td>
<td>58-4</td>
<td>120-3</td>
<td>161-8</td>
</tr>
<tr>
<td>Assay temperature</td>
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<td>20°C‡</td>
<td>10°C</td>
<td>30°C‡</td>
</tr>
<tr>
<td><em>Km</em> isocitrate (mM)</td>
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<td>0-47</td>
<td>0-76</td>
<td>2-07</td>
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<tr>
<td><em>Vmax</em> isocitrate (s⁻¹)</td>
<td>5-76</td>
<td>13-9</td>
<td>0-62</td>
<td>14-3</td>
</tr>
<tr>
<td><em>kcat/Km</em> (s⁻¹ M⁻¹)</td>
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<td>2-65</td>
<td>0-08</td>
<td>0-54</td>
</tr>
<tr>
<td><em>Ea</em> (kJ mol⁻¹)§</td>
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<td>89-2</td>
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<td><strong>ΔG#</strong> (kJ mol⁻¹)‖</td>
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<td>68-8</td>
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<td>105-5</td>
<td>86-9</td>
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<tr>
<td><strong>TD</strong> (kJ mol⁻¹)‖</td>
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<td>36-7</td>
<td>17-6</td>
<td>-4-52</td>
</tr>
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</table>

*Activities were assayed at each optimum temperature as described in Methods, except that activity of *Cm*-Q207H was assayed by using 17 mM isocitrate.
†The three CmICLs and Ec-WT were incubated at 30 and 40°C, respectively.
‡Optimum temperature for activity.
§*Ea* values were calculated by Arrhenius plotting.
‖Values at 10°C.

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**Table 1. Properties of Cm-WT, Cm-Q207H, Cm-Q217K and Ec-WT**

![Fig. 4. Effect of temperature on *Km* isocitrate (a) and *kcat/Km* isocitrate (b). Symbols are the same as Fig. 3.](http://mic.sgmjournals.org)

![Image 1](http://mic.sgmjournals.org)
encoding this protein (Thomas & Cavicchioli, 2000). However, this is not the case for the C. maris icl gene. Since the recombinants of CmICL were more sensitive to tryptic digestion than EcICL (Fig. 5), the poor expression of the CmICLs might be ascribed to their post-translational proteolysis in E. coli cells. On the other hand, Cm-WT showed an obviously lower optimum temperature for activity and thermostability than the mesophilic Ec-WT (Fig. 3), indicating that the sensitivities of Cm-WT, Cm-Q207H, Cm-Q217K and Ec-WT in assay mixtures were calculated as 17.2, 20.1, 20.1 and 16.1 µM, respectively. The ellipticity of each sample was measured by scanning five times between 200 and 250 nm at 1 nm increments.

Two amino acid residues, His184 and Lys194, of EcICL are known to be conserved in many prokaryotic and eukaryotic ICLs, but both of the corresponding residues in CmICL are Gln (Fig. 1a). Lys194 of EcICL should be essential for the catalytic function, because an ‘active-site loop’ including this residue moves flexibly when the enzyme binds to the substrate and releases the reaction products (Britton et al., 2000, 2001; Sharma et al., 2000; cf. Fig. 7). Between 30 °C and 40 °C, Cm-Q207H has a specific activity similar to that of Cm-WT (Fig. 3a, b) and the $K_m$ value of the isocitrate value of the former was lower than that of the latter (Table 1 and Fig. 4a). These results suggest that, in this temperature range, this mutation strengthens the affinity for isocitrate

Fig. 5. Digestion of recombinant ICLs with trypsin. (a) SDS-PAGE: all recombinant ICLs (100 µg) were digested at 4 °C with 0.1 µg trypsin. At indicated times, 10 µg enzyme was withdrawn and subjected immediately to electrophoresis. After electrophoresis, proteins on the gel were stained by Coomassie Brilliant Blue. (b) Densitometric analysis of the amounts of ICL proteins after digestion. Symbols are the same as Fig. 3. Values are expressed as a percentage of the amount of ICL protein at t=0.

Fig. 6. UV-CD spectra of Cm-WT (solid line), Cm-Q207H (---), Cm-Q217K (----) and Ec-WT (--). Based on the tetrameric configurations, the protein concentrations of Cm-WT, Cm-Q207H, Cm-Q217K and Ec-WT in assay mixtures were calculated as 17.2, 20.1, 20.1 and 16.1 µM, respectively. The ellipticity of each sample was measured by scanning five times between 200 and 250 nm at 1 nm increments.
without diminishing the catalytic function. As reported for Ecl (Rehman & McFadden, 1997b), the substitution of the cationic Lys residue for Gln should facilitate the binding to the Mg\(^{2+}\)-isocitrate complex as substrate. On the other hand, the catalytic rates of Cm-Q217K between 10 \(^\circ\)C and 25 \(^\circ\)C were obviously decreased (Figs 3a and 4b), indicating that the Gln residue is important for cold adaptation of CmICL. This may be due to a decrease in the structural flexibility of the mutated enzyme, revealed as an increase in thermostability and a change in the CD spectrum (Figs 3c and 6, Table 1). The high flexibility enhances the accommodation of enzyme with substrate at low temperature, but is also responsible for poor binding to ligand. It has been reported that the \(K_m\) values of cold-adapted enzymes are often higher than those of mesophilic and/or thermophilic counterparts (Fields & Somero, 1998). In fact, between 10 \(^\circ\)C and 30 \(^\circ\)C, the \(K_m\)isocitrate of Cm-WT was larger than those of Ec-WT and Cm-Q217K (Fig. 4a).

Eubacterial and eukaryotic ICLs contain the triose phosphate isomerase (TIM) barrel fold consisting of \((\beta/\alpha)_7\beta\). The molecular modelling of CmICL revealed that the Gln207 of CmICL is present in the barrel fold, and its side chain is directed toward the inner space of the barrel (Fig. 7), in the same way as the His184 of Ecl (Britton et al., 2000, 2001; Sharma et al., 2000). It has been reported that four Ecl mutants, in which His184 was replaced by Leu, Lys, Arg or Gln, are not able to form the tertiary structure, and that only the H184Q mutant shows activity (0-28 % of the wild-type Ecl activity; Diehl & McFadden, 1994). On the other hand, Cm-Q207H showed significant enzyme activity at moderate temperatures between 20 \(^\circ\)C and 30 \(^\circ\)C, while its activities at low temperatures were diminished (Fig. 3a, b), indicating that Gln207 of CmICL is also involved in the cold adaptation of the enzyme. Although this mutation resulted in only a small conformational change of the enzyme protein (Fig. 6), the thermostability and affinity for isocitrate as a substrate were lower than those of Cm-WT (Figs 3c and 4a). These results suggest that Gln207 of CmICL contributes to cold adaptation by a different mechanism from that of the Gln217 residue.

A decrease of \(\Delta H^\#\) (or \(E_a\)) is one of the common strategies for psychrophilic enzymes to weaken the temperature dependence of \(k_{cat}\) (Lonhienne et al., 2000). Furthermore, as seen in the comparison of Cm-WT and Ec-WT, the T\(\Delta S^\#\) values of psychrophilic enzymes are also generally known to be smaller than those of their mesophilic counterparts, because the enhanced flexibility of psychrophilic enzyme proteins allows them to diversify the transition states of intermediate enzyme–substrate complexes during the reaction more than mesophilic ones (Fields & Somero, 1998; Lonhienne et al., 2000). Conversely, such a relationship was not observed in the comparison of Cm-Q207H and Cm-Q217H with Ec-WT (Table 1). Therefore, with regard to their thermodynamic parameters also, both the mutated CmICLs are judged to be inferior to Cm-WT as a cold-adapted enzyme, although they still exhibit a lower optimum temperature for activity and higher thermostability than Ec-WT.

**Fig. 7.** Molecular model of CmICL. A loop region containing the active site is indicated by orange. The side chains of Gln207 and Gln217 are indicated by red.
The phylogenetic analysis revealed that, among the subfamily 3 ICLs from a limited number of eubacteria including *C. maris*, Gln207 and Gln217 of *CnmICL* are conserved, while the putative ICLs of other psychrophilic and psychrotrophic bacteria are classified into subfamily 1, separate from *CnmICL*, and possess His and Lys, identical to EICL, at the corresponding positions (Fig. 1). This implies that the importance of the two Gln residues to cold adaptation is specific to *CnmICL*, and agrees with the finding that more than one pattern of amino acid substitution contributes to the cold and/or thermal adaptation of proteins (Fields & Somero, 1998; Sheridan et al., 2000; Suzuki et al., 2001; Lönn et al., 2002; Gerike et al., 2001). On the other hand, the possibility remains that additional amino acid residue(s), interacting and/or cooperating with the two Gln residues, are involved in the cold adaptation of *CnmICL*. Therefore, further study is in progress to confirm this possibility.

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**REFERENCES**


