Uncoupling the MgATP-induced inhibition and aggregation of *Escherichia coli* phosphofructokinase-2 by C-terminal mutations

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Abstract Binding of MgATP to an allosteric site of *Escherichia coli* phosphofructokinase-2 (Pfk-2) provoked inhibition and a dimer–tetramer (D–T) conversion of the enzyme. Successive deletions of up to 10 residues and point mutations at the C-terminal end led to mutants with elevated $K_{Mapp}$ values for MgATP which failed to show the D–T conversion, but were still inhibited by the nucleotide. Y306 was required for the quaternary packing involved in the D–T conversion and the next residue, L307, was crucial for the ternary packing necessary for the catalytic MgATP-binding site. These results show that the D–T conversion could be uncoupled from the conformational changes that lead to the MgATP-induced allosteric inhibition.

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1. Introduction

Substrate inhibition induced by MgATP is an extended characteristic of phosphofructokinases characterized from different sources. In *Escherichia coli*, two non-homologous phosphofructokinases, Pfk-1 and Pfk-2 [1], have been described to show MgATP-induced substrate inhibition [2–5]. In the case of Pfk-2, this enzymatic inhibition has been related with the presence of an allosteric site for the nucleotide [3], a relevant feature, since *E. coli* strains that express a non-inhibited form of the enzyme presented a diminished growth rate in gluconeogenic media [6].

The quaternary structure of the free form of Pfk-2 has been described as a permanent homodimer whose aggregation state is an obligated conformation necessary for catalysis and stability [7,8]. However, the fact that the presence of MgATP favored the tetrameric form of the enzyme [1,2] prompted the idea that the dimer is the active form of Pfk-2 and the tetramer the inhibited one [9,10]. Nevertheless, the precise role of the tetramer in the inhibition mechanism is still unknown, mainly because of the absence of a model that describes the kinetics of substrate inhibition on the one hand, and the absence of structural support for the biochemical data on the other.

Binding of MgATP to the free enzyme (measured by fluorescence titration) has been interpreted as an allosteric binding [10] because the kinetic mechanism of addition of substrates predicts that MgATP binding to the active site is forbidden in the absence of fructose-6-phosphate (fructose-6-P) [11]. In this regard, a fragment of Pfk-2 obtained by limited proteolysis retained the allosteric-binding capacity, as indicated by fluorescence titration experiments performed in presence of MgATP, but it was not able to form tetramers [12]. Since the Pfk-2 fragment was characterized as a ~28 kDa polypeptide with an intact N-terminal sequence, it was suggested that residues from C-terminal region are important for the MgATP-induced tetramerization [12]. However, the relationship between tetramer formation and the MgATP-induced inhibition of the enzyme could not be addressed since the Pfk-2 fragment was inactive.

Pfk-2 belongs to the ribokinase superfamily [13], an extensive family of kinases that phosphorylates a broad range of vitamins and sugars [14]. In this family, the active sites are mostly contained in a structurally conserved $\alpha/\beta/\alpha$-fold and the structural determinants for the nucleoside phosphate donors binding. ATP or ADP, are located at the C-terminal region of this fold [14,15]. Thus, besides the Pfk-2 inactivation provoked by the proteolytic digestion, structural alterations of the catalytic MgATP-binding site can be predicted to occur upon limited proteolysis of the C-terminal end of enzyme.

In this work, we have performed successive C-terminal deletions and point mutations in Pfk-2 in order to characterize the structural determinants of the catalytic ATP-binding site and the structural determinants of the MgATP-induced dimer–tetramer conversion, whose localization at the C-terminal region is predicted by the limited proteolysis experiments mentioned above. The C-terminal mutants were evaluated with respect to their kinetic parameters, sensitivity towards MgATP inhibition and aggregation state. Finally, the results are discussed in the light of the crystal structure of Pfk-2 obtained recently by our group [PDB ID, 3CQD].

2. Materials and methods

2.1. Site-directed mutagenesis; mutants’ expression and purification

Site-directed mutagenesis was carried out using the GeneTailor™ Site-Directed Mutagenesis System (Invitrogen) using a pET21d plasmid (Novagen) containing the *pfk-2* gene as template. Oligonucleotides encoding stop codons were used to introduce new translational stop

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Abbreviations: Pfk-2, phosphofructokinase-2; $R_h$, hydrodynamic radius; $K_{Mapp}$, apparent Michaelis–Menten constant; $k_{cat}$, catalytic constant; fructose-6-P, fructose-6-phosphate
sites at the appropriate positions in the C-terminal coding region. The mutations were verified by DNA sequencing of both strands. The mutant Pfk-2 enzymes were produced in E. coli Bl21 (DE3). Wild-type and mutant enzymes were purified as described by Parducci et al. [16]. However, the D299 mutant could not be purified because of its poor binding to the Cibacron Blue-Sepharose column. In this case, the overexpressed mutant represents about 80% of the total protein, so its \(k_{cat}\) values were corrected accordingly.

2.2. Size exclusion chromatography

Experiments were performed with a Waters 1525 HPLC binary pump system, with a Bio-Sil SEC-250 (7.8 mm × 30 cm) column (Bio-Rad, Hercules, CA, USA) equilibrated in a buffer containing 25 mM Tris–HCl, pH 7.6, 200 mM KCl and 1 mM dithiotreitol at the indicated Mg\(^{2+}\), ATP and fructose-6-P concentrations. Protein elution was recorded with the use of an online Waters 2487 UV dual detector. All runs were carried out at a flow rate of 0.8 ml/min. The column was calibrated with the following molecular-mass markers: Vitamin B-12 (1.35 kDa, 8.5 Å hydrodynamic radius, \(R_h\)), horse myoglobin (17 kDa, 19 Å \(R_h\)), chicken ovalbumin (44 kDa, 30.5 Å \(R_h\)), bovine gamma globulin (158 kDa, 41.8 Å \(R_h\)) and bovine thyroglobulin (670 kDa, 85 Å \(R_h\)). Protein elution volumes were converted to \(R_h\) values using the linear relationship obtained with the molecular-mass markers,\[ R_h = 51.8 \sqrt{-\log Kav} - 9.5 \]

\[ Kav = \frac{V_e - V_o}{V_t - V_o} \]

where \(Kav\) represents the partition constant calculated from the elution volume (\(V_e\)), the void volume (\(V_o\)) and the total volume of the column (\(V_t\)) [17].

2.3. Enzyme assay

Phosphofructokinase activity was determined spectrophotometrically by coupling the fructose-1,6-bisphosphate production to the oxidation of NADH [1]. The assay was slightly modified to avoid a kinetic lag observed at the initial time of the measurements. Decreasing the Tris–HCl concentration from 100 to 25 mM together the omission of NH\(^+\)\(_4\) salt from the assay mixture, eliminates the kinetic lag that complicates the initial velocity measurements at low substrate concentrations, but does not modify the regulatory properties and the \(k_{cat}\) values of Pfk-2. The apparent \(K_M\) value (\(K_{Mapp}\)) for MgATP was calculated by using an uncompetitive substrate inhibition function, and the \(K_{Mapp}\) for fructose-6-P by using a non-linear curve fit to a hyperbolic function. The fit procedure was performed by a non-linear regression tool provided by the Sigmaplot package 9.0 (SYSTAT Software Inc).

3. Results

3.1. Dimer–tetramer conversion induced by MgATP and kinetic characterization of the mutants generated by successive removal of residues from the C-terminal end

Four mutants with 2–10 residues deletions from the Pfk-2 C-terminal end (Fig. 1) were characterized with respect to their apparent \(K_M\) values for fructose-6-P, MgATP and their catalytic constants \(k_{cat}\). Compared to the wild-type enzyme, all mutants, with the exception L307, presented increased was recorded with the use of an online Waters 2487 UV dual detector. All runs were carried out at a flow rate of 0.8 ml/min. The column was calibrated with the following molecular-mass markers: Vitamin B-12 (1.35 kDa, 8.5 Å hydrodynamic radius, \(R_h\)), horse myoglobin (17 kDa, 19 Å \(R_h\)), chicken ovalbumin (44 kDa, 30.5 Å \(R_h\)), bovine gamma globulin (158 kDa, 41.8 Å \(R_h\)) and bovine thyroglobulin (670 kDa, 85 Å \(R_h\)). Protein elution volumes were converted to \(R_h\) values using the linear relationship obtained with the molecular-mass markers,

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\[ \text{WT} \quad \text{L307} \quad \text{Y306} \quad \text{A305} \quad \text{D299} \]

| \(K_{Mapp}\) fructose-6-P (\(\mu M\))^a\b | 57 ± 3 | 60 | 45 | 36 | 20 |
| \(K_{Mapp}\) MgATP (\(\mu M\))^a | 15 ± 3 | 10 | 880 | 800 | 2000 |
| \(k_{cat}\) (s\(^{-1}\))^c | 58 ± 3 | 46 | 38 | 35 | 4 |

\(a\) Determinations performed with 1 mM co-substrate.

\(b\) The concentration of the co-substrate was chosen in order avoid the effect of MgATP binding at the allosteric site to the binding of fructose-6-P to the catalytic site, which is observed above 1 mM MgATP (unpublished observations).

\(c\) Determinations performed as a function of the MgATP concentration with 1 mM fructose-6-P.

Table 1 Kinetic parameters measured for the wild-type enzyme and the C-terminal deletion mutants

![Fig. 1. Successive C-terminal residue deletions of Pfk-2. The C-terminal end \(\alpha\)-helix of Pfk-2 is represented with the primary sequence expected for each mutant used in this study.](image)

![Fig. 2. Activity of the Y306 and A305 mutants as a function of the MgATP concentration. Each panel shows the initial velocity measured as a function of the MgATP concentration using a fixed concentration of 2 mM (■), 0.2 mM (○) and 0.05 mM (■) fructose-6-P. Y306 mutant (panel A) and A305 mutant (panel B).](image)
Deletion of three residues (mutant Y306) resulted in a ~1.5 orders of magnitude increase of the $K_{M_{\text{app}}}$ value for MgATP, with respect to the L307 mutant and the wild-type enzyme. The elimination of L307 seems to be necessary and sufficient to provoke a large alteration in the pseudo-affinity for MgATP, since subsequent deletion of four residues (mutant A305) and 10 residues (mutant D299) lead to a progressive increase, from 1.5 to 1.8 orders of magnitude, of the $K_{M_{\text{app}}}$ value for MgATP (Table 1). Among the characterized mutants, only Y306 and A305 show a specific alteration in their $K_{M_{\text{app}}}$ value for MgATP since both mutants retain about the 80% of the wild-type activity without variations of the $K_{M_{\text{app}}}$ for fructose-6-P (Table 1). Further elimination of up to 10 residues provokes a decrease of the enzymatic activity to 10% of the value of the wild-type enzyme (Table 1).

Substrate inhibition induced by MgATP is a property of the wild-type enzyme that has been related with the presence of an allosteric site that negatively regulates the enzymatic activity of Pfk-2 [3,10]. Fig. 2 shows the initial velocity of mutants Y306 and A305 measured as a function of the MgATP concentration by using several fixed concentrations of fructose-6-P. As can be seen, the enzymatic activity goes through a maximum value and then decreases at elevated concentrations of MgATP, indicating that the allosteric inhibition induced by MgATP is still operative, although the $K_{M_{\text{app}}}$ value for the nucleotide of both mutants was about 1.5 orders of magnitude higher than the wild-type enzyme.

To determine the effect of the C-terminal deletions in the MgATP-induced tetramer formation, size exclusion chromatography was performed in the presence of 0.2 mM MgATP. As shown in Fig. 3, with the exception of the L307 mutant that presents a wild-type phenotype, neither mutant modified significantly its $R_h$ in the presence of 0.2 mM MgATP, suggesting that the structural alterations that resulted in an elevated $K_{M_{\text{app}}}$ for MgATP are linked with the tetramer formation induced by MgATP binding at the allosteric site. In absence of MgATP the hydrodynamic radiiuses of the C-terminal mutants (Y306, A305 and D299) show a slight increment, between 0.6 and 1 Å with respect the value observed for the wild-type dimer (see Fig. 3). Although the observed size increments could be due to alterations in the shape of the mutants or to increments in the tetramer populations, the latter seems unlikely since the dimer–tetramer conversion does not take place in the wild-type enzyme in absence of MgATP [8].

The impediment of those mutants to form tetramers in presence of MgATP, as Y306 and A305, seems not to be related with the enzymatic inhibition induced by MgATP, since both mutants show this feature when the initial velocity is assayed at elevated concentrations of the nucleotide (Fig. 2). In order to study the failure in tetramer formation in a kinetic background closer to the wild-type, the side chains of L307 and Y306 were replaced by alanine to generate the point mutants Y306A and L307A.

### 3.2. Dimer–tetramer conversion induced by MgATP and kinetic properties of the Y306A and L307A mutants

The MgATP-induced inhibition of Pfk-2 has been described as a biphasic behavior of the initial velocity, measured as function of the MgATP concentration, when the assay was performed at low fructose-6-P concentrations (0.1 mM), which tended to become hyperbolic at elevated concentrations (1 mM) [2,3]. Fig. 4 shows the initial velocity as a function of the MgATP concentration for the wild-type enzyme and the Y306A and L307A mutants using two fixed concentrations of fructose-6-P. At 1 mM fructose-6-P (Fig. 4A), the $K_{M_{\text{app}}}$ for MgATP for the L307A mutant was about 1.5 orders of magnitude higher than the wild-type enzyme while the $k_{\text{cat}}$ and $K_{M_{\text{app}}}$ for fructose-6-P remain essentially unchanged (Table 2). At 0.1 mM fructose-6-P (Fig. 4B) the MgATP-induced inhibition is favored since a decrease of the initial velocity is observed at elevated concentrations of the nucleotide. As shown, the L307A mutant presents its activity maximum at higher MgATP concentrations compared to the wild-type.
enzyme, probably because of its lower pseudo-affinity for MgATP; the inhibition is still observed although at higher concentrations of the nucleotide. As pointed out above, the C-terminal deletion mutants, characterized by elevated $K_{\text{Mapp}}$ values for MgATP, were unable to form tetramers in presence of MgATP. The same situation is observed with the L307A mutant when injected to a size exclusion column in the presence of 0.2 mM MgATP (data not shown). Thus, the origin of the conformational change that leads to an increase in the $K_{\text{Mapp}}$ value for MgATP and the impairment in tetramer formation can be mapped to the side chain of L307.

On the other hand, the change of Y306 by alanine does not modify the $K_{\text{Mapp}}$ value for MgATP and the other kinetic parameters compared to the wild-type enzyme (Fig. 4A and Table 2). This result seems reasonable since the L307 side chain is present in the protein. However, size exclusion chromatography performed with the Y306A mutant did not show the presence of a tetrameric species even at 0.8 mM MgATP (data not shown). Thus, the origin of the conformational change that leads to an increase in the $K_{\text{Mapp}}$ value for MgATP and the impairment in tetramer formation can be mapped to the side chain of L307.

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### Table 2
Kinetic parameters measured for the wild-type enzyme and the Y306A and L307A mutants

<table>
<thead>
<tr>
<th></th>
<th>Wild-type</th>
<th>Y306A</th>
<th>L307A$^b$</th>
</tr>
</thead>
<tbody>
<tr>
<td>$K_{\text{Mapp}}$ fructose-6-P ($\mu$M)$^a$</td>
<td>57 ± 3</td>
<td>62 ± 11</td>
<td>29</td>
</tr>
<tr>
<td>$K_{\text{Mapp}}$ MgATP ($\mu$M)$^b$</td>
<td>15 ± 3</td>
<td>13.7 ± 0.9</td>
<td>954</td>
</tr>
<tr>
<td>$k_{\text{cat}}$ (s$^{-1}$)$^c$</td>
<td>58 ± 3</td>
<td>53 ± 3.7</td>
<td>59</td>
</tr>
</tbody>
</table>

$^a$Determinations performed with 1 mM co-substrate.

$^b$Average of two measurements.

$^c$Determinations performed as a function of the MgATP concentration with 1 mM fructose-6-P.

### 4. Discussion

#### 4.1. Effect of the successive C-terminal deletions on the kinetic parameters of Pfk-2

Despite of the low sequence identity among the members of the ribokinase superfamily, the nucleotide-binding site is a structurally conserved feature [14,15]. Specifically, the adenine ring of the phosphate donor appears into a hydrophobic pocket created by two loops, the small and large ATP-binding loops, in the ribokinase fold [18]. Fig. 6 shows the C-terminal portion of the crystallographic structure of Pfk-2 where the last segment of eleven residues form an amphipathic $\alpha$-helix whose hydrophobic side chains are in close contacts with the large ATP-binding loop atoms and with a structurally con-
served α-helix that participates in the network of the ATP-binding site in the ribokinase superfamily members. Particularly, L307 is the last hydrophobic residue of the amphipathic α-helix whose apolar side chain is buried with atoms of the large ATP loop and of the conserved α-helix (Fig. 6). This feature correlates well with the abrupt increase of 1.5 orders of magnitude of the $K_{M,app}$ value for MgATP that occurs upon deletion of a three residue segment that includes L307 (mutant Y306) or by a point mutation that replaces its side chain by alanine (mutant L307A). It is known that cavities created by substituting apolar buried side chains by apolar smaller ones, provoke structural rearrangements driven by an unfavorable loss of contacts in hydrophobic cores [19]. The structural rearrangements induced by “cavity-creating” mutants, Y306 and L307A, seems to affect mostly the tertiary packing around the ATP-binding site since, besides the large increase in the $K_{M,app}$ value for MgATP, no significant changes were observed in the $K_{M,app}$ for fructose-6-P or $k_{cat}$ values in these mutants.

Chinese hamster adenosine kinase is a monomeric member of the ribokinase superfamily that shares a 12% sequence identity with Pfk-2, but presents a good alpha carbons structural superposition (1.7 Å R.M.S). Maj and coworkers [20] have performed successive deletions of the C-terminal region of adenosine kinase reporting a gradual decrease of the $k_{cat}$ value together with a progressive increase of the $K_{M,app}$ for MgATP, while the $K_{M,app}$ value for the adenosine phosphate acceptor was unaltered, independent of number of deleted residues. Likewise, the $K_{M,app}$ of the phosphate acceptor of Pfk-2 (fructose-6-P) is maintained with the number of deleted residues, suggesting that conformational changes at the C-terminal region of the ATP-binding site can be uncoupled from the phosphate acceptor-binding site in the ribokinase fold. However, in contrast with the abrupt increase of 1.5 orders of magnitude observed upon deletion of L307 in Pfk-2, in adenosine kinase the $K_{M,app}$ value for MgATP suffers a gradual increase with the number of deleted residues. The superposition of both structures shows that the deleted residues, in the case of adenosine kinase, form a loop whose side chains do not show direct contacts with atoms involved in the ATP-binding site [21]. Conversely, the hydrophobic side chains of the amphipathic α-helix in Pfk-2 are packed against atoms of the ATP loop and with a helix that forms part of the adenosine-binding moiety (Fig. 6A).
4.2. Effects of the successive C-terminal deletions on the MgATP-induced tetramer formation

A striking characteristic of the C-terminal mutants is that, besides their elevated $K_{M}^{app}$ for MgATP, they present impairment in the MgATP-induced tetramer formation. The biologic unit of Pfk-2 present into the crystal structure obtained in the presence of MgATP was assumed to be a homotetramer created by a surface of interaction that hides an extensive area of 2154 Å² for each dimer (unpublished data). Despite of this extensive contact area, only 4 side chains are expected to decrease the free energy of subunit interaction ($\Delta G_{bind}$) by at least 1.2 kcal/mol upon alanine replacement. This value was predicted by using an in silico alanine scanning method [22] available in the Web (http://robetta.bakerlab.org/alaninescan). Between the selected side chains, Y306 presents a calculated $\Delta G_{bind}$ of 4.4 kcal/mol (1.15 kcal/mol per subunit). As shown in Fig. 6, Y306 is the only modified residue in this work whose side chain makes contact with the adjacent homodimer. This suggests that the same conformational change that is responsible for the increased $K_{M}^{app}$ value for MgATP, upon replacement or deletion of L307, could modify the correct orientation of the Y306 side chain, which in turn could explain the failure of the dimer–tetramer conversion observed in the “cavity-creating” mutants. In agreement with this idea, the replacement of Y306 by alanine generates a mutant unable to form tetramers, but with a kinetic behavior almost invariant with respect to the wild-type enzyme. The importance of this result is the indication that conformational changes that lead to allosteric inhibition can be uncoupled from tetramer formation at least in the case of the Y306A mutant. For the wild-type enzyme, Cabrera and coworkers [23] reported that, besides tetramer formation, the presence of MgATP promotes subunit conformational changes, defined as rigid movements between the mayor and the minor domains, which result in the closure of the active site of each subunit. Therefore, taking into account that in presence of MgATP the mutant Y306A does not show the dimer–tetramer conversion but presents an inhibition pattern almost indistinguishable from the wild-type enzyme, the enzymatic inhibition induced by MgATP can be explained exclusively by conformational changes at the dimeric or monomeric level of Pfk-2. Further experiments are required to determine the biologic relevance of the dimer–tetramer equilibrium observed in presence of MgATP.

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