Structure of the Protein Phosphatase 2A Holoenzyme

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DOI 10.1016/j.cell.2006.11.033

SUMMARY

Protein Phosphatase 2A (PP2A) plays an essential role in many aspects of cellular physiology. The PP2A holoenzyme consists of a heterodimeric core enzyme, which comprises a scaffolding subunit and a catalytic subunit, and a variable regulatory subunit. Here we report the crystal structure of the heterotrimeric PP2A holoenzyme involving the regulatory subunit B0/PR61. Surprisingly, the B0/PR61 subunit has a HEAT-like (huntingtin-elongation-A subunit-TOR-like) repeat structure, similar to that of the scaffolding subunit. The regulatory B0/PR61 subunit simultaneously interacts with the catalytic subunit as well as the conserved ridge of the scaffolding subunit. The carboxyterminus of the catalytic subunit recognizes a surface groove at the interface between the B0/PR61 subunit and the scaffolding subunit. Compared to the scaffolding subunit in the PP2A core enzyme, formation of the holoenzyme forces the scaffolding subunit to undergo pronounced conformational rearrangements. This structure reveals significant ramifications for understanding the function and regulation of PP2A.

INTRODUCTION

Controlled protein phosphorylation and dephosphorylation are a fundamental regulatory mechanism in all aspects of biology (Hunter, 1995). Protein phosphatase 2A (PP2A) is a major serine/threonine phosphatase involved in many essential aspects of cellular function (Janssens and Goris, 2001; Virshup, 2000). PP2A plays an important role in cell-cycle regulation, cell-growth control, development, regulation of multiple signal transduction pathways, cytoskeleton dynamics, and cell mobility.

The PP2A core enzyme consists of a 65 kDa scaffolding protein known as A or PR65 subunit and a 36 kDa catalytic subunit or C subunit. To gain full activity towards specific substrates, the PP2A core enzyme associates with a variable regulatory subunit to form a heterotrimeric holoenzyme. The variable regulatory subunits have four subfamilies: B (PR55), B′ (B56 or PR61), B″ (PR72), and B‴ (PR93/PR110), with at least 16 members in these subfamilies (Janssens and Goris, 2001; Lechward et al., 2001). In mammalian cells, the scaffolding and the catalytic subunits each have two isoforms, which share high sequence similarity (Arino et al., 1988; Green et al., 1987; Hemmings et al., 1990; Stone et al., 1987). In contrast, the sequence similarity among the four subfamilies of regulatory subunits is very low, and the expression levels of various regulatory subunits are highly diverse depending upon cell types and tissues (Janssens and Goris, 2001; Lechward et al., 2001). In this regard, the B subunits determine the substrate specificity as well as the spatial and temporal functions of PP2A.

PP2A is also an important tumor suppressor protein (Janssens et al., 2005). Mutations in the scaffolding subunit that result in compromised binding to the regulatory or catalytic subunit of PP2A, or a total absence or substantial reduction of the scaffolding subunit, had been reported to be linked to a variety of primary human tumors (Calin et al., 2000; Colella et al., 2001; Ruediger et al., 2001a; Suzuki and Takahashi, 2003; Takagi et al., 2000; Wang et al., 1998). In addition, an amino-terminally truncated form of the B subunit PR61/B′γ1 was found to be associated with a higher metastatic state of melanoma cells (Ito et al., 2000, 2003; Koma et al., 2004), while gain-and loss-of-function experiments firmly established this regulatory subunit as a tumor suppressor (Chen et al., 2004).

The PP2A scaffolding subunit contains 15 tandem repeats of a conserved 39-residue sequence known as a HEAT (huntingtin-elongation-A subunit-TOR) motif (Hemmings et al., 1990; Walter et al., 1989). These 15 HEAT repeats are organized into an extended, L-shaped molecule (Groves et al., 1999). The catalytic subunit recognizes one end of the elongated scaffolding subunit by interacting with the conserved ridges of HEAT repeats 11–15 (Ruediger et al., 1992, 1994; Xing et al., 2006). Formation of the PP2A core enzyme results in significant bending of the HEAT repeats 12–15 towards the amino terminus of the scaffolding subunit (Xing et al., 2006).
Despite recent advances, mechanistic understanding on the function and assembly of the PP2A holoenzyme has been slow to emerge. There is no structural information on any of the regulatory subunits, and it remains unclear how the regulatory subunit specifically recognizes the scaffolding subunit or the catalytic subunit. Methylation of the carboxyterminus of the catalytic subunit is thought to play an important role in the formation of a holoenzyme (Bryant et al., 1999; Kloeker et al., 1997; Tolstykh et al., 2000; Wei et al., 2001; Wu et al., 2000; Xing et al., 2006; Yu et al., 2001), although the molecular basis underlying these observations remains unclear.

In this manuscript, we report the crystal structure of a PP2A holoenzyme, involving the α isoform of the scaffolding subunit (αz), the β isoform of the catalytic subunit (Cz), and the γ1 isoform of the regulatory B/B56/PR61 subunit (B-γ1). This crystal structure reveals a number of unexpected findings that have significant biological implications for understanding the mechanisms and cellular functions of PP2A.

RESULTS

Assembly and Crystallization of the PP2A Holoenzyme

The PP2A core enzyme, involving the full-length αz and Cz, was assembled as described (Xing et al., 2006). The PP2A core enzyme was methylated by a PP2A-specific leucine carboxyl methyltransferase (LCMT) in the presence of S-adenosyl methionine (SAM). The fully methylated PP2A core enzyme was incubated with a stoichiometric amount of the full-length B-γ1. The heterotrimeric PP2A holoenzyme was purified to homogeneity by gel-filtration chromatography.

The PP2A holoenzyme was incubated with 1.2 molar equivalence of microcystin-LR (MCLR) prior to crystallization. We succeeded in obtaining small crystals of the PP2A holoenzyme and were able to generate large, diffraction-quality crystals using macroseeding. These crystals were sensitive to radiation damage and diffracted X-rays to about 3.3 Å resolution at synchrotron. The structure was determined by a combination of molecular replacement and single-wavelength anomalous dispersion. The atomic model of the native holoenzyme has been refined to 3.3 Å resolution (Table 1).

Overall Structure of the PP2A Holoenzyme

The structure of the PP2A holoenzyme exhibits a compact architecture, measuring 90 × 90 × 70 Å3 (Figure 1). αz has an extensive, C-shaped structure characterized by double-layered α helices. Compared to the free form (Groves et al., 1999) and the PP2A core enzyme (Xing et al., 2006), αz has undergone significant conformational rearrangements (discussed in detail later). There are 15 HEAT repeats in αz, with each HEAT repeat comprising a pair of antiparallel α helices. The interhelical region within each HEAT repeat forms a contiguous ridge—hereafter referred to as the ridge. As previously shown (Xing et al., 2006), Cz binds to one end of the scaffold through interactions with the ridge of HEAT repeats 11–15 (Figure 1), consistent with an earlier prediction (Ruediger et al., 1992). Interactions between αz and Cz result in the burial of 2072 Å² exposed surface area.

The regulatory B/PR61-γ1 subunit has an elongated, superhelical structure comprising 18 α helices (Figure 1A). The superhelical structure has an apparent curvature, with the convex side binding to αz and the concave side facing up and tilted towards Cz. Strikingly, despite a complete lack of sequence homology, the structure of B-γ1 closely resembles that of αz. There are eight HEAT-like repeat motifs in B-γ1.

B-γ1 makes extensive interactions with both αz and Cz (Figure 1). On one hand, B-γ1 binds to the conserved ridge of HEAT repeats 2–8 in αz. On the other hand, B-γ1 makes direct interactions with three surface areas of Czα, including helix ∆φ and the carboxyterminus. A total surface area of 4300 Å² is buried as a result of interaction between the regulatory B-γ1 subunit and the PP2A core enzyme. Approximately 55% of the buried surface area is due to interactions between B-γ1 and Czα, with the rest coming from interactions between B-γ1 and αz. Interestingly, the carboxyterminus of Czα recognizes a surface groove at the interface between B-γ1 and αz (Figure 1B).

Structure of the Regulatory B/PR61 Subunit

The regulatory B/PR61-γ1 subunit consists exclusively of α helices (Figures 2 and 3). Eighteen α helices stack against each other laterally to create an elongated, superhelical structure. The concave side of the curved superhelical structure is enriched with negatively charged amino acids, whereas the convex side contains a number of conserved hydrophobic residues (Figure 2A). These hydrophobic amino acids make important contributions in binding to the scaffolding subunit αz.

One unanticipated finding from the structural investigation is that the structure of B-γ1 resembles that of αz. Systematic structure-based search of the Protein Data Bank using the DALI server (Holm and Sander, 1993) led to the identification of a number of close structural homologs, among which the nuclear import factor karyopherin α (Conti et al., 1998), the armadillo repeat protein β-catenin (Huber et al., 1997), and αz of PP2A (Groves et al., 1999) have the highest Z scores (similarity score) (Figure 2B). Intriguingly, both karyopherin α and β-catenin have the ability to interact with peptides using a concave surface (Conti et al., 1998; Graham et al., 2000). This analysis suggests the possibility that the regulatory B/PR61 subunit may interact with PP2A substrates or other regulatory proteins using the acidic, concave surface (Figure 2B).

Structure of B-γ1 can be superimposed with that of αz with a root-mean-square deviation (rmsd) of 5.1 Å over 254 aligned backbone alpha carbon atoms. This alignment identified 8 HEAT-like repeats in B-γ1. Unlike the HEAT repeats in αz or the armadillo motifs in β-catenin, these
HEAT-like motifs exhibit little sequence homology and appear to lack a strong consensus sequence (Figure 3). Nonetheless, these motifs can be superimposed with each other with 0.9–2.5 Å pairwise rmsd (Figures 2C and 3). HEAT-like motif 7 of B₀–g₁ is most homologous to the HEAT repeat of Aα. For the ease of discussion, the HEAT-like motifs of B₀–g₁ are hereafter referred to as HEAT repeats.

### Interface between the Catalytic and the Regulatory Subunits

The interface between the catalytic subunit and the regulatory B₀/PR61 subunit is extensive and consists of two major areas (Figure 4A). In one area, amino acids from HEAT repeats 6–8 of B₀–γ₁ interact with residues in the a5 region of Cα, mainly through hydrogen bonds (Figure 4B). At the center of this interface, the polar side chain of Gln125 in Cα makes a pair of hydrogen bonds to the main chain atoms in B₀–g₁. The aliphatic portion of Gln125 side chain also mediates van der Waals contact with His339, Phe340, and Trp382 in B₀–g₁. These interactions are further buttressed by three hydrogen bonds between the side chain of Asp131 in Cα and the amide nitrogen and side chain of Ser298 in B₀–g₁ (Figure 4B).

In the other area, the hydrophobic carboxyl terminus of Cα is nestled in a surface groove at the interface between Aα and B₀–g₁ (Figure 4C). There are numerous van der Waals contacts at this interface. Three residues from Cα, Pro305, Tyr307, and Phe308, stack closely against each other and are surrounded by the aliphatic portion of the

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Table 1. Statistics from Crystallographic Analysis

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<th>Protein components</th>
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<td>100–3.8 (3.94–3.80)</td>
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<td>0.116 (0.446)</td>
<td>0.109 (0.432)</td>
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</table>

### Refinement

| Resolution (outer shell) | 100–3.3 (3.42–3.30)         | 100–3.6 (3.73–3.60) | 100–3.8 (3.94–3.80) |
| Number of reflections (| 70,199                      | 53,472            | 46,983             |
| Data coverage           | 98.4%                       | 97.4%             | 99.6%              |
| R_work (outer shell)    | 0.252 (0.392)               | 0.267 (0.364)     | 0.282 (0.332)      |
| R_free (outer shell)    | 0.297 (0.434)               | 0.331 (0.439)     | 0.335 (0.418)      |
| Total number of atoms   | 20,474                      | 20,212            | 20,212             |
| Number of waters        | 0                           | 0                 | 0                 |
| Rmsd bond length (Å)    | 0.0083                      | 0.010             | 0.011              |
| Rmsd bond angles (degree) | 1.38                        | 1.53              | 1.56              |

### Average B factor

| Az—two copies per a.u.  | 69.6, 60.1                   | 70.3, 48.1        | 81.3, 67.3         |
| Cα—two copies per a.u.  | 72.7, 70.1                   | 69.2, 59.5        | 80.3, 76.4         |
| B₀–γ₁—two copies per a.u.| 85.8, 79.1                   | 84.3, 70.5        | 95.2, 86.3         |

### Ramachandran Plot

| Most favored (%) | 80.6 | 75.3 | 73.2 |
| Additionally allowed (%) | 17.6 | 22.5 | 23.8 |
| Generously allowed (%) | 1.4  | 2.0  | 2.5  |
| Disallowed (%) | 0.3  | 0.3  | 0.4  |

R_sym = (Σ|I_h,i|/ΣP_h,i) / (ΣP_h,i), where I_h,i is the mean intensity of the i observations of symmetry-related reflections of h. R = Σ|Fo — Fc| / ΣFo, where Fc is the calculated protein structure factor from the atomic model (R_free was calculated with 5% of the reflections). Rmsd in bond lengths and angles are the deviations from ideal values. In protein components, Cα(T) represents the trypsin-digested Cα (residues 1–294), which has the C-terminal 15 amino acids removed.
side chains of Lys256, Lys 258, Tyr292, and Lys295 in B'-γ1 and Asp63, Glu64, and Glu101 of Az (Figure 4C). These van der Waals interactions are strengthened by six additional hydrogen bonds, one of which occurs between the side chain of Glu64 in Az and the main chain amide nitrogen of residue 309 in Cz (Figure 4C). Importantly, mutation of Glu64 to Asp or Gly in Az has been observed in cancer cells (Ruediger et al., 2001b). Supporting an important role for Glu64, the mutant proteins E64D and E64G exhibited significantly compromised ability to interact with the B'/PR61 subunit (Ruediger et al., 2001b). Our structural observation provides a rational basis for the mutagenesis result and for the cancer link. Although the carboxylate of Leu309 is fully methylated as judged by Figure 1. Overall Structure of the Hetro-trimeric PP2A Holoenzyme Bound to Microcystin-LR (MCLR)

(A) Overall structure of the PP2A holoenzyme bound to MCLR. The scaffolding (Aα), catalytic (Cα), and regulatory B'/PR61 (B'-γ1) subunits are shown in green, blue, and yellow, respectively. MCLR is shown in magenta. B'-γ1 interacts with both Aα and Cα through extensive interfaces. Cz interacts with Aα as described (Xing et al., 2006). Three views are shown here to reveal the essential features of the holoenzyme. Surprisingly, B'-γ1 adopts a structure that closely resembles that of the scaffolding subunit (discussed in detail later).

(B) A surface representation of the PP2A holoenzyme. Aα and B'-γ1 are shown in surface representation. Cα is shown in backbone worm to highlight the observation that the carboxyl terminus of Cα binds to a surface groove at the interface between Aα and B'-γ1. Figures 1A, 2A, 4A, and 4E were prepared using GRASP (Nicholls et al., 1991); all other structural figures were made using MOLSCRIPT (Kraulis, 1991).

Figure 2. Structure of the Regulatory B'/PR61 Subunit

(A) A surface representation of the structure of B'-γ1 showing electrostatic potential. Three perpendicular views are shown here. Note that the elongated B'-γ1 protein contains a highly acidic, concave surface patch (top panel). B'-γ1 uses its less-charged, convex surface to interact with Aα.

(B) The regulatory B'/PR61 subunit is structurally similar to the scaffolding subunit of PP2A and other binding proteins. B'-γ1 (yellow) contains eight HEAT repeat-like structural motifs. These HEAT-like motifs can be superimposed well with the scaffolding subunit of PP2A (left panel) or karyopherin α (importin α, right panel). The peptide binding region of karyopherin α coincides with the acidic, concave surface of B'-γ1.

(C) Structural overlay of all eight HEAT-like motifs in the regulatory B'/PR61 subunit. HEAT-like motif 7 is most similar to the HEAT repeat of the PP2A scaffolding subunit. The majority of the HEAT-like motifs have helices almost parallel with each other.
Western blot, we chose not to build the methyl group, because the electron density for the methyl group is weak. Nonetheless, the binding groove for the carboxyl terminus of Ca is very acidic (Figure 4A); methylation of the carboxyl terminus of Ca removes a negative charge and presumably facilitates its docking into the acidic groove.

In addition to the two major interfaces, the extended loop within HEAT repeat 2 of B0-g makes additional interactions with Ca. Arg268 of Ca, which mediates van der Waals contacts with MCLR, donates three hydrogen bonds to residues in B0-g: two to the main chain carbonyl oxygen atoms of residues 119 and 122 and one to the side chain of Asp123 (Figure 4D).

Figure 3. Sequence Alignment of the Regulatory B/PR61 Subunits across Species

Secondary structural elements are indicated above the sequences. Each HEAT repeat is represented by a distinct color. Conserved residues are highlighted in yellow. Residues that interact with Aα and Ca are identified by green and blue circles, respectively, above the sequences. The sequences shown include all five isoforms of B/PR61 from humans: alpha (GI:18490281), beta (GI:30795206), gamma1 (GI:1418819), delta (GI:31083279), and epsilon (GI:31083295). The other sequences are the following: Saccharomyces cerevisiae Rts1p (GI:532526), Chizosaccharomyces pombe CpB (GI:19075706), Caenorhabditis elegans CeB (GI:17557914), Drosophila melanogaster DmB (GI:60777747), and Arabidopsis thaliana AtB (GI:2244898). Residues 626–651 were not shown in this alignment.

In addition to the two major interfaces, the extended loop within HEAT repeat 2 of B-γ1 makes additional interactions with Ca. Arg268 of Ca, which mediates van der Waals contacts with MCLR, donates three hydrogen bonds to residues in B-γ1: two to the main chain carbonyl oxygen atoms of residues 119 and 122 and one to the side chain of Asp123 (Figure 4D).
Interface between the Regulatory and the Scaffolding Subunits

B₀⁻γ1 uses its convex surface to sit on the ridge of HEAT repeats 2–8 in Ax (Figure 4E). Although the buried surface area is large, the intensity of interaction does not appear to be high. The interface between B₀⁻γ1 and Ax is scattered into a large area and a small area. The large area involves HEAT repeats 4 and 5 of B₀⁻γ1 and the ridge of HEAT repeats 2–5 of Ax (Figure 4F). Trp140 and Phe141 of Ax make multiple van der Waals interactions with...
hydrophobic residues in B'-γ1. In addition, Arg183 of Aγ donates a pair of charge-stabilized hydrogen bonds to Glu214 of B'-γ1 (Figure 4F). Lys256 of B'-γ1 forms a salt bridge with three acidic residues in Aα, Asp63, Glu100, and Glu101 (Figure 4F). The small area involves the extended loop in HEAT repeat 2 of B'-γ1 and the ridge of HEAT repeats 7 and 8, where Trp257 of Aγ hydrogen bonds to main chain carbonyl oxygen of residue 107 in B'-γ1 (Figure 4F).

All residues in B'-γ1 that mediate important contacts with residues in Cα and Aα are highly conserved. For example, Glu214, which accepts two hydrogen bonds from Arg183 of Aγ (Figure 4F), and Asp123, which hydrogen bonds to Arg268 of Cα (Figure 4D), are invariant in all B'/PR61 family members across species (Figure 3). The other three B'-γ1 residues that contribute to hydrogen bonds at the interfaces, Lys295, Ser298, and His339, are conserved in all but one B'/PR61 family members (Figure 3). This analysis indicates that the binding interactions reported in this study are likely conserved in holoenzymes involving other B'/PR61 isoforms.

**Biochemical Analysis of Subunit Interaction**

PP2A is a tumor suppressor protein. Mutations in both the α and β isoforms of the scaffolding subunit have been found in cancer. Glu64 of Aα was mutated to Asp in lung cancer and to Gly in breast cancer (Ruediger et al., 2001b). Several cancer-derived missense mutations have been identified in Aβ, including P65S, L101P, K343E, D504G, and V545A (Wang et al., 1998). These five mutations correspond to P53S, L89P, K331E, D492G, and V533A, respectively, in Aα. To examine the effect of these mutations on PP2A subunit interaction, we generated mutant Aα proteins, each containing one such mutation. Using a published protocol (Li and Virshup, 2002) and homogeneous recombinant proteins, we examined the effect of each mutation on interaction with the B'/PR61 regulatory subunit. Consistent with the observed important function of Glu64 in binding to B'-γ1 (Figure 4C), mutation of Glu64 to Gly or Asp significantly compromised its binding to B'-γ1 (Figure S1A). In contrast, mutation of Pro53, Leu89, and Lys331 of Aα, which are not involved in direct interaction with B'-γ1 (Figure S1A).

We also examined the effect of mutations on the interaction between Aα and Cα. The conserved ridge of HEAT repeats 11–15 in Aα is responsible for binding to Cα. Consequently, mutation of residues (Pro53, Glu64, Leu 89, and Lys331) outside of the ridge of HEAT repeats 11–15 had no impact on the interaction between Aα and Cα (Figure S1B). Interestingly, however, the mutations Y456A, Y495A, and V533A, which affect residues at the interface with Cα, did not significantly weaken the interaction with Cα (Figure S1B). This is likely due to the fact that the interactions between Aα and Cα are extremely strong with a dissociation constant of about 0.1 nM (Kamibayashi et al., 1992); such interactions might be able to withstand the mild Ala mutations. Supporting this notion, the mutation of Val533 to a charged residue Asp resulted in abrogation of the interaction between Aα and Cα (Figure S1B).

It is noteworthy that Val533 in Aα corresponds to Val545 in Aβ, which was mutated in colon cancer (Wang et al., 1998). Finally, mutation of Asn535, which mediates two hydrogen bonds at the interface, to Lys also abolished the interaction between Aα and Cα (Figure S1B).

**Conformational Changes in the Scaffolding Subunit**

The conformation of the free Aα is well defined, with only minor variation in the carboxy-terminal HEAT repeats (Groves et al., 1999). Formation of the PP2A core enzyme resulted in a pronounced conformational change in Aα, which pivots around HEAT repeat 13 (Xing et al., 2006). The core enzyme associates with a variable regulatory subunit to form a holoenzyme. To assess additional conformational changes brought about by binding to the regulatory subunit, we attempted to superimpose the core structure of the PP2A core enzyme—the entire Cα and HEAT repeats 11–15—with the same fragment from the PP2A holoenzyme. This analysis revealed a drastic conformational rearrangement (Figure 5A). Formation of the holoenzyme forces the N-terminal HEAT repeats of Aα to twist and to move by as much as 50–60 Å. As a result of this dramatic change, the amino and carboxyl termini of Aα are now 53 Å apart from each other, compared to 70 Å in the PP2A core enzyme.

To locate the site of the conformational rearrangement, we compared the result of two alignments of Aα between the core enzyme and the holoenzyme. One alignment was based on the carboxy-terminal 5 HEAT repeats (Figure 5B, left panel) whereas the other alignment relied on the amino-terminal 10 HEAT repeats (Figure 5B, right panel). This comparison revealed that, in both cases, drastic conformational rearrangement originates in the amino-terminal helix of HEAT repeat 11 (Figures 5B and 5C). Hydrophobic packing within protein interior remains unchanged for HEAT repeats 1–10 and 12–15. However, packing interactions both within HEAT repeat 11 and between HEAT repeats 11 and 12 have changed.

To accurately identify the pivoting residues, we examined this helix in atomic detail and found one significant difference. The distance between the carboxyl oxygen atom of residue 401 and the amide nitrogen atom of residue 405 is 2.9 Å in the core enzyme, suggesting a normal hydrogen bond (Figure 4D). In contrast, this distance is enlarged to 4.7 Å in the holoenzyme, which is beyond the range of a hydrogen bond (Figure 4D). Thus, the breakage of the main chain hydrogen bond between residues 401 and 405 in HEAT repeat 11 appears to make a dominant contribution to the observed conformational rearrangement. Interestingly, Pro406 is located immediately carboxy-terminally to the broken hydrogen bond; this residue likely endows the helix in HEAT repeat 11 the ability to bend.

**Role of Methylation in Holoenzyme Assembly**

Methylation of the carboxyl terminus of the catalytic subunit is thought to play an important role in the formation
Formation of a heterotrimeric PP2A complex involving the lytic subunit facilitates but may not be required for the observations strongly argue that methylation of the catalytically core enzyme (Tolstykh et al., 2000). These PP2A holoenzyme involving the B/PR55 and PR61 subunits were found to associate with the scaffolding subunit alone (Li and Virshup, 2002) or with the unmethylated PP2A core enzyme (Tolstykh et al., 2000). These observations strongly argue that methylation of the catalytic subunit facilitates but may not be required for the formation of a heterotrimeric PP2A complex involving the B/PR61 subunits. This notion is consistent with the structural observation that the carboxy-terminal residue Leu309 does not mediate significant interactions in the PP2A holoenzyme (Figure 4C).

To investigate the role of methylation in the assembly of a PP2A holoenzyme involving the B/PR61 subunits, we prepared three variants of the catalytic subunit: the fully methylated Cα, the unmethylated Cα, and the truncated Cα (residues 1–294) that is missing the carboxy-terminal 15 amino acids. These Cα variants were individually used to associate with Aα to form the core enzymes, which were then incubated with Bαγ1 for holoenzyme assembly as assessed by gel filtration. As anticipated, the fully methylated PP2A core enzyme formed a stable heterotrimeric complex with Bγ1 (Figure 6A, top panel). Interestingly, the unmethylated PP2A core enzyme and that involving the truncated Cα also formed stable complexes with Bγ1 (Figure 6A, middle and bottom panels). These results unambiguously show that formation of the heterotrimeric PP2A complex in vitro does not require methylation of Cα. Using isothermal titration calorimetry (ITC), the binding affinity between Bγ1 and the unmethylated PP2A core enzyme was determined to be approximately 52.6 ± 7.9 nM (Figure 6B).

Next, we crystallized the heterotrimeric PP2A complexes involving Aα, truncated Cα (residues 1–294), and Bγ1 or Bγ3. We determined both crystal structures and refined the atomic models to 3.8 and 3.6 Å resolution, respectively (Table 1 and Figures 6C and S2). These two structures are nearly identical to each other, with an rmsd of 0.6 Å for all backbone alpha carbon atoms. Thus, we focus our discussion on one such complex. Notably, compared to the methylated PP2A holoenzyme, the absence of methylation does not appear to cause any significant structural rearrangement in the holoenzyme involving the truncated Cα, as judged by the relatively small rmsd of 0.66 Å over 1256 aligned alpha carbon atoms. The primary protein–protein interfaces between Aα and Cα and between Aα and Bγ are also exactly preserved in the heterotrimeric PP2A complexes involving the truncated Cα. The only apparent difference is that interactions mediated by the carboxy-terminal 15 amino acids of Cα are now absent (Figure 6C).

DISCUSSION

The heterotrimeric PP2A holoenzyme is believed to determine the substrate specificity as well as the spatial and temporal functions of PP2A. For example, the PP2A holoenzyme involving the B/PR61 family plays an essential role...
Figure 6. Role of Methylation in the Assembly of PP2A Holoenzyme Involving B-/PR61 Subunit

(A) In vitro assembly of a heterotrimeric PP2A complex does not depend on methylation of the catalytic subunit. Compared to the fully methylated Cα (top panel), the unmethylated Cα (middle panel) and the carboxy-terminally truncated Cα (residues 1–294) (bottom panel) retained the ability to form a stable complex with Aα and Bγ-γ1. Shown here are representative gel-filtration runs, with the peak fractions visualized by SDS-PAGE and Coomassie blue staining.

(B) Measurement of the binding affinity between Bγ-γ1 and the unmethylated PP2A core enzyme using isothermal titration calorimetry (ITC).

(C) Structure of the heterotrimeric PP2A complex involving Aα (green), Bγ-γ3 (yellow), and the truncated Cα (blue, residues 1–294). The red oval circle denotes the absence of interactions involving the carboxyl terminus of Cα.

(D) A structure-based model for the assembly and regulation of the PP2A holoenzyme. The first step in holoenzyme assembly is the association of the free scaffolding subunit with the catalytic subunit to form the PP2A core enzyme. This association results in significant conformational changes in HEAT repeats 13 and to a lesser extent HEAT repeat 12 (Xing et al., 2006). The PP2A core enzyme then associates with a regulatory B-/PR61 subunit to form a holoenzyme. This association results in more drastic conformational rearrangements in the scaffolding subunit. The regulatory B-/PR61 subunit recruits substrate proteins using its acidic, concave groove. The red star in the catalytic subunit denotes the active site.

Cell 127, 1239–1251, December 15, 2006 ©2006 Elsevier Inc. 1247
in cell-cycle progression, through direct interaction with the protein Shugoshin (Kitajima et al., 2006; Riedel et al., 2006; Tang et al., 2006). In this manuscript, we report the crystal structures of the PP2A holoenzyme involving the regulatory B’/PR61 subunits.

These structures reveal a number of unexpected findings. First, the structure of B’/PR61 closely resembles that of a number of other superhelical proteins, including karyopherin, β-catenin, and the scaffolding subunit of PP2A. This unexpected structural mimicry raises a number of interesting scenarios. The scaffolding subunit uses its ridge to interact with the catalytic and the regulatory subunit. Similarly, the B’/PR61 subunit uses its own ridge to bind to the catalytic subunit; in the meantime, the convex side of the B’/PR61 subunit recognizes the ridge of the scaffolding subunit. This arrangement leaves the highly acidic, concave side of B’/PR61 unoccupied. It is possible that the B’/PR61 subunit relies on the acidic, concave surface to interact with substrate proteins or peptide motifs. Consistent with this hypothesis, the concave surface is tilted towards the active site pocket of the PP2A catalytic subunit. Additional support comes from the observation that both karyopherin and β-catenin contain a charged, concave surface groove that interacts with peptides (Conti et al., 1998; Graham et al., 2000), although importin-β can also bind to peptides using its convex surface (Bayliss et al., 2000). We propose that, following the assembly of a PP2A holoenzyme involving a B’/PR61 subunit, the concave, acidic surface is used to recruit substrate proteins, which may contain a positively charged epitope (Figure 6D). Supporting this hypothesis, Shugoshin, which interacts with the B’/PR61 subunit in cell-cycle regulation, is a basic protein.

Although conformational flexibility was evident from the study of the PP2A core enzyme (Xing et al., 2006), the pronounced structural changes upon formation of the holoenzyme is still unanticipated. When the core domains of the PP2A core enzyme are aligned, the amino-terminal HEAT repeats of the scaffolding subunit swing by as much as 50–60 Å. Through structural analysis, we located the switch point to be within the first helix of HEAT repeat 11 and mapped the helix-breaking amino acids to be within residues 401–406. The characteristic presence of proline in the HEAT repeats is believed to cause the bending of helices in the outer layer and subsequent formation of the horse-shoe shaped scaffolding subunit (Groves et al., 1999). Here, the presence of proline in HEAT repeat 11 is likely to be a major source of the observed conformational flexibility, which is likely essential for PP2A function. The PP2A core enzyme must specifically associate with four different classes of the regulatory subunits and must act to remove phosphate groups in different substrates; having conformational latitude may facilitate binding to regulatory subunits and help accomplish catalysis.

Our structural analysis provides a mechanistic basis for a body of published data on the assembly of PP2A holoenzyme (Kamibayashi et al., 1992, 1994; Li and Virshup, 2002; Ruediger et al., 1992, 1994). HEAT repeats 1–10 were found to be required for interaction with the regulatory subunit (Ruediger et al., 1994). Our structural analysis shows, however, only HEAT repeats 2–8 of Aε are directly involved in binding to B’/PR61. One plausible explanation for the difference is that HEAT repeats 1, 9, and 10 might be important for the structural integrity of HEAT repeats 2–8. Supporting this explanation, a mutant Aε with truncation of the first HEAT repeat exhibited poor solution behavior and failed to assemble into a holoenzyme (data not shown).

Two regions of the regulatory B’/B55/PR61 subunits were found to mediate binding to the scaffolding subunit of PP2A (Li and Virshup, 2002). One of these regions includes residues 200–303, which correspond to HEAT-like repeats 4 and 5 (Figure 3). This region contains seven amino acids that directly interact with Aε. One of the seven residues, Glu214, makes a pair of hydrogen bonds to Arg183 in Aε. Interestingly, the two regions of the B’/PR61/B56 subunits appeared to be conserved in the regulatory B/B55/PR55 and B’/PR72 subunits, with an invariant Glu corresponding to Glu214 in B’/B56/PR61 (Li and Virshup, 2002). This observation raises the possibility that some of the interactions between the scaffolding and the regulatory B’/B56/PR61 subunits may be conserved for the other two major families of regulatory subunits (Li and Virshup, 2002). However, this possibility does not necessarily imply that the other two families of regulatory subunits have similar helical structures as the B’/B56/PR61 family. In fact, B/B55/PR55 is predicted to contain multiple WD40 repeats, whereas B’/PR72 is thought to contain two calcium binding EF hands (Janssens et al., 2003).

While this paper was under review, a manuscript describing the structure of the PP2A holoenzyme was published online (Cho and Xu, 2006). This holoenzyme contains the full-length Aε, Cε, and an amino-terminally truncated B’-γ1 (Cho and Xu, 2006). Both structures show unambiguously that the carboxy-terminal residues of Cε contribute to the formation of PP2A holoenzyme (Figure 4C). Intriguingly, however, these interactions do not appear to be required for the assembly of a heterotrimeric PP2A complex in vitro (Figure 6). In fact, the B’/PR61 subunit can bind to Aε in the absence of Cε (Li and Virshup, 2002) (Figure S1A). Although these in vitro experiments may not necessarily recapitulate the circumstances in cells, the structures of the PP2A holoenzyme provide powerful references for understanding the function and regulation of PP2A.

EXPERIMENTAL PROCEDURES

Protein Preparation and Assembly of PP2A Holoenzyme

All constructs and point mutations were generated using a standard PCR-based cloning strategy. B’-γ1 and B’-γ3 were overexpressed in E. coli as a fusion protein with glutathione S transferase (GST) and purified as described (Xing et al., 2006). The PP2A core enzyme, involving the full-length Cε (residues 1–309) and Aε (residues 8–589), was assembled as described (Xing et al., 2006). The PP2A core enzyme was methylated by a PP2A-specific leucine carboxyl methyltransferase...
Crystallization and Data Collection

Diffraction crystals were obtained for the three PP2A holoenzyme complexes described above, which were individually incubated with 1.2 molar equivalence of MCLR prior to crystallization. We also generated crystals of the holoenzyme using selenomethionine-substituted scalding and regulatory subunits. Crystals were grown by the hanging-drop vapor-diffusion method by mixing the protein (−8 mg/ml) with an equal volume of reservoir solution containing 10%–15% PEG8000, 0.1 M Tris-Cl, pH 8.5, and 0.2 M magnesium sulfate. Small crystals appeared within a few hours. Macroseeding was used to generate single, large crystals. The crystals belong to the space group P212121, with a = 109.29 Å, b = 159.05 Å, and c = 269.17 Å. There are two complexes per asymmetric unit, and the solvent content is approximately 80%. Crystals were slowly equilibrated in a cryoprotectant reservoir buffer plus 20% glycerol (v/v) and were flash frozen in a cold nitrogen stream at −170 °C. The native and selenium Sad data sets were collected at NSLS beamline X29 and processed using the software Denzo and Scalepack (Otwinowski and Minor, 1997).

Structure Determination

The structure of human PP2A holoenzyme was determined by molecular replacement using three models: Cα and two fragments of Aα (residues 9–415 and 416–589) from the structure of the previously determined PP2A core enzyme (accession code 2IE3) (Xing et al., 2006). Fragments were located using the program PHASER (McCoy et al., 2005). The backbone of Bγ was built into model-phased two-fold-averaged Fobs − Fc and Fc − Fobs electron density maps. Side-chain interpretation of SeMet Sad data collected from complexes containing SeMet-labeled Aα and Bα components. Model building was performed using O (Jones et al., 1991) and refined using CNS (Brüger et al., 1998). Tight NCS restraints between the two complexes (50 kcal/mol) were applied throughout most of the refinement and relaxed in the final cycles. Positional refinement was performed against maximum likelihood target with NCS- and geometrically restrained individual B factor refinement, with weights adjusted on the basis of Rfree. In tests, this refinement method consistently gave a lower Rfree than grouped B factor refinement (which cannot be restrained by geometry in CNS) and was considerably better than adopting a single overall or per-domain B factor model. The final atomic model of the native holoenzyme has been refined to 3.3 Å resolution. Due to nearly identical unit cells, structures of the two heterotrimeric PP2A complexes involving the truncated Cα were directly refined at 3.6 and 3.8 Å resolution after removal of the carboxy-terminal 15 amino acids from Cα. Similar positional and B factor refinement protocols were used as described above. For the native holoenzyme, the atomic model contains amino acids 2–309 for Cα, residues 8–589 for Aα, and residues 38–66 and 68–425 for Bγ−1. There is no electron density for residues 1–37 and 426–439 of Bγ−1; we presume these regions are disordered in the crystals. The methyl group on the methylated carboxyl terminus of Cα was not modeled, because there is no clear electron density for this group at this resolution. For the holoenzyme involving the truncated Cα residues 1–294 and Bγ−3 or Bγ−1, the atomic models contain amino acids 2–294 for Cα, residues 8–589 for Aα, and residues 38–66 and 68–425 for Bγ−3 or Bγ−1.

Methylation of PP2A Core Enzyme by LCMT

LCMT and PP2A core enzyme, at a 1:2 molar ratio, was incubated on ice. Methylation was initiated by addition of SAM (S-adenosyl methionine) to a final concentration of 0.75 mM. The reaction was carried out at 22 °C and reached completion after 2–3 hours. The methylated PP2A core enzyme was purified away from LCMT by anion exchange chromatography. The extent of methylation was examined using an antibody that only recognizes the unmethylated carboxyl terminus of Cα (Xing et al., 2006).

GST-Mediated Pull-Down Assay

Approximately 30 μg of GST-Aα were bound to 30 μl of glutathione resin. The resin was washed with 200 μl assay buffer for three times to remove excess unbound Aα. Then 20 μg of wild-type (WT) or mutant Bγ−1 were allowed to bind the resin in a 125 μl volume. After four times of washing with an assay buffer containing 25 mM Tris, pH 8.0, 150 mM NaCl, and 2 mM dithiothreitol (DTT), the remaining protein and resin were mixed with 15 μl SDS sample buffer and applied to SDS-PAGE. The results were visualized by SDS-polyacrylamide gel electrophoresis (SDS-PAGE) with Coomassie blue staining.

Isothermal Titration Calorimetry (ITC)

To obtain a direct binding affinity between PP2A core enzyme and Bγ−1, 0.1 mM Bγ−1 were titrated against 9 μM unmethylated PP2A core enzyme using VP-ITC microcalorimeter (MicroCal). All proteins were prepared in a buffer containing 25 mM HEPES, pH 8.0, and 150 mM NaCl. The data were fitted by software Origin 7.0.

Supplemental Data

Supplemental data include two figures and can be found with this article online at http://www.cell.com/cgi/content/full/127/6/1239/DC1/.

ACKNOWLEDGMENTS

We thank A. Saxena at BNL NSLS beamlines for help. This work was supported by grants from the US National Institutes of Health (Y.S.).

Received: October 19, 2006
Revised: November 20, 2006
Accepted: November 28, 2006
Published: December 14, 2006

REFERENCES


**Accession Numbers**

The atomic coordinates of the three PP2A holoenzyme complexes have been deposited in the Protein Data Bank with the accession codes 2NPP (full-length Ca), 2NYM (truncated Cα and Bβγ0γ3), and 2NYL (truncated Cα and Bβγ1).