Comparative Structural Analysis of the Erbin PDZ Domain and the First PDZ Domain of ZO-1

INSIGHTS INTO DETERMINANTS OF PDZ DOMAIN SPECIFICITY

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We report a structural comparison of the first PDZ domain of ZO-1 (ZO1-PDZ1) and the PDZ domain of Erbin (Erbin-PDZ). Although the binding profile of Erbin-PDZ is extremely specific ([D/E][T/S][W/Y][V/L/I]COOH), that of ZO1-PDZ1 is similar ([R/K/S/T][T/S][W/Y][V/L/I]COOH), but broadened by increased promiscuity for three of the last four ligand residues. Consequently, the biological function of ZO-1 is also broadened, as it interacts with both tight and adherens junction proteins, whereas Erbin is restricted to adherens junctions. Structural analyses reveal that the differences in specificity can be accounted for by two key differences in primary sequence. A reduction in the size of the hydrophobic residue at the base of the site0 pocket enables ZO1-PDZ1 to accommodate larger C-terminal residues. A single additional difference alters the specificity of both site−1 and site−3. In ZO1-PDZ1, an Asp residue makes favorable interactions with both Tyr−1 and Lys/Arg−3. In contrast, Erbin-PDZ contains an Arg at the equivalent position, and this side chain cannot accommodate either Tyr−1 or Lys/Arg−3 but, instead, interacts favorably with Glu/Asp−3. We propose a model for ligand recognition that accounts for interactions extending across the entire binding site but that highlights several key specificity switches within the PDZ domain fold.

PDZ2 (PSD-95/Discs-large/ZO-1) domains are compact globular modules that typically recognize specific C-terminal motifs and, in so doing, assemble multicomponent protein complexes inside eukaryotic cells (1, 2). PDZ domains are usually embedded in larger multidomain scaffold proteins that often include multiple PDZ domains and other protein-binding modules. Thus, the biological function of each PDZ domain is determined by its intrinsic ligand specificity and also by the context in which it interacts with other cellular components. To understand how the diverse members of the PDZ domain family have adapted to their particular biological roles, it will be important to define accurately the specificity of each domain and the structural basis for ligand recognition. To this end, we have conducted comparative structural and functional studies of the PDZ domains of the human LAP (leucine-rich repeats and PDZ domains) family member Erbin (3, 4) and the MAGUK (membrane-associated guanylate kinase) family member zonula adherens-1 (ZO-1) (5), as these two proteins offer an opportunity to examine both the similarities and differences between domains that have adapted to similar but distinct biological functions.

In an accompanying article (6), we used phage-displayed peptide libraries to define the binding specificities of the single PDZ domain of Erbin (Erbin-PDZ) and two of the three PDZ domains of ZO-1, along with those of the other human LAP family members Densin-180 and Scribble (or Scrib) (Fig. 1). We found that Erbin-PDZ exhibits a strict preference for particular types of residues at each of the last four ligand positions and thus recognizes a highly conserved core binding motif ([E/D][T/S][W/Y]COOH). Interestingly, the first PDZ domain of ZO-1 (ZO1-PDZ1) recognizes a core binding motif ([T/S/K/R][T/S][W/Y][V/L/I]COOH) that resembles that of Erbin-PDZ but also exhibits significant differences. In particular, it appears that the ligand binding profile of ZO1-PDZ1 is expanded in comparison with that of Erbin-PDZ by virtue of increased promiscuity for three of the last four ligand residues. Specifically, both domains accept ligands containing Val at position, but only ZO1-PDZ1 is able to accept the larger Leu/Ile residues with comparable affinity. At site−1, Erbin-PDZ exhibits a strict preference for Trp side chains, whereas ZO1-PDZ1 is bispecific, accepting either Trp or Tyr but discriminating against other residues. Finally, the two domains differ dramatically in their preferences at site−3. Erbin-PDZ exhibits a preference for acidic residues, whereas ZO1-PDZ1 exhibits bispecific character at this site, preferring either basic residues or small, polar Thr/Ser residues.

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The atomic coordinates and structure factors (codes 2H2B, 2H2C, 2H3L, 2H3M) have been deposited in the Protein Data Bank, Research Collaboratory for Structural Bioinformatics, Rutgers University, New Brunswick, NJ (http://www.rcsb.org/).

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3 The abbreviations used are: PDZ, PSD-95/Discs-large/ZO-1; GUK, guanylate kinase; LAP, leucine-rich repeats and PDZ; MAGUK, (membrane-associated guanylate kinase) family member zonula occludens-1 (ZO-1); r.m.s.d., root mean square deviation.

4 The C terminus of a PDZ domain ligand is designated as position0, and the remaining positions are numbered with negative integers for which the absolute value decreases toward the C terminus. The corresponding binding sites on the PDZ domain are numbered in a corresponding manner as site0, site−1, etc.
These differences and similarities in ligand specificity reflect the overlapping yet distinct subcellular localizations and biological interactions of Erbin and ZO-1. Erbin and the other LAP family members are involved in the organization of junctional complexes and in the establishment and maintenance of cell polarity; and in polarized epithelia, they are localized on the lateral membrane and at adherens junctions (7–9). Notably, we and others have shown that Erbin-PDZ interacts strongly with three members of the p120-like catenin family (H9254-catzenin, ARVCF (armadillo protein deleted in velo-cardiofacial syndrome), and p0071) (10–12), which contain a conserved C-terminal sequence (DSWVCOOH) that closely matches the optimal binding motif defined by phage display. These p120-like catenins also associate with other catenins and E-cadherin to form the cadherin-catenin complex, a major component of adherens junctions (13).

In non-epithelial cells and nonpolarized epithelial cells, ZO-1 also associates with adherens junction markers (14–18), and its localization resembles that of Erbin. Indeed, it has been shown that, under these conditions, ZO-1 co-localizes with E-cadherin, and like Erbin-PDZ, ZO1-PDZ1 binds to the C terminus of ARVCF (11, 16). However, as polarization proceeds and distinct junctional complexes form, ZO-1 segregates away from E-cadherin and localizes exclusively to the tight junctions that form apical to the adherens junctions (14, 19). The recruitment of ZO1 to the tight junctions is mediated by interactions between ZO1-PDZ1 and the C termini of claudins (20, 21), which are the major structural proteins of tight junctions and for the most part share a common C-terminal motif (YVCOOH) (6). In contrast, Erbin and the other LAP family members are excluded from the tight junctions and remain confined to the adherens junctions and the basolateral layer (7–9).

Thus, it appears that the bispecific character of the site0 and site1 sites of ZO1-PDZ1 reflects the bifunctional nature of ZO-1 in the cellular environment. In nonpolarized cells, ZO1-PDZ1 is able to interact at adherens junctions with p120-like catenins that contain Trp1 (16), whereas in polarized cells it is able to interact at tight junctions with claudins that contain Tyr1 (6, 21). On the other hand, Erbin-PDZ has a strict preference for Trp1, and even in polarized cells, Erbin does not interact with claudins but, rather, remains associated with p120-like catenins at the adherens junctions (10–12). These differences in cellular localization are also likely to be enforced by differences in site3. Erbin-PDZ prefers acidic residues at position3, and all three p120-like catenins described above contain an Asp at this position. In contrast, acidic residues are conspicuously absent at position3 among the 20 known human claudins, which typically contain either basic residues or Thr/Ser at this position, in good agreement with the binding specificity of ZO1-PDZ1 (6).

In contrast with earlier studies that emphasized the importance of site0 and site1 (22–24), more recent studies, including the work described above, make it clear that PDZ domains recognize ligands through interactions that extend across the entire surface of the binding groove (25–33). According to the established PDZ domain classification system based on site0 and site1, both Erbin-PDZ and ZO1-PDZ1 exhibit typical class I specificities (X[T/S]XΦCOOH, where Φ is a hydrophobic residue). However, it is clear that the ligand-binding specificities of these domains depend critically on interactions with all four C-terminal residues and, importantly, that differences in
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the specificities of site$^{-1}$ and site$^{-3}$ are critical determinants of functional convergence and divergence.

Herein we report a comparative structural analysis aimed at elucidating the molecular basis for the similarities and differences between the binding specificities of Erbin-PDZ and ZO1-PDZ1. We solved the unliganded crystal structures of both domains, as well as high resolution structures of ZO1-PDZ1 in complex with two different phage-derived ligands designed to investigate the broadened specificity of ZO1-PDZ1. Together with a previously reported structure of Erbin-PDZ bound to an optimal ligand (29), these structures allow for a detailed comparison of the ligand-binding sites of the two domains. Remarkably, the analysis reveals that the differences in the site$^{-1}$ and site$^{-3}$ specificities of Erbin-PDZ and ZO1-PDZ1 are likely caused by a single amino acid difference between the two domains, and furthermore, the differences in site$^9$ can be accounted for by one additional difference. Overall, the results support a model for PDZ domain-ligand interactions that takes into account interactions that extend across the entire binding site but depends critically on several key specificity switches within the PDZ domain fold.

EXPERIMENTAL PROCEDURES

Protein Purification—Erbin-PDZ (residues 1314–1422) was purified as described previously (29). For ZO1-PDZ1, a DNA fragment encoding for residues 18–110 of human ZO-1 was cloned into the NdeI/BamHI sites of the pET22d expression vector, and this created an open reading frame encoding for ZO1-PDZ1 with an N-terminal His tag and a thrombin cleavage site. In addition, standard molecular biology techniques were used to fuse 10-residue extensions to the C terminus of ZO1-PDZ1-YL (0.1M sodium acetate, 0.2M ammonium sulfate, 30% ethylene glycol 4000 (pH 7.5–8.5)), ZO1-PDZ-WV (0.1M sodium acetate, 0.2M ammonium sulfate, 30% polyethylene glycol 2000 monomethyl ether (pH 4.6)). For cryoprotection, each crystal was soaked briefly in a mixture of 75% reservoir solution and 25% ethylene glycol and subsequently flash-frozen in liquid nitrogen.

Diffraction data were collected at beamline 5.0.1 (Advanced Light Source, Berkeley, CA) for the ZO1-PDZ1-WV and ZO1-PDZ1-YL crystals, at beamline 9-2 (Stanford Synchrotron Radiation Laboratory, Palo Alto, CA) for the Erbin-PDZ crystal, or on an in-house Cu-Ko source for the ZO1-PDZ1 crystal (Table 1). Data were processed with DENZO and SCALEPACK from the HKL suite (34). The structures were solved by molecular replacement with AMoRe (35) or Phaser (36) using the PDZ domain from a liganded Erbin-PDZ structure (PDB code 1MFG) (37) as the search model for the unliganded Erbin-PDZ structure, our refined Erbin-PDZ structure as the search model for the unliganded ZO1-PDZ1 structure, and the refined coordinates of the ZO1-PDZ1 structure as the search model for the liganded ZO1-PDZ1 structures. Atomic models were built with multiple rounds of manual inspection using XtalView (38) or O (39) and refined by the maximum likelihood method with REFMAC (40).

Each of the three ZO1-PDZ1 structures contains one molecule per asymmetric unit; the Erbin-PDZ structure contains two molecules per asymmetric unit. The ZO1-PDZ1-WV and ZO1-PDZ1-YL structures form dimers. The 2-fold axes relating the molecules within these dimers to each other coincide in both cases with a crystallographic 2-fold axis. In each dimer, the two PDZ monomers are arranged in a head-to-tail fashion where each PDZ domain binds to the C-terminal peptide of its symmetry-related binding partner. For all of the ZO1-PDZ1 structures, the entire PDZ domain is well defined in the electron density. In the Erbin-PDZ structure, the entire PDZ domain is visible in the first molecule, but the last residue of the C terminus is disordered in the second.

RESULTS

Crystallographic Analysis of Erbin-PDZ and ZO1-PDZ1—We determined the crystal structures of unliganded ZO1-PDZ1 and Erbin-PDZ, and the structures of ZO1-PDZ1 in complex with two high affinity heptapeptides (WRRTTWV$\text{COOH}$ and WRRTTYL$\text{COOH}$) (Table 1) identified by C-terminal phage display (6). The heptapeptides differ at the two C-terminal positions and were chosen for structural analysis to investigate the subtle differences between the specificities of ZO1-PDZ1 and Erbin-PDZ. To facilitate structural studies, we used a construct in which the peptides of interest were fused to the C terminus of ZO1-PDZ1 via a tri-glucylic linker. This strategy was used previously to determine the crystal structures of various PDZ domain-ligand complexes (27, 28, 37) and, in the case of ZO1-PDZ1, resulted in the formation of crystallographic dimers in which each fused ligand binds to the apposing PDZ domain (Fig. 2A). The crystal structures for the resulting fusion proteins are subsequently referred to as ZO1-PDZ1-WV or ZO1-PDZ1-YL for
the structures of the complex with WRRTTWV<sub>COOH</sub> or WRRTTYL<sub>COOH</sub>, respectively. All four structures presented here were solved by molecular replacement and are well refined with R-values ranging from 14.8 to 22.7% and with good geometry. In particular, the ZO1-PDZ1-YL and -WV structures were solved at 1.6 and 2 Å (Table 1), respectively, and thus provide high enough resolution to allow commenting on critical differences of the bound ligands. This enables us to suggest how specificity among related PDZ domains is achieved on a structural basis.

**Overall Structure**—The canonical PDZ domain fold is characterized by a six-stranded β-sandwich that is capped with two α-helices. Helix α1 usually follows strand β3. However, similar to previously reported structures of Erbin-PDZ (29, 37), ZO1-PDZ1 lacks helix α1 (Fig. 2B). Helix α2 follows strand β5 and is present in all PDZ domains. Helix α2 and strand β2 form either side of the conserved ligand-binding cleft within the PDZ domain fold. In the liganded ZO1-PDZ1 structures, the heptapeptides are bound in this cleft and are well defined in the electron density. The temperature factors of the bound ligands are among the lowest in the entire structure with the exception of the guanidinium groups of Arg<sup>−5</sup> and Arg<sup>−6</sup>, which are mobile in both structures.

To facilitate the comparison of PDZ structures, we have adopted a naming convention that recognizes structurally equivalent residues (Fig. 1). In this numbering scheme, residue β2-1 for example, represents the first residue in strand β2, whereas β2:β3-1 indicates the first amino acid in the loop between strands β2 and β3. Because of numerous insertions and deletions within the loop regions of many PDZ domains, it is not possible to unequivocally assign structurally equivalent residues in these regions. Therefore, the numbering represents the lengths of the longest loops used in our alignment.

At the high protein concentrations used during crystallization, the ligand-binding grooves of PDZ domains are often occupied by the C termini of symmetry-related molecules.
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**A** Erbin-PDZ  
ZO1-PDZ1

**B**



- **β2:β3-8**
- **β2:β3-9**
- **Trp<sup>6</sup>**

**FIGURE 3.** Superposition of ZO1-PDZ1 and Erbin-PDZ structures. A, the left panel shows the superposition of structures for unliganded (light orange) and liganded Erbin-PDZ (PDZ, green; peptide, magenta; PDB 1MFL). The right panel displays the superposition of structures for unliganded (yellow) and liganded ZO1-PDZ1 (ZO1-PDZ-YL: PDZ, gray; peptide, green; ZO1-PDZ-WV: PDZ, light blue; peptide, magenta). The arrow indicates the position of the β2:β3 loop, B, the conformation of the β2:β3 loop of ZO1-PDZ1 differs between the unliganded (top) and liganded (bottom) forms. This difference is due to rotation about the peptide backbone at residue β2:β3-9 and results in the formation of a hydrophobic pocket that is occupied by Trp<sup>6</sup> in the liganded form.

Within the crystal lattice (23, 26, 41). Fortunately, the peptide-binding grooves in the unliganded ZO1-PDZ1 and Erbin-PDZ structures presented here are empty and thus provide an opportunity to compare the unliganded and liganded forms of these domains. Crystallographic analysis of the sixth PDZ domain of glutamate receptor-interacting protein (GRIP-PDZ6) revealed that ligand binding induces a reorientation of helix α2 (26), suggesting that PDZ domains might undergo conformational changes to accommodate ligands. In contrast, superposition of the unliganded and liganded structures of the third PDZ domain of PSD-95 (22) and those of the Shank1 PDZ domain (25) did not show significant conformational changes upon ligand binding. Similar to the Shank1 PDZ domains, our unliganded Erbin-PDZ structure has root mean square deviation (r.m.s.d.) values of 0.6 to 0.7 Å when compared with two Erbin-PDZ structures in complex with low affinity peptides (Fig. 3A) (37), indicating that no major structural changes are induced upon ligand binding. Similarly, the three ZO1-PDZ1 structures presented here superimpose with root mean square deviation values of only 0.6 to 0.9 Å for 93 equivalent Ca atoms, and aside from subtle movement in the β2:β3 loop, there are no structural differences. The movement in the β2:β3 loop is induced by the rotation of Phe(β2:β3-9) around the peptide backbone, which results in a 5-Å movement of its Ca atom (Fig. 3B). In summary, it seems that ligand binding does not induce conformational changes in most PDZ domains studied thus far.

**Comparison of Site<sup>0</sup> and Site<sup>-2</sup> of Erbin-PDZ and ZO1-PDZ1—**

To better understand the molecular basis for ligand recognition and selectivity by PDZ domains, we compared the liganded structures of ZO1-PDZ1 bound to peptides WRRTTWV<sub>COOH</sub> or WRRTTWV<sub>COOH</sub> with an NMR structure of Erbin-PDZ bound to a high affinity peptide (TGWETTWV<sub>COOH</sub>) (29). In each structure, the PDZ domain forms extensive contacts with the bound ligand. Some of these interactions are observed in most PDZ domains, whereas other contacts seem to be specific to class I PDZ domains preferring Thr/Ser at position -2. In all three structures, the Thr<sup>-2</sup> hydroxyl group is able to form a hydrogen bond to N<sup>-2</sup> of His(α2-1), whereas the aliphatic portion of the side chain is in van der Waals distance to Val(α2-5) (Fig. 4A). These interactions are consistent with the functional analyses of these PDZ domains; they are also in agreement for the PDZ domains of the other LAP family members Densin-180 and Scribble, which also contain His and Val at α2-1 and α2-5, respectively, and show a strong preference for Thr/Ser<sup>-2</sup> (6). Notably, the third PDZ domain of ZO-1 (ZO1-PDZ3) exhibits no specificity at site<sup>-2</sup>; this is the only domain that contains an Arg at α2-1 (Fig. 1).

At site<sup>-2</sup>, all three structures are very closely related in the way that the C-terminal carboxylate group of the ligand forms hydrogen bonds with the main chain amides of the carboxylate binding loop (β1:β2-7, β1:β2-8, and β2-1). Interactions of this type are conserved among all PDZ domains that engage ligands with free C termini (22, 28, 29). However, the structures differ significantly in terms of the interactions that mediate recognition of the C-terminal side chain (Fig. 4A). In each liganded ZO1-PDZ1 structure, the side chain of Val<sup>0</sup> or Leu<sup>0</sup> is buried in a deep hydrophobic pocket that is formed predominantly by residues in the β1:β2 loop and strand β2 (Phe(β1:β2-7), Ile(β2-1), Ile(β2-3)) but also involves residues in helix α2 (Val(α2-5), Leu(α2-8)). However, depending on the bound ligand, Ile(β2-3) adopts different rotamers around the Ca–Cβ bond to accommodate either Val<sup>0</sup> or Leu<sup>0</sup>. Erbin-PDZ also contains an Ile at β2-3, but position β2-1 is occupied by a larger Phe rather than an Ile. In comparison with ZO1-PDZ1, the bulkier Phe(β2-1) side chain reduces the volume of the site<sup>0</sup> pocket of Erbin-PDZ and restricts the movement of the Ile side chain at β2-3, making the pocket less accommodating for ligands with larger side chains at position<sup>2</sup>. It is striking that the PDZ domains that have a stringent preference for a Val<sup>0</sup> side chain (Erbin-PDZ, Densin-PDZ, and Scribble, which also contain His<sup>1</sup> at α2-1) have Phe in position<sup>2</sup> (Scrib-PDZ1, Scrib-PDZ2) that restricts the movement of the Ile side chain at β2-1 (Fig. 1).

**Comparison of Site<sup>1</sup> and Site<sup>-3</sup> of Erbin-PDZ and ZO1-PDZ1—**

Erbin-PDZ displays a strict preference for Trp at site<sup>-5</sup>, whereas in contrast, ZO1-PDZ1 is bispecific and can accommodate Trp or Tyr equally well. A comparison of the various PDZ domain structures bound to ligands suggests a likely explanation for this biologically important difference in specificity (Fig. 4B). In both domains, Trp<sup>-5</sup> occupies a very similar position, and the side chain extends across strand β2 and inserts between the side chains of β3-5 and β3α1-1. Alanine-
scanning mutagenesis of Erbin-PDZ has shown that favorable recognition of the Trp⁻¹ side chain does not derive from contacts with PDZ domain side chains but, rather, from contacts with the main chain of strands β2 and β3 (29). The crystal structure of ZO1-PDZ1-WV shows as well that most interactions of ZO1-PDZ with the Trp⁻¹ side chain are mediated by main chain interactions. In the ZO1-PDZ1-YL structure, the Tyr⁻¹ side chain occupies a position that is similar to that of site⁻³; however, as the Tyr side chain is somewhat smaller than the Trp side chain, the overall amount of buried surface and hydrophobic interactions is decreased. In ZO1-PDZ1, this loss in binding energy is compensated for by the formation of a hydrogen bond between the hydroxyl group of Tyr⁻¹ to the carboxylate group of Asp(β3-5). In our accompanying article (6), we observed that ZO1-PDZ1 has similar affinities for peptides with Trp⁻¹ or Tyr⁻¹. However, the replacement with Phe⁻¹ results in a 10-fold loss in affinity, emphasizing the importance of the hydrogen bond between Tyr⁻¹ and Asp(β3-5). In contrast, Erbin-PDZ contains an Arg at position β3-5, which is not suited for the formation of a similar interaction. Thus, it appears that recognition of Tyr⁻¹ by ZO1-PDZ1 is dependent on both interactions with the main chain and hydrogen bond formation with the side chain of Asp(β3-5). Among the PDZ domains that we investigated by phage display (Fig. 1), only ZO1-PDZ1 contains an Asp in position β3-5, and it is the only one with a preference for both Trp⁻¹ and Tyr⁻¹. All of the other domains either strongly prefer Trp⁻¹ or accommodate any amino acid in this position. More detailed analysis of binding preferences in other PDZ domains will be required to determine whether an Asp in position β3-5 is a signature for PDZ domains that are bispecific at site⁻³ with a preference for both Trp and Tyr.

Erbin-PDZ and ZO1-PDZ1 differ significantly in their specificity at site⁻³, with Erbin-PDZ exhibiting a strong preference for acidic residues and ZO1-PDZ1 preferring either Thr/Ser or Arg/Lys (6). A careful analysis of the crystal structures reveals that the specificity of site⁻³ is closely tied to the residue at position β3-5. This residue, therefore, is involved in the formation of interactions at both site⁻¹ and site⁻³ (Fig. 4B). In the case of Erbin-PDZ, Arg(β3-5) is well positioned to make favorable electrostatic interactions with a negatively charged ligand side chain at site⁻³. This explains the preference of this domain and other LAP family PDZ domains to bind peptides that contain negatively charged residues at position⁻³, as they all contain Arg or

FIGURE 4. Comparison of the ligand-binding subsites of ZO1-PDZ1 and Erbin-PDZ. In each horizontal gallery, the three structures are shown in the same relative orientation. Main chain traces are rendered as gray or green tubes for the PDZ domain or peptide ligand, respectively. Side chains are rendered as sticks, and carbon atoms are colored green or yellow for the ligand or PDZ domain, respectively, except where noted. Side chain oxygen and nitrogen atoms are colored red and blue, respectively. Hydrogen bonds are shown schematically as blue dashes. A, the interactions at site⁻³ and site⁻¹ involve residues from the β1:β2 loop, strand β2, and helix α2. ZO1-PDZ1, which contains an Ile at β2-1 (orange), can accommodate Val, Leu, or Ile at site⁻³, whereas Erbin-PDZ (PDB 1N7T), which contains a Phe at β2-1, shows a strong preference for Val². Additionally, Ile(β2-3) (orange) of ZO1-PDZ1 adopts different rotamers to accommodate either Val² or Leu². B, both Erbin-PDZ and ZO1-PDZ1 accommodate Trp⁻¹, which extends over strand β2. However, ZO1-PDZ1 also recognizes Tyr⁻¹ because of a hydrogen bond formed with the carboxylate group of Asp(β3-5). In contrast, Erbin-PDZ contains an Arg at position β3-5, which instead mediates a salt bridge with Glu⁻³. The ZO1-PDZ1 structures also display a hydrogen bond between the hydroxyl atoms of Ser(β2-4) and Thr⁻³. C, ZO1-PDZ1 and Erbin-PDZ display a preference for a Trp at positions⁻⁵ and -⁴, respectively. Although these interactions are dependent on residues in the β2:β3 loop of both domains, the molecular details are different (see “Results,” under “Contributions of the β2:β3 Loop to Ligand Recognition,” for details).
Lys at position β3-5 (Fig. 1). In contrast, the Asp in position β3-5 of ZO1-PDZ1, which is responsible for the bispecific nature of this domain to accommodate either Trp or Tyr in site-1, conducts the preference of this domain toward the binding of a positively charged residue in site-1. Alternately, ZO1-PDZ readily accepts Ser or Thr in this position, small residues that are able to form hydrogen bonds with the Ser in position β2-4 (Fig. 4B). This Ser is conserved in Erbin-PDZ (Fig. 1), but the interaction between Arg(β3-5) and a Glu/Asp is likely to be stronger than the formation of a hydrogen bond between Ser(β2-4) and Thr/Serβ3. In summary, the specificity at site-3 is conducted by interactions with residues in position β2-4 or β3-5 for short or long ligand side chains, respectively.

**Contributions of the β2:β3 Loop to Ligand Recognition—** Erbin-PDZ and ZO1-PDZ1 exhibit preferences for ligands that contain hydrophobic side chains as well as arginines and lysines upstream of position-1. In both cases, recognition is mediated predominantly by residues of the β2:β3 loop, but the structural details of these interactions differ considerably (Fig. 4C). In the case of Erbin-PDZ, the side chain of Trpβ2-4 packs against strand β2 and interacts with residues in strands β2 and β3 (Ser(β2-4), Thr(β3-4)) and the β2:β3 loop (Pro(β2:β3-8) and Phe(β2:β3-9)). In ZO1-PDZ1, the β2:β3 loop forms a number of contacts with side chains at the -4 and -6 positions, and Trpβ6 appears to be especially important for binding. Unlike for Erbin-PDZ, the side chain of Argβ4 points toward helix α2, and the side chain of Trpβ6 lies toward strand β2 and occupies a position similar to that of Trpβ4 in the Erbin-PDZ1 ligand.

In both liganded ZO1-PDZ1 structures, the interactions with the peptide are virtually identical in this region, and it appears that the last three residues of the ligand function as a unit that interacts with the PDZ domain through both side chain and main chain interactions. The main chain nitrogen and oxygen atoms of Argβ4 are hydrogen-bonded to Oα and Nα of Asn(β2:β3-6), respectively, whereas the main chain nitrogen of Argβ5 participates in coordination of a water molecule with Nα of the same Asn and the main chain oxygen of Pro(β2:β3-7). In addition, the aromatic side chain of Phe(β2:β3-9) is in van der Waals contact with the peptide bond between Argβ5 and Trpβ6, highlighting the rearrangement within the β2:β3 loop that occurs upon ligand binding (Fig. 3B). In contrast to their main chain atoms, the side chains at the -4 and -5 positions are not intimately involved in contacts with ZO1-PDZ1, and the guanidinium groups of both arginines have very high temperature factors, indicating that these side chains are rather mobile. The side chain of Argβ5 is completely exposed to solvent and does not form close interactions with the PDZ domain, whereas the aliphatic portion of Argβ4 is in proximity to the side chains of Asn(β2:β3-6) and Hisα(α2-1). A lack of interactions involving Argβ5 is in good agreement with the promiscuity of ZO1-PDZ1 with respect to this ligand position, but the structural data do not explain the preference for residues with a general hydrophobic character in position-1 (6). The aromatic side chain of Trpβ6 is buried in a deep pocket formed by Val(β3-2) and the aliphatic portions of Ser(β2-4), His(β2:β3-8), and Ser(β3-4). Interestingly, this pocket is not visible in the unliganded structure but, rather, forms upon ligand binding as a result of rearrangements in the β2:β3 loop (Fig. 3B).

**DISCUSSION**

It has become apparent that PDZ domains recognize C-terminal ligands through interactions with a core ligand motif consisting of the last four residues and that affinity can be further enhanced by auxiliary interactions with residues upstream of the core motif (25–33). According to the structural and functional analyses described here and in the accompanying article (6), we propose a division of the functional elements within the PDZ domain structural framework into three groups on the basis of their contributions to ligand recognition. We have designated these groups as “primary,” “secondary,” and “tertiary” determinants of specificity (Fig. 5). The primary functional determinants are contained in an eight-residue stretch that includes the carboxylate binding loop and strand β2 (β1:β2-6 to 8 and β2-1 to 5). This functional element forms interactions with the C-terminal carboxylate and the main chain of the four C-terminal residues of the bound peptide. In addition, the hydrophobic side chains at β2-1 and β2-3 contribute to the recognition of the ligand side chain at position-1. The main chain of strand β2 is important for recognition of the side chain at the -1 position, and the side chain at β2-4 can also interact with the side chain at position-3. Importantly, the sequence of this primary functional element is highly conserved among all PDZ domains, and thus it represents an essentially invariant core of PDZ domain function.

The primary functional element is sandwiched on either side by secondary functional elements that account for the remaining interactions necessary for recognition of the core ligand...
motif. On one side, the side chain at a key position within strand $\beta 3$ ($\beta 3-5$) is poised to contribute to both site $^{-1}$ and site $^{-3}$. Furthermore, the side chain at $\beta 3:a1-1$ can contribute to site $^{-1}$ recognition. In the crystal structure of the PDZ domain from Shank1, a salt bridge is observed between Asp($\beta 3:a1-1$) and Arg$^{-1}$ of its ligand (25). In essence, interactions involving these side chains expand upon the “default” ligand recognition provided by the primary functional element. For example, Erbin-PDZ and many other PDZ domains prefer a Trp$^{-1}$ (6, 42). This is the default residue in this site, as it forms interactions mostly with the main chain of strand $\beta 2$ and does not rely on contacts with side chains in strand $\beta 3$. On the other hand, Erbin-PDZ prefers a Glu/Asp$^{-3}$ because of favorable interactions with Arg($\beta 3-5$). In contrast, an exchange of Arg to Asp at position $\beta 3-5$ in ZO1-PDZ1 relative to Erbin-PDZ expands the specificity of ZO1-PDZ1 site $^{-1}$ to include Tyr$^{-1}$ in addition to Trp$^{-1}$, as the Asp($\beta 3-5$) side chain is able form favorable interactions with the side chain of Tyr$^{-1}$. This exchange also alters site $^{-3}$ specificity to favor either Thr/Ser$^{-3}$ through interactions with Ser($\beta 2-4$) or Arg/Lys$^{-3}$ by the formation of electrostatic interactions with Asp($\beta 3-5$). On the other side of the binding groove, the secondary functional elements are contained within helix a2. The residues in positions a2-5 and a2-8 cooperate with the primary functional elements to form the site$^0$ pocket, whereas positions a2-1 and a2-5 are the key binding determinants for site $^{-2}$. Residues upstream of position $^{-3}$ constitute an auxiliary ligand motif, as interactions involving this region modulate binding affinity for ZO1-PDZ1 and all of the LAP family PDZ domains (6). In both Erbin-PDZ and ZO1-PDZ1 (Fig. 4C), and likely many other PDZ domains (31, 33), the recognition of this auxiliary ligand motif is mediated primarily by the $\beta 2$:$\beta 3$ loop. This region constitutes a tertiary functional element (Fig. 5) that interacts with the auxiliary motif and supports the binding of the core motif. The $\beta 2$:$\beta 3$ loop is highly divergent in sequence, length, and main chain conformation among different PDZ domains, making it difficult to develop general rules that correlate primary sequence with specificity. The structures of Erbin-PDZ and ZO1-PDZ1 explain the particular specificities of each domain for residues upstream of position $^{-3}$ reasonably well. However, these structures are not sufficient to discern the structural and functional roles of $\beta 2$:$\beta 3$ loops in other PDZ domains.

In summary, the structural and functional analyses presented here and in the accompanying article (6) provide a detailed view of how subtle changes in PDZ domain structure can have a profound impact on ligand specificity and biological function. In the particular case of Erbin and ZO-1, it is striking how a few key sequence differences have endowed ZO-1 with an expanded binding profile that in turn broadens its range of biological interactions. As a result, ZO-1 assumes a bifunctional role that involves localization to either adherens junctions or tight junctions under different conditions, whereas Erbin is restricted to adherens junctions. In more general terms, our study provides a structural framework for understanding how the diverse members of the PDZ domain family have adapted to different biological roles.

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