**Complete integrin headpiece opening in eight steps**

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Carefully soaking crystals with Arg-Gly-Asp (RGD) peptides, we captured eight distinct RGD-bound conformations of the \(\alpha_{\beta_3}\beta_3\) integrin headpiece. Starting from the closed \(\beta I\) domain conformation, we saw six intermediate \(\beta I\) conformations and finally the fully open \(\beta I\) with the hybrid domain swung out in the crystal lattice. The \(\beta I-\alpha I\) backbone that hydrogen bonds to the Asp side chain of RGD was the first element to move followed by adjacent to metal ion-dependent adhesion site \(\text{Ca}^{2+}\), \(\alpha I\) helix, \(\alpha I'\) helix, \(\beta 6-\alpha 7\) loop, \(\alpha 7\) helix, and hybrid domain.

We define in atomic detail how conformational change was transmitted over long distances in integrins, 40 Å from the ligand binding site to the opposite end of the \(\beta I\) domain and 80 Å to the far end of the hybrid domain. During these movements, RGD slid in its binding groove toward \(\alpha_{\beta_3}\), and its Arg side chain became ordered. RGD concentration requirements in soaking suggested a >200-fold higher affinity after opening. The thermodynamic cycle shows how higher affinity pays the energetic cost of opening.

**Introduction**

Among cell surface receptors, integrins undergo the most complex and longest-range conformational changes currently known. These changes function to transmit bidirectional signals over long distances between the ligand-binding integrin headpiece and the actin cytoskeleton. Integrins thus are able to communicate binding to the extracellular matrix or ligands on other cells to the actin cytoskeleton, to discriminate against soluble ligands, and to bind only with high affinity to cell surface or matrix-bound ligands (Luo et al., 2007; Springer and Dustin, 2012).

Integrins contain \(\alpha\) and \(\beta\) subunits. The \(\alpha\) subunit \(\beta\)-propeller and thigh domains and the \(\beta\) subunit \(\beta I\) hybrid, PSI (plexin, semaphorin, and integrin), and I-EGF-1 domains form the ligand-binding headpiece, i.e., the head and the upper legs (Fig. 1). The \(\alpha\) subunit calf-1 and calf-2 and the \(\beta\) subunit I-EGF-2 to I-EGF-4 and \(\beta\) tail domains form the lower legs (Fig. 1; Xiong et al., 2001; Zhu et al., 2008).

Two distinct types of global conformational changes occur in integrin extracellular domains. Extension at the knees releases integrin from a compact bent conformation (Fig. 1, A, B, D, and E). In integrin headpiece opening, the hybrid domain swings out, the \(\beta I\) domain changes from closed to open conformation, and affinity for ligand increases (Fig. 1, B, C, E, and F; Takagi et al., 2002; Luo et al., 2007; Springer and Dustin, 2012).

\(\beta I\) domains, present in all integrin \(\beta\) subunits, transmit conformational change from their interface with the swinging hybrid domain to a ligand binding site on an interface with the \(\alpha\) subunit \(\beta\)-propeller domain (Fig. 1, A–F; Xiao et al., 2004). The \(\beta I\) domain divides the hybrid domain into N- and C-terminal sequence segments. Activation at the metal ion-dependent adhesion site (MIDAS) in the \(\beta I\) domain ligand binding site is communicated to the opposite end of the \(\beta I\) domain by \(\alpha 7\) helix pistoning at the C-terminal connection to the hybrid domain (Fig. 1, B, C, E, and F). Pivoting about the N-terminal connection causes the hybrid domain to swing away from the \(\alpha\) subunit thigh domain, with an increase in separation at the \(\alpha\) and \(\beta\) knees of 70 Å (Takagi et al., 2002; Xiao et al., 2004). Swing out is readily visualized in solution by small angle x-ray scattering for both headpiece fragment (Mould et al., 2003b) and intact, detergent-soluble integrin (Eng et al., 2011) and at typical negative-stain EM or tomography resolution of ~25 Å for intact integrins or their ectodomain or headpiece fragments (Takagi et al., 2002, 2003; Iwasaki et al., 2005; Eng et al., 2011; Shi et al., 2011; Wang et al., 2012; Yu et al., 2012).

The evidence for integrin headpiece opening and its association with the high affinity state of integrins is extensive.
RGD opens the headpiece of αβ₁ ectodomain (Takagi et al., 2002). RGD and fibronectin open the αβ₁ headpiece (Mould et al., 2003; Takagi et al., 2003), and an allosteric, inhibitory Fab lowers affinity for fibronectin by stabilizing the closed headpiece (Luo et al., 2004b; Nagae et al., 2012). The αmβ₃ headpiece cocrystallizes with ligands and pseudoligands with an open headpiece (Xiao et al., 2004) and in their absence with a closed headpiece (Zhu et al., 2010). RGD mimetics increase the Stokes radius of the αmβ₃ headpiece by the amount predicted from the hydrodynamic radii of the crystal structures of the closed and open headpieces (Zhu et al., 2010). Headpiece opening induced by ligand mimetics has been demonstrated with detergent-soluble intact αmβ₃ using EM (Eng et al., 2011), small angle x-ray scattering (Eng et al., 2011), and electron tomography (Iwasaki et al., 2005). Antibodies that stabilize the closed and open headpieces of β₃ integrins show that the open conformation is required for cell adhesion and high affinity (Chen et al., 2010; Schürpf and Springer, 2011).

The equilibria between integrin extension and headpiece opening are linked because the hybrid domain, as part of the upper β leg, participates in interfaces with the lower legs that stabilize the bent conformation (Fig. 1). Thus, induction by ligand of headpiece opening in intact integrins or ectodomain fragments is always accompanied by integrin extension. Furthermore, because the lower α and β legs are very close in the ectodomain, and the α and β subunit transmembrane (TM) domains associate with one another, the equilibria governing integrin TM domain association, extension, and headpiece opening are all linked (Takagi et al., 2002).

The complexities of linked equilibria, the lack of a clear order to the steps in integrin activation (Takagi et al., 2002), and the complexity of the structure of the integrin ectodomain itself have all contributed to the difficulty of comprehending the structural basis of integrin function in the cell biology community. Currently, the concept that the open integrin headpiece corresponds to the conformation with high affinity for ligand is well accepted for β₁, β₂, β₆, and β₃ integrins, as referenced in the preceding paragraphs. However, this concept remains controversial in the β₃ integrin field (Adair et al., 2005; Xiong et al., 2009) and has resulted in a lively dialog in the literature on αmβ₃ and αβ₃ integrins. One idea, based on the observation that more metal ion sites in the βI domain are occupied in αβ₁ crystals after soaking with ligand than in absence of ligand, is that metal ion binding could somehow regulate ligand binding (Xiong et al., 2002). However, protonation of metal-coordinating Asp and Glu residues at the low pH at which αβ₃ crystallizes is probably responsible for variable absence of bound metal (Dong et al., 2012), and unliganded, closed αmβ₃, which crystallizes at higher pH, has all metal sites bound (Zhu et al., 2008, 2010). Arguments for regulation by a deadbolt and then an interface between the α and β knees have been raised (Gupta et al., 2007; Xiong et al., 2009) and contradicted (Zhu et al., 2007a; Smagghe et al., 2010; Xie et al., 2010; Dong et al., 2012). In contrast, mutational studies that shorten the βI α7 helix (Yang et al., 2004) or introduce disulfide bonds (Luo et al., 2004c; Kamata et al., 2010), N-glycan wedges (Luo et al., 2003, 2004b), or point mutations (Mould et al., 2003a; Barton et al., 2004; Luo et al., 2009) consistently show that the closed and open headpiece conformations of intact β₃ integrins on cell surfaces have low and high affinity for ligand, respectively.

Soaking RGD peptides into closed, bent αβ₃, ectodomain and α7β₁, closed headpiece crystals has in each case revealed only a single intermediate and a conformation nearer to closed than open (Xiong et al., 2002; Nagae et al., 2012). Although the βI domain adopted an intermediate conformation in these studies, little α1 helix movement occurred, the α7 helix did not piston, and the hybrid domain did not swing out; thus, the headpiece remained in an overall closed conformation. If ligands added to crystals were able to drive conformational change all the way to the open headpiece, this would provide incontrovertible thermodynamic evidence that RGD peptides bind with higher affinity to the open than closed headpiece conformation.

With purified integrins, conformational change is most readily studied by adding an excess of ligand; however, because of the principle of reversibility of chemical reactions, the same pathway will mediate change in both the outside-in and inside-out directions. Because the difference is so large between the closed and open headpiece, structures trapped at various positions along the pathway between these states would provide important information about how such a large conformational
change is accomplished. In this study, we soaked RGD peptide into closed αmβ3 headpiece crystals under different conditions. We captured RGD peptide bound to the closed headpiece, six intermediate states between closed and open, complete conversion of the βI domain to the open state, and swing out of the hybrid domain in the crystal lattice.

Results

Soaking RGD peptide into closed headpiece crystals of αmβ3

In the closed αmβ3 headpiece crystal form studied here, two independent molecules are present in the asymmetric unit. Each molecule contains the αth subunit β-propeller domain bound to 10E5 Fab and the β3 subunit PSI, hybrid, βI, and I-EGF-1 domains (Zhu et al., 2010). Crystals were soaked with or without GRGDSP peptide in buffer containing either 5 mM Mg2+ or 2 mM Mn2+ (Mn/Ca or 2 mM Mn2+/0.1 mM Ca2+ (Mn/Ca; Table 1). Crystals were stable after soaking for up to 24 h in 10 mM GRGDSP and Mg/Ca; furthermore, diffraction revealed bound peptide. However, soaking with 0.34 mM peptide for 72 h did not show bound ligand. In contrast, strong electron density for ligand was observed after soaking a preformed open αmβ3 headpiece crystal with 0.05 mM peptide (Springer et al., 2008).

Soaking in 10 mM GRGDSP and Mn/Ca for 4 h resulted in no visible cracks in some crystals and microscopically visible cracks in others. Some of these crystals yielded excellent diffraction (Table 1). All crystals had cracks after 6 h, and some began to dissolve. At 12 h, all crystals dissolved. Soaking for 0.5, 1, 1.5, and 2 h failed to yield similarly good diffraction, perhaps because conformational state was not homogenous in the crystal. Therefore, in Mn/Ca, we chose 4 h as an optimum time point and varied the concentration of RGD peptide from 1 to 10 mM (Table 1). Diffraction data from a large number of soaked crystals were examined. A single round of refinement was sufficient to show whether electron density at the ligand binding site corresponding to bound peptide was present. The best diffracting crystals from six soaking conditions were fully refined from 2.35- to 3.00-Å resolution with Rfree ~0.22 (Table 2).

All crystals were isomorphous with starting crystals, except for crystals soaked with 10 mM GRGDSP in Mn/Ca. These crystals shrunk 26 Å along the a axis (Table 2), correlating with swing out of the hybrid domain in one molecule in the asymmetric unit (see Hybrid domain swing out in the crystal lattice in Results). An almost identical change in unit cell dimension occurred in a crystal soaked with 10 mM of a one-residue-longer GRGDSPK peptide in Mn/Ca, and it also showed hybrid domain swing out; however, because resolution was lower at 2.75 Å and the C-terminal Lys residue of the GRGDSPK peptide was not resolved, this crystal was not fully refined.

Molecules 1 and 2 in the asymmetric unit in each of six crystals result in 12 examples of the αmβ3 headpiece bound to GRGDSP peptide (Table 1). Comparisons among these, with a previously published native closed-headpiece structure (Zhu et al., 2012) and with an open-headpiece structure that was co-crystallized with a ligand mimetic (Springer et al., 2008), define eight conformational states. States 1 and 8 correspond to closed and open, respectively, and states 2–7 correspond to intermediate conformations (Table 1).

Overall, there was a close correlation between soaking condition and conformational state (Table 1). The shift in conformation in Mn/Ca was clearly dependent on RGD because no conformational change was observed in Mn/Ca alone (Table 1; Fig. S1, A and B; and Fig. S2, A and B). In Mn/Ca, the conformation of molecule 2 progressively shifted more toward open between 1, 3, 5, and 10 mM RGD (Table 1). In contrast, the conformation of molecule 1 shifted immediately to state 7 at 1 mM RGD in Mn/Ca and continued to be in state 7 in 3 and 5 mM RGD until it shifted to state 6 in 10 mM RGD, coincident with the change in lattice dimensions and the swing out of the hybrid domain in molecule 2.

Insights into ligand binding

RGD binds at the interface between the αth β-propeller and β3 βI domains (Fig. 2). The Arg and Asp side chains of RGD extend linearly in opposite directions toward αth and β3, respectively (Fig. 2, B–J). A binding pocket is formed by aliphatic and aromatic side chains, water-mediated interaction with the Arg backbone carbonyl oxygen, and specific interactions with the Arg and Asp side chains. The Arg’s positively charged guanidino moiety forms a salt bridge and, in states 4–8, also forms hydrogen bonds to the side chain of αth residue Asp-224 (Fig. 2). One Asp carboxyl oxygen coordinates to the MIDAS metal ion, and depending on the conformation, the two Asp carboxyl oxygens hydrogen bond with different metal ions.

| CRITICAL | Table 1. Relation of crystal soaking conditions to conformational state |
|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|
| **GRGDSP** | **Metal ions** | **Soaking time** | **Resolution** | **Chains A + B** | **Chains C + D** | **PDB accession no.** |
| Mn or Mg | Ca | mM | mM | h | Å | (molecule 1) | (molecule 2) | |
| 0 | Mn, 2 | 0.1 | 4 | 2.45 | State 1 | State 1 | 3ZDX |
| 10 | Mg, 5 | 1.0 | 24 | 2.45 | State 2 | State 1 | 3DZ |
| 1 | Mn, 2 | 0.1 | 4 | 2.75 | State 6 | State 7 | 3DZ |
| 3 | Mn, 2 | 0.1 | 4 | 2.95 | State 7 | State 7 | 3DZ |
| 5 | Mn, 2 | 0.1 | 4 | 3.00 | State 7 | State 7 | 3DZ |
| 10 | Mn, 2 | 0.1 | 4 | 2.35 | State 6 | State 8 | 3DZ |

PDB, Protein Data Bank.

*Representative state shown in the main text figures.
to one to three backbone nitrogens of βI domain residues Tyr-122 and Ser-123 in the β1-α1 loop and Arg-214 (Fig. 2, B–J).

Electron density for the ligand in general correlated with the ligand concentration used in soaking and the amount of conformational change (Fig. 3). The real space correlation coefficient (RSCC) of electron density for Arg, Gly, and Asp residues in RGD is an estimate of occupancy and order of the ligand at the ligand binding site. The overall trend in increase of RSCC formational change (Fig. 3). Arg side chain conformations were evident in molecules 1 and 2 after soaking with 10 mM peptide for 24 h in Mg/Ca (Fig. 2, B and C). Electron density for Arg was markedly weaker in molecule 2, and a single side chain conformation was modeled, after soaking with 1, 3, and 5 mM peptide for 4 h in Mn/Ca (Fig. 2, D–F). Molecule 1 has greater accessibility of its ligand binding pocket in the crystal lattice than molecule 2, and more completely bound ligand when soaking was limited to 4 h (Fig. 3).

Strong electron densities were present for the metals at the three βI domain metal ion binding sites. When Mn$^{2+}$ was present, it largely replaced the Mg$^{2+}$ at the MIDAS and the Ca$^{2+}$ at the synergistic metal binding site (SyMBS) and adjacent to MIDAS (ADMIDAS); electron density was fit best when metals at all three sites were modeled as Mn$^{2+}$ (Fig. 2, D–I; and Fig. S1). In crystals soaked with Mn/Ca alone, the RGD-binding pockets of both molecule 1 and 2 were occupied with solvent molecules, which often occupied the same positions as polar atoms of the ligand (Fig. S1, A and B).

Remarkably, the structures reveal movements in position of the bound RGD. One of the most important movements during opening is the strengthening of hydrogen bonds between the RGD Asp side chain and the βI β1-α1 loop backbone as the distance between these elements decreases (Fig. 2 and 4).
Figure 2. The RGD-binding pocket. (A–J) The \( \alpha_{IIb}\beta_3 \) headpiece states are native closed (A; Zhu et al., 2012); the states indicated in Table 1 (B–I), and native open (J; Springer et al., 2008) are shown. Residues that contribute to the RGD-binding pocket are shown both as sticks and transparent surfaces in light blue (\( \alpha_{IIb} \)) and wheat (\( \beta_3 \)). Metal ions are shown as yellow (SyMBS and ADMIDAS) or cyan (MIDAS) spheres. Waters are smaller red spheres. GRGDSP peptides are shown in stick with green carbons. Oxygens and nitrogens are red and blue, respectively. Composite omit simulated-annealing electron density is in black mesh contoured at 3 \( \sigma \) for SyMBS and MIDAS metal ions, 1 \( \sigma \) (except 0.5 \( \sigma \) in E and F) for ADMIDAS metal ion, and 0.5 \( \sigma \) for waters and GRGDSP peptide. Hydrogen bonds and metal ion coordination bonds are dashed.
formed in the absence of ligand (closed; Protein Data Bank accession no. 3T3P; Zhu et al., 2012) and the presence of ligand (open; Protein Data Bank accession no. 2VDR; Springer et al., 2008) are included for comparison. The key transitions in each step are described as follows.

The structure in state 1 of molecule 2 in 10 mM RGD in Mg/Ca (Fig. 2 B) is remarkable because despite strong density for bound RGD, there is no significant difference from the closed conformation in absence of RGD (Fig. 5 A). One RGD Asp carboxyl oxygen directly coordinates the MIDAS Mg$^{2+}$ (Fig. 2 B) by replacing a water molecule (Fig. 2 A), whereas the other Asp carboxyl oxygen hydrogen bonds to the backbone nitrogen of Tyr-122 (Fig. 2 B and Fig. 6 A, image 1). However, the Asp side chain orients differently than in all other states (Fig. 6 A).

In state 2, the RGD Asp side chain rotates to the same orientation as in states 3–8 (Fig. 6 A, image 2). Furthermore, the $\beta$I domain $\beta$1-1 loop moves 0.5 Å toward the Asp of RGD. In states 1–3, the distances are too great for hydrogen bonds, and intervening water molecules are explicitly visible in states 1 and 2 in which RGD density is strong (Fig. 2, B and C). Arg density is weak in states 3–5 (Fig. 2, D–F), and two Arg conformations are present in state 6 (Fig. 2 G). It is only by state 7 that two strong hydrogen bonds develop between the RGD Arg and $\alpha_{\text{IIb}}$ Asp-224 side chains (Fig. 2 H and Fig. S1, C and D).

The pathway of RGD-induced headpiece opening

Our eight states reveal a detailed pathway for headpiece opening (Fig. 5 and Fig. 6). Movements occur only in the $\beta$I subunit. From state 1 to 8, the Ser-123 backbone in the $\beta$1-1 loop moves 2.2 Å, the ADMIDAS metal ion moves 3.9 Å, and the carbonyl oxygen of Met-335 moves 9.0 Å (Fig. 4). Fig. 5 shows a view of each state (Fig. 5, S1–S8) that includes the ligand Asp and the moving portions of the $\beta$I domain. Fig. 6 shows comparisons between each consecutive state, with more detail in the most relevant region. In each figure, structures from crystals formed in the absence of ligand (closed; Protein Data Bank accession no. 3T3P; Zhu et al., 2012) and the presence of ligand (open; Protein Data Bank accession no. 2VDR; Springer et al., 2008) are included for comparison. The key transitions in each step are described as follows.

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In state 3, the $\beta$1-1 loop moves still closer to the Asp of RGD (Fig. 6 A, image 3). The closer approach of the $\beta$1-1 backbone nitrogens, together with the rotation of the Asp side chain in preceding state 2, enables formation of a second backbone hydrogen bond to the RGD Asp (Fig. 6 A, image 3). These two hydrogen bonds, between each of the two Asp oxygens and the Tyr-122 and Ser-123 backbone NH groups, remain intact in all subsequent states and grow in strength with closer approach of the $\beta$1-1...
backbone to the Asp side chain carboxyl oxygens. The approach of the Ser-123 backbone to the Asp side chain is accompanied by approach of the Ser-123 side chain oxygen to the MIDAS metal ion. These two atoms form a direct coordination for the first time in state 3 (Fig. 6 A, image 3), which is maintained in all subsequent states. The Ser-123 side chain takes the position of a water, which in states 1 and 2 intervenes between the MIDAS metal ion and the Ser-123 side chain (Fig. 6 A, images 1 and 2). To follow the movement of the β1-α1 backbone, the ADMIDAS metal ion shifts 0.6 Å to maintain coordination to the Ser-123 carbonyl oxygen (Fig. 6 A, image 3). Furthermore, the β6-α7 loop moves with the ADMIDAS metal to maintain close coordination with the Met-335 carbonyl oxygen (Fig. 6 A, image 3).

In state 4, the β1-α1 loop and the ADMIDAS metal ion continue to approach the RGD Asp and MIDAS metal ion (Fig. 6 A, image 4). The RGD Asp side chain also moves, allowing RGD backbone sliding toward α5. The α1 helix is divided by a nonhelical segment into α1 and α1’ helices (Fig. 5). In state 4, the α1 portion begins to follow the movements of the β1-α1 loop and the ADMIDAS metal ion (Fig. 5 D and Fig. 6 A, image 4). The ADMIDAS metal ion continues to follow the movement of Ser-123 carbonyl oxygen and shifts an additional 1.0 Å, and ADMIDAS-coordinating residues in the α1 helix, Asp-126 and Asp-127, begin to follow the movement of the ADMIDAS metal ion (Fig. 6 A, image 4).

State 5 marks further movements of the ADMIDAS metal ion and a change in its coordination status. This metal ion now loses its coordination to the β6-α7 loop backbone; however, the β6-α7 loop maintains the position adopted since state 3 (Fig. 6 A, image 5). Furthermore, the ADMIDAS gains a direct coordination to the side chain of Asp-251 (Fig. 6 A, image 5). This direct coordination remains all the way to open state 8 and replaces an indirect coordination through a water molecule to Asp-251 in states 1–4 (Fig. 6 A).

State 6 marks the completion of the movements of the β1-α1 loop and ADMIDAS metal ion, which bring them into positions that are essentially the same as in states 7 and 8 (Fig. 5, F–H; and Fig. 6 A, image 6). The α1 helix moves in position more between states 5 and 6 than between previous states (Fig. 5, A–F), accompanied by changes in positions of ADMIDAS-coordinating α1 helix side chains Asp-126 and Asp-127 (Fig. 6 A, image 6). However, the ADMIDAS metal ion moves further than these side chains, and direct ADMIDAS metal ion coordinations are lost to Asp-126 and Asp-127. In their place, two water molecules appear that mediate indirect ADMIDAS metal ion coordinations to Asp-126 and Asp-127 (Fig. 6 A, image 6). The β6-α7 loop reverses its movement toward the ADMIDAS metal ion seen in state 3 and moves away from the metal ion, making way for the α1 helix to move toward the ADMIDAS (Fig. 6 A, image 6).

The three examples of state 7 each show a single RGD Arg side chain conformation (Fig. 2 H and Fig. S1, C and D) and essentially identical βI domain conformations (Fig. 5 G and Fig. S2, C and D). In state 7, the epicenter of conformational change shifts to the α1 helix and the β6-α7 loop (Fig. 6 B, image 2). The α1 helix with its Asp-126 and Asp-127 residues moves almost as a rigid body toward the ADMIDAS metal ion, enabling direct coordinations of each Asp side chain to reform with the ADMIDAS metal ion. Additionally, Trp-129 in the α1 helix changes rotamer and invades the space of the β6-α7 loop even more than the α1 helix backbone (Fig. 6 B, image 2). These invasions completely displace the β6-α7 loop, which flips far away to make way for the α1 helix (Fig. 6 B, image 2).

In state 8, the action shifts to the α1’ and α7 helices. Between states 7 and 8, the α1’ helix moves as a rigid body toward the α1 helix, with large movements of its buried, hydrophobic Leu-134 and Leu-138 side chains (Fig. 6 B, image 3). The loop between the α1 and α1’ helices becomes helical, and the α1 and α1’ helices with a bend between their axes merge into a single α1 helix with a single helical axis. Trp-129, near the merger, changes rotamer for the final time (Fig. 6 B, image 3). To make way for the merger of the α1 and α1’ helices, the β6-α7 loop...
reshapes, and the α7 helix pistons along its helical axis toward the hybrid domain (Fig. 6 B, image 3).

Thus, as in a series of falling dominoes, successive movements of the β1-α1 loop, the ADMIDAS metal ion, the α1 helix, and the α1′ helix propagate to the β6-α7 loop and α7 helix. Remarkably, the conformation of the β1 domain in state 8, achieved by soaking closed headpiece crystals with 10 mM RGD peptide for 4 h in Mn/Ca, is indistinguishable from
Figure 6. Detailed comparisons between nearest-neighbor states. [A] The region around the ligand Asp, β1-α1 loop, α1-helix, and β6-α7 loop where movement is greatest between states 1 and 6 (S1-S6). [B] The region around the α1 helix, α1” helix, β6-α7 loop, and α7 helix where movement is greatest between states 6 and 8. Each panel compares two nearest-neighbor states. For economy, and to compare the two rotamers of W129 in states 1–5, image 1 in B compares states 4 and 6. The carbons and metal ions of each state are in the same colors as the names of each state or the reference structures 3T3P (closed) and 2VDR (open). For clarity, water molecules as spheres and metal coordination bonds as red dashes are shown only for the second named structure in each image. Nitrogens and oxygens are shown in blue and red, respectively.
Figure 7. Hybrid domain swing out. (A–D) One integrin molecule is shown as a Ca trace, with different colors for each domain. The hybrid domain (red) and PSI and I-EGF-1 domains (yellow) are shown as thicker traces for emphasis. Other integrin molecules and all Fabs in the crystal lattice are shown as white, semitransparent, solvent accessible surfaces. The label hybrid is placed in identical positions in A–D. (A) Molecule 2 before soaking (3T3P closed structure). (B) Molecule 1 after soaking with 10 mM RGD and Mn/Ca. (C) Molecule 2 after soaking with 10 mM RGD and Mn/Ca. Composite omit simulated-annealing electron density contoured at 0.5 σ around the hybrid domain is shown as purple mesh. PSI and I-EGF-1 domains are missing in density, and superposition on the hybrid domain is used to show their approximate location in the lattice. (D) The native open headpiece (Protein Data Bank [PDB] accession no. 2VDJ) superimposed based on the β-propeller and β1 domains in C and shown in the same lattice as in C. Severe clashes are evident. (E) Superposition of αIIbβ3 headpieces. Similar regions in gray and colored shape-shifting portions in cartoon; metal ions are shown as spheres, and RGD is shown in stick. Structures are molecule 2 after soaking with 10 mM RGD and Mn/Ca (red), native closed (PDB accession no. 3T3P; blue), and native open (PDB accession no. 2VDJ; green).
the fully open state seen after cocry stallization with a ligand mimetic in Mg/Ca (Fig. 6 B, image 4). Even details of loop positions and rotamers of key side chains are identical. Opening is complete.

**Hybrid domain swing out in the crystal lattice**

β1 domain α7 helix pistoning is accommodated by swing out of the hybrid domain. Fig. 7 (A–C) shows the environments in the crystal lattice, with one integrin molecule shown as a Ca trace with each domain in a different color. Fab fragments and neighboring integrin molecules in the crystal lattice are shown as white, semitransparent, solvent-accessible surfaces. The native crystal lattice has room for the hybrid domain of molecule 2 to move to the right in the orientation shown in Fig. 7 A. In contrast, similar movement of molecule 1 is prevented by its lattice (Fig. 7 B). Indeed, soaking with 10 mM RGD in Mn/Ca for 4 h induced swing out of the hybrid domain of molecule 2 (Fig. 7 C) but not molecule 1 (Fig. 7 B). The change in position of the hybrid domain after swing out is emphasized by placing the label “hybrid” in identical positions in Fig. 7 (A–D). Although the dominant position of the hybrid domain in the lattice was clearly evident from simulated-annecing composite omit map (mesh; Fig. 7 C), this density was weaker than for the hybrid domain of molecule 1, showing some variation in the position in the crystal lattice of the swing out hybrid domain in molecule 2. Density for the PSI and I-EGF-1 domains was insufficient to include them in the molecular model. Enough space is present in the lattice to place the PSI and I-EGF-1 domains based on the position of hybrid domain, as shown by models of the PSI and I-EGF-1 in Fig. 7 C. Further, outward swing of the hybrid, PSI, and I-EGF-1 domains appears to be constrained by the crystal lattice (Fig. 7 C).

Fig. 7 E compares the positions of the hybrid domain obtained in previous closed (Fig. 7 E, blue) and open (Fig. 7 E, green) crystal structures with that attained here by soaking (Fig. 7 E, red). Binding of RGD has clearly induced swinging of the hybrid domain toward its position in open crystal structures. However, further movement to the same open position as seen when the headpiece is cocry stallized with ligand results in severe clashes with the lattice. This is illustrated by superposition of the open conformation seen in cocry stallization into the lattice, which shows that much of the ribbon trace of the hybrid domain is obscured by the semitransparent surface of the lattice (Fig. 7 D). Clashes would be even more severe for the other molecule in the lattice (Fig. 7 B) were its hybrid domain to swing out, consistent with cracking and dissolution of crystals soaked for 6 and 12 h.

**Discussion**

Previous crystal structures of integrin α7β3 defined two conformational states of its headpiece, closed and open. The same closed headpiece conformation was seen in crystals of the complete α7β3 ectodomain without Fab (Zhu et al., 2008) and in crystals of the α7β3 headpiece complex with 10E5 Fab identical to those studied here (Zhu et al., 2010). Similarly, the same open conformation of the α7β3 headpiece was seen with a cacy clolate pseudoligand or RGD mimetics, with or without bound 10E5 Fab (Xiao et al., 2004; Springer et al., 2008). Fabs have no carbohydrate and minimal flexibility compared with integrins and can stabilize crystal lattices and improve resolution. 10E5 Fab binds to the β-propeller domain, far from shape-shifting portions of the β subunit, and has no influence on β-propeller or integrin conformation as shown by comparisons among these many views of the α7β3 structure.

Conformational changes have previously been visualized by soaking RGD into crystals containing the α7β3 ectodomain (Xiong et al., 2002) or α7β3 headpiece (Nagae et al., 2012) with closed conformations. The previous RGD/α7β3 structure in Mn2+ is closest to our state 5. The RGD/α7β3 structure in Mg/Ca is closest to our states 3 and 4. These structures showed one intermediate trapped at a particular point in the shape-shifting pathway, in contrast to the eight RGD-bound states in a conformational continuum studied here.

Although the term hinge opening was used in molecular dynamics (MD) simulations of the α7β3 (Puklin-Faucher et al., 2006; Puklin-Faucher and Vogel, 2009), the simulations did not result in the open β1 domain or open headpiece conformations seen here or previously (Xiao et al., 2004). Compared with a 62° increase in angle between β1 and hybrid domains in the open headpiece crystal structure, MD showed a 23° increase starting with a model of Fn3 domain 10 bound to a RGD-ligated state 5–like structure and a 13° increase starting with an unliganded state 1 structure lacking metal ions at the MIDAS and SyMBS. The α1 and α1 helices merged and approached the α7 helix; however, merger occurred in the absence of α7 helix pistoning and displacement of β6–α7 loop residue Val-340 from its ratchet pocket and occupation of this pocket by Leu-134 in the α1/α1 helix. Because the simulations began with state 5, they were silent on RGD sliding and β1–α1 loop movement, which are largely complete by states 5–6. Dependence on RGD was not demonstrated by omitting RGD from the state 5–like structure. RGD was not sufficient for α1 and α1 helix merger because it required a modeled interaction between the body of Fn3 domain 10 and the Trp-129 side chain in the α1 helix itself. Large changes in position and rotamer of Trp-129 between our states 4, 6, 7, and 8 (Fig. 6 B) are incompatible with maintenance of a side chain hydrogen bond to the Fn3 domain 10 in the MD simulations. MD simulations of integrins are challenging because the force fields lack descriptions of the highly directional nature of octahedral metal coordination and hydrogen bonds. MD lost physiological coordinations at the MIDAS and SyMBS (Craig et al., 2004) and an invariant water at the MIDAS (Puklin-Faucher et al., 2006). Additionally, the α7β3 structures used in simulations lacked cis-Pro at β1 domain residues 163 and 169 and contained a sequence-to-structure frameshift at specifcity-determining loop residues 168–176 (Dong et al., 2012) that lie near the Fn3 docking site and α1 helix and could have affected MD results.

Surprisingly, β1 domain reshaping is related to a change in position of RGD in the ligand-binding pocket. Movement of the β1–α1 loop toward the Asp of RGD permitted the entire RGD moiety to slide in its groove away from the β3 subunit and toward the α7β3 subunit. Sliding enabled a water-mediated interaction between the RGD Arg side chain and α7β3 Asp-224 to be converted...
to a much stronger direct, charged interaction through multiple hydrogen bonds. Sliding also correlated with a shift from multiple Arg side chain conformations to a single conformation. Sliding and disorder or multiple conformations of the Arg side chain show that binding of the Asp of RGD to the MIDAS metal ion and the β1-α1 loop backbone is energetically more important than Arg interactions with the αtβ3 subunit.

Placing our structural snapshots into sequence in a movie provides a plausible shape-shifting pathway for conformational change in integrins (Videos 1, 2, and 3). In contrast to previous soaking studies, our state 1 structure shows that RGD can bind to αtβ3 with no appreciable change in structure. The early parts of the pathway from state 1 to 6 are captured in atomic detail as movements of <1 Å, whereas those in states 7 and 8 involve large concerted changes. Ligand-induced integrin shape shifting begins in the β1-α1 loop and then works its way more C-terminally in the α1 and α1’ helices until they finally merge and push the β6-α7 loop and α7 helix out of the way toward the hybrid domain.

Linkage between the MIDAS and ADMIDAS coordination shells grows between states 3 and 6 (Video 2). In state 3, the Ser-123 side chain replaces a water molecule in the inner MIDAS coordination sphere; the Ser-123 backbone oxygen directly coordinates the ADMIDAS through states 1–8. In state 5, the ADMIDAS metal ion directly coordinates the Asp-251 side chain. In state 6, the same Asp-251 oxygen indirectly coordinates the MIDAS through the Ser-123 side chain; the other Asp-251 oxygen indirectly coordinates the MIDAS through a water molecule throughout states 1–8 (Fig. 6). By state 6, interaction between the MIDAS and ADMIDAS coordination shells is strong enough to leave behind the direct ADMIDAS coordinations to the Asp-126 and Asp-127 side chains, which are reformed in state 7.

Saltatory motions between states 1 and 7 thus include numerous steps of removal, addition, and removal yet again of waters that provide indirect MIDAS and ADMIDAS coordinations to side chains. Another movement not predicted by structural interpolation is that of the β6-α7 loop. Its coordinating Met-335 carbonyl oxygen moves toward the fleeing ADMIDAS in state 3 and stays in this new position until state 6, when it moves in the opposite direction.

Presumably, the movement of the β1-α1 loop places strain on the α1 helix, which contains ADMIDAS-coordinating residues Asp-126 and Asp-127. In state 7, there is a large rigid-body movement of the α1 helix, as Asp-126 and Asp-127 catch up with and reform direct coordinations to the ADMIDAS. In turn, the movement of the α1 helix is likely to place strain on the α1’ helix. In final state 8, the α1’ helix catches up with and joins to the α1 helix. Alignment of the α1 and α1’ helices squeezes at their junction the β6-α7 loop. The hydrophobic ratchet pocket occupied by β6-α7 residue Val-340 in states 1–6 is occupied by α1 helix residue Leu-134 in state 8, with Val-340 in an intermediate position in state 7 (Fig. 6). Merging of the α1 and α1’ helices appears to be the final straw that breaks the camel’s back and pushes the α7 helix toward the hybrid domain, causing it to swing out. The side chain of Trp-129 near the end of the α1 helix appears to buffer the large α1 and α1’ conformational movements between states 6 and 8. Between states 1 and 6, the Trp-129 side chain can occupy either of the two rotamers shown in Fig. 6 B (image 1). In state 7, it adopts a very different buried rotamer and acts as a placeholder for the α1’ helix. Then, in state 8, Trp-129 moves outward again and adopts yet another rotamer to make way for the merge of the α1 and α1’ helices and the side chain of Leu-134.

These movements illustrate the separability yet interdependence of the MIDAS and ADMIDAS coordination shells, the β1-α1 loop, α1 helix, α1’ helix, and β6-α7 loop. The final movements of the α1’ helix and α7 helix are highly concerted (Video 3). Compared with the earlier movements in states 1–6, those of the α1 helix and β6-α7 loop in state 7 are much larger, whereas the movements in the α1’ helix, β6-α7 loop, and α7 helix in state 8 are cataclysmic. The extraordinary movements that follow state 7 likely correspond to a large decrease in free energy and suggest that state 7 must be close to the transition state for conformational change.

Among the caveats in our study is the difficulty of modeling the segments undergoing conformational change as single conformations. Particularly for the β6-α7 loop in state 7, density is consistent with multiple backbone conformations. Resolution of most of our structures was sufficient to model alternative conformations of the side chains of some residues; however, attempts to include alternative backbone conformations did not improve model fit to density. In general, the quality of the density in our structures is high. It is difficult to explain the gradual shift in positions described here in each structure as an averaging of structural ensembles that change in occupancy and not in position. Further evidence against this idea is the saltatory nature of many of the movements.

Another caveat is that the states seen here are trapped in crystals and may be perturbed by lattice contacts. However, changes occurred in two molecules in different lattice environments. Furthermore, despite presence of states 2, 6, and 7 in molecule 1 and states 1, 3, 4, 5, and 8 in molecule 2, these states fall on a single pathway of headpiece opening. Constraints limiting conformational change differ in crystals and on cell surfaces. Nonetheless, hybrid domain swing out is the largest motion in headpiece opening and thus may be rate limiting for conformational change on cell surfaces as well as in crystals. Furthermore, the different moving elements, i.e., α1 helix, α1’ helix, β6-α7 loop, α7 helix, and hybrid domain are key on cell surfaces as well, as shown by mutational studies in these elements (Luo et al., 2003, 2004b,c, 2009; Mould et al., 2003a; Barton et al., 2004; Yang et al., 2004; Kamata et al., 2010).

It is extremely difficult to obtain information on conformational change on cell surfaces at any similar resolution, and structures determined here will likely offer the best models for understanding integrin allostery on cell surfaces for many years. The principle of the reversibility of chemical reactions implies that inside-out activation of integrins may proceed by a similar pathway as outside-in activation studied here but in the opposite direction.

Two alternative mechanisms for achieving conformational change in proteins, selection by ligand of preexisting conformational states and ligand-induced fit (Henzler-Wildman and Kern, 2007), are linked in a thermodynamic cycle (Fig. 8). In the crystals studied here, no conformational change is required.
for RGD binding to state 1, and thus, induced fit (Fig. 8, k₁ and k₃) is the mechanism driving conformational change. However, on cell surfaces, interconversion between conformational states driven by thermal motion (Fig. 8, k₁ and k₃) will be far faster than in crystals, and inside-out signals may also increase k₃. Therefore, the relative fluxes of integrins to ligand-bound states on cell surfaces through induced fit (k₁ and k₃), and selection of preexisting states (k₁ and k₃) will depend on their relative rates, as well as ligand concentration, and remains to be determined.

RGD must bind with higher affinity to the open than closed headpiece to drive conversion to the higher energy open headpiece conformation. This may be formally demonstrated using the thermodynamic cycle (Fig. 8). The difference in energy between any two states in this cycle is identical, whether conversion occurs by clockwise or counterclockwise routes. Therefore, ∆Gopen+RGD = ∆Gclosed+RGD + ∆Ginduced fit - ∆Gconf. Crystal soaking experiments demonstrate that ∆Ginduced fit is highly negative. A large number of experiments referenced in the Introduction demonstrate that in absence of ligand, [Closed] >> [Open]; therefore, ∆Gconf is highly positive. It follows from the aforementioned equation that ∆Gopen+RGD < ∆Gclosed+RGD; i.e., that the affinity of the open headpiece is much higher than that of the closed headpiece for RGD.

Conclusions based on soaking experiments in Mn/Ca and the thermodynamic cycle are supported by soaking results in Mg/Ca (Fig. 3). Based on soaking with 10 mM RGD in Mn/Ca for different times, near-equilibrium concentrations of RGD should have been reached inside the crystal lattice after soaking with 10 mM RGD in Mg/Ca for 24 h. Incomplete occupation of the ligand binding site in Mg/Ca as shown by the RSOC values in Fig. 3 suggests an affinity insufficient to saturate binding in 10 mM RGD, consistent with lack of detectable binding after soaking for 72 h with 0.34 mM RGD in Mg/Ca. In contrast, when crystals with the open headpiece were soaked for 96 h with 0.05 mM RGD in Mg/Ca, RGD peptide completely replaced a cacodylate pseudo-ligand (Fig. 3; Springer et al., 2008). Complete saturation of the open headpiece at 0.05 mM RGD and incomplete saturation of the closed headpiece at 10 mM RGD support the conclusion that the affinity of the open α₃β₃ headpiece for RGD peptide is ≥200-fold higher than that of the closed headpiece in Mg/Ca.

The ability of the lattice around molecule 2 to hold the integrin headpiece in the closed conformation after binding RGD in Mg/Ca provides a thus far unique estimate of affinity of the closed headpiece for RGD. This affinity is between 0.34 and 10 mM and closer to 10 mM. It is not possible to estimate the affinity of the closed headpiece for RGD from measurements of binding to integrins in solution or on cell surfaces because, as discussed in the previous two paragraphs, the higher affinity for the open headpiece is more than sufficient to pay the energetic penalty of inducing or selecting the open headpiece conformation. Conversion to the open headpiece is consistent with 50% effective concentration estimates in the range of 7 µM to 1 mM RGD in Mg/Ca for binding of antibodies to ligand-induced binding sites or increased protease sensitivity using isolated, intact α₃β₃ or α₅β₃ and αvβ₅ on cell surfaces (Parise et al., 1987; Frelinger et al., 1988, 1990). 50% inhibitory concentration values for inhibition by RGD of activated platelet binding to ligands are in the range of 0.01–0.2 mM (Plow et al., 1985), suggesting that the headpiece has been opened.

The thermodynamic cycle characterized here for headpiece fragments has wide applicability. In extended integrins (Fig. 1, B, C, E, and F), as in headpiece fragments, none of the headpiece domains are in buried interfaces, and the lower β leg is highly flexible (Fig. 1, dashed lines). Therefore, the relative energies of the four states in the cycle (Fig. 8) are expected to be very similar in headpiece fragments and in extended integrins on cell surfaces. Bent integrins on cell surfaces have extensive interfaces that are exposed upon extension; the hybrid domain is in one of these buried interfaces (Fig. 1, A and D). Also, the α and β subunit C termini are close to one another in the bent conformation, and the TM domains associate (Luo et al., 2007; Springer and Dustin, 2012). Because headpiece opening exposes most of the same interfaces that become exposed upon integrin activation, ∆Ginduced fit and ∆Gconf in Fig. 8 (which each require headpiece opening) are higher (by similar amounts) for cell surface integrins than for headpiece fragments. If one is willing to accept that ligand-induced binding site epitopes in β₃ integrins measure either integrin extension induced by headpiece opening or headpiece opening itself, one could conclude that ∆Ginduced fit is negative and ∆Gconf is positive (as for integrin headpieces) and deduce that the open headpiece conformation has higher affinity for RGD than the closed headpiece conformation in intact integrins, just as demonstrated here for headpiece fragments.

A remaining question is whether headpiece opening upon ligand binding is general for integrins or is dependent on the integrin or the ligand. MAdCAM-1 binding to the αβ₃ headpiece
can yield either an intermediate or open state as seen by EM (Yu et al., 2012), which appear to mediate rolling adhesion (intermediate affinity) and firm adhesion (high affinity), respectively (Chen et al., 2004). Thus, there may be differences among integrins. However, headpiece opening has been demonstrated for all integrins yet tested, including those containing the β1, β2, β3, β6, and β3 subunits (Shi et al., 2011; Springer and Dustin, 2012; Yu et al., 2012). An obvious exception would be when a carboxyl group is absent in the ligand because this is the moiety that interacts with the integrin β subunit in which allostery occurs. Thus, antagonists that mimic the Arg moiety of RGD and do not bind the MIDAS, or instead displace Mg2+ from the MIDAS, do not activate opening and stabilize the closed headpiece against opening, respectively (Zhu et al., 2010, 2012).

Is binding to the MIDAS alone sufficient to open the headpiece, or must another moiety such as the Arg of RGD pull the βI domain β1-α1 loop toward the α subunit to open the headpiece? The primary role of the Asp shown here in allostery suggests that Arg might not be required. Some small molecules based on RGD, selected by the pharmaceutical industry for their ability to bind equally well to the low and high affinity states of αIIbβ3, are reported not to induce reactivity with antibodies to ligand-induced binding sites (Aga et al., 2004) and, therefore, appear not to induce headpiece opening. These mimetics have unique chemical features that may set them apart from the Asp and Glu side chains present in physiological integrin ligands.

Further investigation is important of the principles that ligand binding induces integrin headpiece opening and that the open headpiece corresponds to the high affinity state of integrins. The large movement at the integrin knees upon headpiece opening is thought to be important for transmission of allostery through long legs that are flexible except when elongational force is applied in cell adhesion (Zhu et al., 2008; Springer and Dustin, 2012). We have shown here a remarkable pathway for transmission of signals in a cell surface protein with an exceptionally complex and large structure. A movement of only 2 Å at the β1-α1 loop in the ligand binding site is transmitted through an intricate shape-shifting pathway a distance of 40 Å across the βI domain. A 10-Å a 7 helix movement like that of a connecting rod in the βI domain causes the hybrid domain to swing out by pivoting at its other connection to the βI domain. The length of the hybrid domain is 40 Å, and the PSI and I-EGF1 domains attached at the lower end opposite the βI domain make the total length of the upper integrin β leg ~70 Å. The leverlike swing of the upper leg amplifies the 2 Å movement of the β1-α1 loop to a 75-Å increase in separation at the integrin knees (Fig. 1). The integrin headpiece is a marvelous biological machine that appears designed for the difficult job of transmitting allostery in extracellular environments.

If in general the open headpiece is the high affinity state of integrins, induction by inside-out signals of headpiece opening can be a general mechanism for integrin activation on cell surfaces.

**Materials and methods**

**Protein expression, purification, and crystallization**

The soluble αIIbβ3 headpiece construct was expressed in CHO-Lec cells and purified using its His6 tag, treated with chymotrypsin and carboxypeptidase to remove the thigh domain as well as the C-terminal acid-base coded coil and His6 tag, and further purified exactly as described in detail in the supplemental materials in Xiao et al. (2004). Crystallization in complex with 10E5 Fab in the absence of ligand was as previously described (Zhu et al., 2010); purified αIIbβ3-10E5-Fab complex at 10 mg/ml was crystallized at 4°C in hanging drops by adding an equal volume of 11–13% PEG 8000, 0.2 M ammonium sulfate, and 0.1 M Tris-HCl, pH 8.9. Crystals did not always appear under these conditions. In this case, crystal seedling was induced. Seeds were obtained by breaking a single crystal with a seed bead (Hampton Research). 0.1-µl diluted crystal seeds were added to 0.9 µl of a hanging drop that had preequilibrated overnight. Crystals were grown at 4°C for >1 mo before soaking.

**Crystal soaking and diffraction data collection**

Crystals were harvested in cryosolution containing 15% PEG 8000, 0.2 M ammonium sulfate, and 0.1 M Tris-HCl, pH 8.9, with the addition of glycerol as a cryoprotectant in 5% increments up to a 20% final concentration. Harvested crystals were soaked in the same cryosolution plus desired concentrations of GRGDSP peptide and metal ions shown in Table 1 for the indicated times at 4°C before plunge vitrification in liquid nitrogen. Diffraction data were collected at beamline 23-ID (Advanced Photon Source) and processed with MOSFLM (Leslie and Powell, 2007) and SCALA (Evans, 2006) or the XDS package (Kabsch, 2001). Resolution cutoff was chosen to give ε(ω) > 1.6 in the highest resolution shell.

**Structure refinement**

For soaked crystals with little difference in the unit cell compared with the nonsoaked crystal, we used the closed-headpiece αIIbβ3-10E5-Fab structure (Protein Data Bank accession no. 3NIQ; 2.25 Å) as the starting structure and subjected each domain to rigid body refinement with PHASER (Adams et al., 2010). Electron densities at the ligand binding pocket corresponding to RGD peptides were readily identified after the first round of rigid body refinement. For soaked crystals with 10 mM RGD peptide in Mn2+, the unit cell shrunk in dimension, and structures were solved by molecular replacement with PHASER (McCoy et al., 2007) using the αIIb-propeller and β3 domains as search models. Structures of soaked ligands were built into their electron densities with COOT (Emsley and Cowtan, 2004). The complete structures were then finalized by several rounds of rebuilding with COOT and refinement with PHENIX (Adams et al., 2010). High resolution αIIbβ3 structures (3T3P, 2.2 Å; or 3NIQ, 2.25 Å) were used as reference structures when rebuilding the lower resolution soaked structures with COOT. Metal ion coordination restraints were used for MIDAS and SynB5, but not ADMAS metals, during refinement with PHENIX. Because the two molecules of the asymmetric unit were trapped in different conformations after crystal soaking, noncrystallographic symmetry restraints were not used during structure refinement. The structures were validated with the MolProbity server (Davis et al., 2007).

**Simulated-annealing composite omit maps and RSCCs**

Both simulated-annealing composite omit maps and RSCCs with simulated-annealing composite omit density were calculated by PHENIX (Adams et al., 2010). When dual conformations were present, the structures were refined with either conformation or a third intermediate conformation as a single conformation, and the highest RSCC values were plotted. Although RSCC values provide a good estimate of occupancy, they also are dependent on the resolution and quality of the diffraction data and refinement.

**Superpositions and figures**

All superpositions were on Cα atoms of the αIIb β-propeller domain and nonmoving portions of the β3 βI domain. Panels in multipanel figures (Fig. 2, Fig. 4, Fig. 6 [A and B], and Fig. 7 [A–D]) are shown in identical orientations and are aligned vertically and horizontally on the page. Figures were prepared with PyMOL (Schrödinger, LLC). All main text figures use the states indicated in Table 1.

**Online supplemental material**

Fig. S1 shows electron density map and the RGD-binding pocket for examples of states 1 and 7 not depicted in Fig. 2. Fig. S2 is an overview of the moving portions of the βI domain for examples of states 1 and 7 not depicted in Fig. 5. Video 1 shows an overview of headpiece opening. Video 2 shows a detailed overview of the ligand and metals. Video 3 focuses on the αI and αI’ helices, β6α7 loop, and α7 helix. Online supplemental material is available at http://www.jcb.org/cgi/content/full/jcb.201212037/DC1.

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Eight snapshots of integrin headpiece opening Zhu et al.


