Eukaryotic protein kinases (EPKs) constitute a class of allosteric switches that mediate a myriad of signaling events. It has been postulated that EPKs’ active and inactive states depend on the structural architecture of their hydrophobic cores, organized around two highly conserved spines: C-spine and R-spine. How the spines orchestrate the transition of the enzyme between catalytically uncommitted and committed states remains elusive. Using relaxation dispersion nuclear magnetic resonance spectroscopy, we found that the hydrophobic core of the catalytic subunit of protein kinase A, a prototypical and ubiquitous EPK, moves synchronously to poise the C subunit for catalysis in response to binding adenosine 5′-triphosphate. In addition to completing the C-spine, the adenine ring fuses the β structures of the N-lobe and the C-lobe. Additional residues that bridge the two spines (I150 and V104) are revealed as part of the correlated hydrophobic network; their importance was validated by mutagenesis, which led to inactivation. Because the hydrophobic architecture of the catalytic core is conserved throughout the EPK superfamily, the present study suggests a universal mechanism for dynamically driven allosteric activation of kinases mediated by coordinated signal transmission through ordered motifs in their hydrophobic cores.

INTRODUCTION
Protein phosphorylation is the most common signaling mechanism in cell biology, regulating growth, differentiation, and metabolism. Aberrant phosphorylation causes many lethal pathologies, making kinases prime targets for drug therapy (1, 2). Cyclic adenosine 5′-monophosphate (cAMP)-dependent protein kinase A (PKA) is a benchmark for the entire eukaryotic protein kinase (EPK) superfamily. The bilobal core of the catalytic subunit (PKA-C), conserved among all Ser/Thr- and Tyr-specific kinases (3), consists of a small N-lobe and a large C-lobe that harbors most of the catalytic residues as well as the substrate docking surface; adenosine 5′-triphosphate (ATP) binds in the cleft between the two lobes (4). The spatial architecture of the kinase core is also highly conserved and consists of two hydrophobic spines (regulatory R-spine and catalytic C-spine) (5), which serve as noncovalent links between the N-lobe and the C-lobe (Fig. 1). The structural assembly of the R-spine is a hallmark for activated kinases (5). Formation of the R-spine is highly regulated and typically requires phosphorylation of the activation loop (Thr197) in PKA-C (6), whereas the C-spine is transiently assembled by ATP binding, which is an absolute requirement for orienting the nucleotide toward the substrate for catalysis (5). Once the spines are assembled, the enzyme binds substrates cooperatively (7, 8) and transfers the γ-phosphate of ATP to Ser/Thr acceptors. The functional relevance of the spines has been validated by the discovery of oncogenic mutations, such as the T334I mutation found in chronic myeloid leukemia that stabilizes the R-spine of the c-Abl kinase, rendering it constitutively active and resistant to imatinib (9). The constitutively activating L485F and L505F mutations found in BRAF (10, 11) further support an active role for the hydrophobic spine architecture in kinase activation and regulation.

RESULTS
Here, we investigated the role of conformational dynamics in the hydrophobic core of PKA-C using methyl-TROSY (transverse-relaxation optimized spectroscopy) nuclear magnetic resonance (NMR) spectroscopy, isothermal titration calorimetry, and site-directed mutagenesis. Methyl groups are ideal probes for monitoring the conformational dynamics of hydrophobic cores because they densely populate the interior of folded proteins (12). A total of 122 methyl groups (~98% of the methyl groups from isoleucine, leucine, and valine [ILV] residues) were assigned, enabling us to comprehensively map the enzyme’s hydrophobic core in the apo, binary (nucleotide-bound), and ternary (substrate or peptide-scaffold) forms (figs. S1 to S3). As a substrate, we chose a peptide encompassing the PKA-C recognition sequence of phospholamban (PLN1–19), a central regulator of heart muscle contractility (7, 13). As a pseudosubstrate, we used a peptide corresponding to the inhibitory sequence of the endogenous protein kinase inhibitor (PKI5–24). Upon nucleotide binding, the chemical shift perturbations (CSPs) follow linear trajectories along the coordinates of the open to closed states (figs. S4 and S5). When mapped onto the three-dimensional (3D) structure (Fig. 2A), we observe several spatially contiguous paths of residues crossing the inner core of the enzyme. Although the CSPs of the amide groups reported on the effects of ligand binding on the catalytic loop and the Mg-positioning loop (Mg loop), they were essentially silent for residues in the hydrophobic core (fig. S6). In contrast, methyl groups are sensitive reporters of changes occurring throughout the entire enzyme, especially across the hydrophobic core (table S1). By fusing the β sheet of the N-lobe with β7–β8 of the C-lobe, nucleotide binding exerts global effects that not only radiate to adjacent residues, as seen in Fig. 2B (for example, L173 in β7, V57 in β2, and V104 in the cC–β4 loop), but also extend to distal sites in the hydrophobic core, including L172 and I174 in β7, L180 and L182 in β8, L95 in αC, I150 in αb, and L167 in the catalytic loop, as well as the eH and eI helices, the substrate-binding cleft, and the hub for the regulatory subunit recognition (table S1). Substrate binding (PLN1–19) shifts the enzyme’s conformational
equilibrium toward a closed state, which is fully reached when the pseudosubstrate inhibitor (PKI\textsubscript{3-24}) is bound (fig. S4).

When analyzed using statistical methods (Materials and Methods), the CSPs display a correlated behavior along the open-to-closed path (fig. S5). Fifty-eight methyl resonances interspersed throughout the enzyme’s structure display correlated linear trajectories, indicating a concerted conformational exchange between the different states (table S1, figs. S4 and S5, and Fig. 2). The correlation map indicates the presence of several clusters of residues in physical contact throughout the kinase core; however, other clusters that are apparently isolated show the same correlated conformational equilibrium. However, when both the CSPs from amide and methyl probes are combined (fig. S6), it is apparent that the conformational changes involve the enzyme globally as predicted by the molecular dynamics simulations and community map analysis (14, 15).

To determine the contribution of the conformational entropy to ligand binding, we calculated the methyl group order parameters (\(S^2\)) from \(^2\text{H}\) spin relaxation times (\(T_2\)) (16). The average \(S^2\) (\(<S^2>\)) systematically increases going from the apo to the intermediate and closed states (Fig. 3, fig. S7, and tables S2 to S5), suggesting that the enzyme’s conformational dynamics in the fast time scale (picosecond to nanosecond) plays a significant role in both nucleotide and pseudosubstrate recognition. Nucleotide binding makes the hydrophobic core of the enzyme more densely packed, with a concomitant increase of the local degree of ordering for the methyl-group bearing side chains. The C-spine and neighboring side-chain residues become less mobile when the adenine ring intercalates between A70 and V57 in the N-lobe and L173 in the C-lobe (Fig. 3B), and this locks the core into a committed conformation that is poised for catalysis. The trend and magnitude of the methyl group (\(S^2\)) do not correlate with the total entropy of binding, as seen for other systems (fig. S8) (17). This is not surprising because the backbone amide groups show a net increase in the conformational dynamics upon ATP binding (18) and solvation-desolvation effects are not accounted for. Upon pseudosubstrate binding, both backbone amide and side-chain methyl group dynamics show similar trends, with a global reduction of the internal motions (that is, dynamically quenched state).

Slow conformational dynamics (microsecond-to-millisecond time scale) of the catalytic loops are directly linked to the catalytic cycle of PKA-C (18). A dynamic knockout mutation (that is, Y204A, far from the active site) shifts the motional regime of the catalytic loops outside the microsecond-to-millisecond time scale, reducing the enzyme’s catalytic efficiency (15, 19). To assess the contribution of the hydrophobic core’s slow time scale motions to the catalytic process, we analyzed the structural fluctuations of the methyl-bearing side chains using Carr-Purcell-Meiboom-Gill (CPMG) experiments (20, 21). CPMG relaxation dispersion curves enable one to determine the exchange contribution to the transverse relaxation (\(R_{ex}\)) and quantify both the kinetics (\(k_{ex}\)) and thermodynamics (that is, populations, \(P_A\) and \(P_B\)) of the conformational equilibria (tables S6 to S10 and figs. S9 and S10). The apo enzyme shows significant \(R_{ex}\) values for several residues interspersed throughout the hydrophobic core, the C-spine (V57 and L172), the R-spine (L95), the R-spine shell (V104), and additional residues neighboring and bridging the spines (L167, I180, and I150) (Fig. 4A). Although it is possible to fit dispersion curves of isolated clusters of residues, they exchange with different rate constants, suggesting that motions in the core of the apo enzyme are essentially asynchronous. Although the nucleotide binding attenuates the magnitude of the \(R_{ex}\) values throughout the entire enzyme, significant conformational dynamics are present for residues at the interface between the two lobes. The shapes of the CPMG dispersion curves indicate that the motions become faster and (Fig. 4A) most of the residues can be clustered into a unique group with a global exchange rate constant \(k_{ex} = 2500 \pm 300 \text{ s}^{-1}\), indicating that the enzyme’s motions in the nucleotide-bound state are faster and synchronous. The residues include the C-spine (L172, L173, and L174), the R-spine (L95), and neighboring hydrophobic residues extending toward the \(\alpha C\) helix (V104, V182, I150, L180, L162, L103, and I194) (fig. S11). Upon PLN\textsubscript{1-19} binding, the conformational dynamics remain persistent for these residues and become prominent for other residues away from the substrate-binding site (table S11 and fig. S9). Residues in the C-spine and R-spine, the hydrophobic shell, and those bridging the spines moving synchronously in the nucleotide-bound state retain significant motions (fig. S11). Significant \(R_{ex}\) values were also observed for residues in and neighboring the C-spine (L227 and L49), the \(\alpha C\) helix (L89), and the activation loop (V191), and throughout the two lobes. These motions exemplify the dynamically committed state of the enzyme in the presence of both nucleotide and substrate. In contrast, PKI\textsubscript{3-24} binding quenches most of the conformational exchange in the microsecond-to-millisecond time scale in agreement with amide measurements (fig. S9) (22).
To verify the importance of these motions in the allosteric network, we individually mutated two residues, V104 and I150, which display significant $R_{ex}$ values. These residues are highly conserved throughout the EPK family (23) and form a bridge between the two spines (Fig. 4B). Community map analysis of molecular dynamics trajectories predicted that these residues are essential to the integrity of the hydrophobic core and actively participate in the dynamically driven allosteric communication (fig. S12) (14). Upon mutating these positions (V104G and I150A), the kinase does not undergo autophosphorylation at Thr197 and S338 (figs. S13 and S14) (24), confirming that the enzyme is completely inactive. Coexpressing the mutants with PDK1, a kinase that phosphorylates Thr197, PKA-C adopts an active conformation of the activation loop, where the R-spine is assembled and the $\alpha$C helix is positioned for catalysis. Remarkably, this is not sufficient to activate the kinase. Because the mutated kinases retain the overall fold, these results suggest that I150 and V104 in the inner hydrophobic core of the kinases must contribute to the intramolecular allosteric signaling for substrate recognition and catalysis.

**DISCUSSION**

The participation of protein hydrophobic cores in the transmission of intermolecular and intramolecular allosteric signaling was hypothesized decades ago based on free-energy considerations (25), and theoretical studies have hypothesized that correlated internal dynamics of hydrophobic cores may be required for biological function (26). In support of this, several proteins have been found to adopt loose packing of the inner cores, yet can still carry out biological functions with high specificity (27). This is reminiscent of the 3D conservation map of PKA-C with a firmly packed core (4) and site for phosphoryl transfer buried in the center. However, our NMR analysis reveals an unprecedented map of the dynamic network of communication within the hydrophobic core of PKA-C that orchestrates signal transmission between distal sites and encapsulates the active site. The apo enzyme with a dismantled C-spine is malleable and relatively accessible to water, displaying uncorrelated motions that are uncommitted for catalysis. On the other hand, the nucleotide-bound state, with the C-spine assembled, features synchronous motions, where the fluidity of the well-packed side chains is due to an

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**Fig. 2. Methyl CSPs define the conformational transition between open and closed states.** (A) Portions of the methyl-TROSY spectra showing the linear chemical shift trajectories upon ligand binding along the major states of the catalytic cycle. The residues with highly correlated chemical shift changes ($R^2 > 0.97$) are shown with purple spheres, whereas residues with lower correlation coefficients ($R^2 < 0.97$) are shown in white. A comprehensive map of the cross-correlation is shown in fig. S5. (B) Zoom-in of the core of the enzyme showing that the chemical shift changes with the highest correlation cluster in the proximity of the C-spine reporting on the open-to-closed conformational transition.
extended network of correlated motions (27). We speculate that both the structure and dynamics of the core and residues surrounding the active site may have coevolved to optimize the intramolecular allosteric communication (fig. S12) (28, 29).

Computer simulations provide us with an unprecedented computational microscope and generate new mechanistic concepts whereas crystallography provides us with static snapshots of the various conformational states that reflect the catalytic cycle. We used these tools to create a community map for PKA-C, which allows us to see how elements of secondary structure are woven together in new ways (14).

Using methyl relaxation dispersion NMR experiments, we can now experimentally capture the correlated internal dynamics of the hydrophobic core. The β structure of the kinase core, which embraces and surrounds the active-site cleft, provides a framework to think about how the adenine ring engages and commits the entire molecule to catalysis. The five-stranded β sheet in the N-lobe has been described extensively. It moves as a rigid body as the cleft opens and closes (30). In the apo conformation, it is not closely engaged with the C-lobe, as confirmed by earlier NMR studies (22). However, when ATP binds, the adenine ring bridges the β sheet of the N-lobe with the small β7-β8 sheets in the C-lobe (Fig. 5A). The β structure of the C-lobe received less attention, but is equally important for function. The β7 and β8 strands are directly linked to the catalytic loop and the Mg loop, respectively. The catalytic loop is anchored to the αF helix through Leu167, and this is one of the correlated residues that is quite prominent. In the assembled and active kinase, the catalytic loop is preceded by β6, whereas the Mg loop is followed by β9. The shell residues (Val104, Met11, and Met120) that bridge the R-spine and C-spine in the N-lobe have been described.

**Fig. 3.** Changes of the fast conformational dynamics of the hydrophobic core upon nucleotide and pseudosubstrate binding. (A) Order parameters of the methyl groups mapped onto the apo, binary (ATP/C-bound), and ternary (ATP/N/PKI5-24) forms of the enzyme, showing an increasing rigidification of the hydrophobic core. (B) Ordering of the C-spine, αF, and the catalytic loop upon nucleotide and PKI binding. (C) Structural details of PKA-C showing the C-spine and the αE helix, highlighting the rigidification of I150, I180, and V182 upon ligand binding and bridging β1-β2 in the N-lobe with β7-β8 in the C-lobe.
earlier, but there are two other residues (Val\textsuperscript{104} and Ile\textsuperscript{150}) that scored high on the local spatial pattern (LSP) alignment that was used to identify the spines (5). At the time, however, their functional significance was not appreciated. Both of these residues are highly correlated in the methyl-TROSY NMR analysis and both serve as additional bridging residues that link the R-spine and C-spine (fig. S13).

One additional element that was not fully appreciated in the initial analyses of the kinase structure is the aC-b4 loop, which is a classic \beta turn and also moves as a rigid body (Fig. 5B). The tip of this loop is the only piece of the N-lobe that remains anchored to the C-lobe in the open conformation (30). Val\textsuperscript{104} and Leu\textsuperscript{103} are part of this loop, and these residues are an important part of the correlated network of residues that

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**Fig. 4.** Synchronization of motions within the hydrophobic core upon nucleotide binding. (A) \textsuperscript{13}C CPMG relaxation dispersion curves carried out at two different magnetic fields (700 MHz, black symbols; 850 MHz, red symbols) for selected residues in the hydrophobic core for the apo, binary (ATP\textsubscript{C}-bound), and ternary (ATP\textsubscript{N/PKi\textsubscript{5}-24}) complexes. (B) Mapping of the methyl groups showing conformational dynamics in the C-spine, R-spine, and bridging residues of the hydrophobic core. V104 bridges the R-spine and C-spine together with the adenine ring of ATP, whereas I150 bridges the C-spine to the \(\alpha E\) helix.

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**Fig. 5.** \beta structure of the kinase core is anchored to the hydrophobic core and recruited for catalysis by ATP binding. (A) The \beta sheet of the N-lobe (b1 to b5) is anchored to the adenine ring of ATP through V57 (b2) and A70 (b3). In the C-lobe, b7 and b8 are anchored to the adenine ring through L173 (b7). The catalytic loop that spans b6 and b7 is anchored to the F helix by L167; the Mg loop (red) bridges b8 and b9. (B) Another conserved structural element in the N-lobe is the aC-b4 loop that is characterized by a stable \beta turn. This loop also contains many key residues with highly correlated chemical shift changes (black spheres).
may constitute a novel target to modulate the allosteric functions of each
kinase (to ATP binding). Here, we show that once the R-spine is assembled, the molecule is
of the switch mechanism that is an inherent feature of every protein ki-
nase. This small, highly dynamic segment encapsulates the essence
of this residue were also detected computationally
(Fig. 6B). This small conformation, particularly the b8-b9 sheet, is further stabilized by phos-
phorylation at Thr197 in the activation loop.

Fig. 6. Dynamic ordering of the hydrophobic core. (A) Most of the elements of the hydrophobic core (teal and gray) are stable in both active and inactive conformational
states. This includes b1 through b8 and the catalytic loop. These elements, as well as the C-spine, are anchored to the F helix by hydrophobic interactions. The assembly of
the R-spine completes the hydrophobic core and positions the Mg loop and b9 (red). (B) This active conformation, particularly the b8-b9 sheet, is further stabilized by phos-
phorylation at Thr197 in the activation loop.

engages one another following the binding of ATP. From earlier work, we
know that Asn99, when replaced with cysteine and modified by fluores-
ccin, is a sensor of ATP binding, and myristylation of the N terminus and
the dynamic feature of this residue were also detected computationally
with methyl-TROSY relaxation dispersion experiments, we now have
sensors for the entire b network and beyond. What is also striking about
this network is that most of it is firmly in place and always anchored to the
hydrophobic core even when the R-spine is not assembled. The one ex-
ception is b9 and the Mg loop that precedes it. This small segment shown
in red in Fig. 6A is only assembled as a consequence of the formation of
the R-spine. One can also appreciate how phosphorylation of the activation
loop contributes to stabilizing both the active site and the R-spine
(Fig. 6B). This small, highly dynamic segment encapsulates the essence
of the switch mechanism that is an inherent feature of every protein ki-
nase. Here, we show that once the R-spine is assembled, the molecule is
poised to be committed to catalysis in a highly correlated way in response
to ATP binding.

Because the hydrophobic core of PKA-C is highly conserved across
the kinome (5), it emerges as a critical element for kinase regulation and
may constitute a novel target to modulate the allosteric functions of each
kinase (33). The drug imatinib, for instance, breaks the R-spine and in-
activates both Src and Abl kinases (33–35), whereas other drugs, such as
PLX4720, deform the conformation of the R-spine in BRAF and acti-

Details on the expression and purification procedures for PKA-C have
been previously reported (35, 36). For the samples used in the NMR
relaxation experiments, the murine Ca subunit of cAMP-dependent
protein kinase (PKA-C) cDNA was cloned into a PRSetb vector.
The kinase was expressed in Escherichia coli BL21(DE3) cells in M9 me-
dium with 15NH4Cl as the only source of nitrogen and 100% 2H-glucose.
The M9 salts were solubilized into 80% 2H2O. For selective 13CH3 labeling,
the M9 medium was supplemented with 2-ketobutyric acid-4-
13C,3,3-d2 (70 mg/liter) and 2-keto-3-(methyl-13C,d2)-butyric acid-4-
13C,d1 (90 mg/liter) 1 hour before induction. For selective 13CH3 labeling,
the growth medium was supplemented with 2-ketobutyric acid-4-
13C,3,3-d2 (70 mg/liter) and 2-keto-3-(methyl-d3)-butyric acid-4-
13C,3-d1 (90 mg/liter) 1 hour before induction. Protein overexpression was
induced by 0.4 mM isopropyl b-D-thiogalactopyranoside (IPTG) and
carried out overnight at 24°C.

PKA-C purification was carried out by affinity chromatography with recombinant RIIa subunit (35, 36). The RIIa(R213K) was cloned into a PRSetb vector and expressed in 4 liters of LB media. Protein expression
was induced by the addition of 0.4 mM IPTG and carried out for 5 hours
at 24°C before harvesting the cells. Cell pellets of the catalytic subunit
and RIIa(R213K) were combined and lysed in 30 mM 3-(N-morpholi-
no)propanesulfonic acid (Mops), 200 mM ATP, 15 mM MgCl2, and
5 mM 2-mercaptoethanol (pH 8.0). After removing cell debris by cen-
trifugation, the supernatant was incubated with Ni2+ nitrilotriacetic acid
resin (Thermo Scientific, 1 ml of resin per liter of culture) at 4°C for
approximately 3 hours. The resin was then washed with 30 mM Mops,
25 mM KCl, 15 mM MgCl2, and 5 mM 2-mercaptoethanol (pH 8.0)
and eluted with 30 mM Mops, 25 mM KCl, 15 mM MgCl2, 5 mM 2-
mercaptoethanol, and 1 mM cAMP (pH 8.0). The fractions containing
PKA-C were collected and dialyzed overnight in 20 mM KH2PO4, 25 mM
KCl, and 5 mM 2-mercaptoethanol (pH 6.5). The three isoforms of
PKA-C (corresponding to the three different phosphorylation states
with identical catalytic parameters) were separated by HiTrap SP col-
umn (GE Healthcare Life Sciences) chromatography, using a linear

MATERIALS AND METHODS
Sample preparation
All reagents used were of analytical grade or better. Synthesis and pu-
rification of PLN1,19 and PKI5–24 have been previously described (7, 22).
gradient from buffer A [20 mM KH₂PO₄ (pH 6.5)] to 30% buffer B [20 mM KH₂PO₄ and 10.0 M KCl (pH 6.5)] with a flow rate of 2.0 ml/min. Isoform II with pThr¹⁹⁷, pSer²³⁸, and pSer¹⁰ was used for all NMR experiments (37).

**NMR experimental conditions**

All NMR samples were prepared in a buffer solution containing 20 mM KH₂PO₄, 90 mM KCl, 10 mM MgCl₂, 10 mM dithiothreitol (DTT), and 1 mM NaN₃ (pH 6.5) at 27°C. The concentration of the methyl-labeled PKA-C sample was 400 μM in a 250-μl volume. The sample was first titrated with adenylyl-imidodiphosphate (ATPγN) (0.1, 0.2, 0.6, 1, 2, 3, and 12 mM) and then with PKI₅₋₂₅ (100, 200, 300, and 400 μM). The titration experiments were monitored by ¹H-¹³C-heteronuclear multiple-quantum coherence experiments. Note that we used a nonhydrolyzable nucleotide analog, ATPγN (Roche LifeScience), for our relaxation measurements, because ATPγN undergoes slow hydrolysis in a period of several days. Residue-specific resonance assignments of PKA-C were previously described (35).

**Analysis of CSPs**

The chemical shift trajectories were monitored using the CONCISE (COordinated Chemical Shifts bEhavior) method (38) to measure the change in structural equilibrium associated with each PKA-C complex (apo, ATPγN, ATPγN/PLN₁₋₁₉, and ATPγN/PKI₅₋₂₅). Using principal components analysis (PCA), the method identifies a set of residues whose chemical shifts respond linearly to the open-to-closed conformational transition of the kinase. Each one of these residues provides a measure of the equilibrium position for all the PKA-C complexes, scoring them along the first principal component. The equilibrium position for a given PKA-C variant is given by the average of the PC scores over all linear residues. To identify the residues whose chemical shifts follow a linear pattern, a threshold of 1.0 to 3.0 for the ratio of the SDs of PC1 over PC2 was used and residues affected by CSPs below 0.05 parts per million were discarded (38). To identify the largest group of residues that responded to ligand binding in a correlated fashion, an adapted version of the chemical shift covariance analysis (CHESCA) (39) was applied to the PCA projection of the chemical shifts. First, the correlation matrix between all linear residues’ PC1 projections was constructed and used to build dendrograms through hierarchical clustering. After a threshold of 0.97 level of correlation coefficient was applied, a total of 58 residues formed the subset to trace the equilibrium position for each state.

**Nuclear spin relaxation experiments**

Side-chain dynamics in the fast time scale (picosecond to nanosecond) were determined by measuring ¹H spin relaxation rates of the ¹³CHD₂ isotopomers (40) using an 850-MHz spectrometer equipped with a Bruker AVANCE III console. For ¹³CHD₂ methyl-labeled PKA-C, we used a 250 μM sample for the apo form, a 230 μM sample for the binary form (saturated with 12 mM ATPγC), and a 200 μM sample for the closed form (saturated with 12 mM ATPγN and 400 μM PKI₅₋₂₅) in a 300-μl volume. All the samples were analyzed by dynamic light scattering to confirm sample homogeneity and determine the global rotational correlation time (τc). All spectra were collected in an interleaved manner with relaxation delays of 0.8, 1.2, 1.6, 2.0, 2.4, 3.2, 4.0, and 4.8 ms for the apo enzyme; 0.5, 1.0, 1.5, 2.0, 2.5, 3.0, 3.5, and 4.0 ms for the nucleotide-bound PKA-C; and 0.4, 0.8, 1.2, 1.6, 2.0, 2.4, 2.8, and 3.2 ms for the ternary complex. The intensities were fit to a single exponential using nonlinear fit, and the order parameter was extracted from the relaxation rate using the following relationship (40)

\[
R(D⁺) \approx R(D₋Dₓ + DₓD₋) \approx \left( \frac{1}{80} \right) (2πQCC)^2 S^2 τ_c
\]

where QCC = \(\varepsilon qQ/h\) is the quadrupolar coupling constant (167 ± 1 kHz) and \(S^2\) is the order parameter. The order parameters were calculated directly using transverse deuterium relaxation rates and the global rotational correlation time based on the dynamic light scattering data.

**Methyl relaxation dispersion experiments**

Single-quantum ¹³C CPMG (41) experiments were carried out on 700- and 850-MHz spectrometers equipped with Bruker AVANCE and AVANCE III consoles, respectively. All the spectra were collected in an interleaved manner with CPMG field strengths of 50, 100, 150, 200, 250, 300, 400, 500, 600, 800, and 1000 Hz. Replicate experiments were performed at field strengths of 50, 300, and 1000 Hz with a constant time delay of 40 ms. Each 2D spectrum was recorded with 2048 × 160 complex points and 48 transients with a recycle delay of 2.0 s. All data were processed using NMRPipe (42), and peak intensities were picked using Sparky (43). The peak intensities were converted to transverse decay rates, \(R_{2\text{eff}}\), as previously described (44). The software GUARDD (45) was used to fit the relaxation data to the Carver-Richards equation (46), which describes the dependence of the relaxation contribution of chemical exchange, \(R_{ex}\), to transverse relaxation. A two-state fit provides the exchange rate, the population of the second state, and chemical shift difference for a two-state exchange process

\[
R_2 \left( \frac{1}{τ_c} \right) = R_2^0 + \frac{1}{2} k_{ex} - \frac{1}{τ_c} \cos^{-1} [Dₓ \cosh(ηₓ) - D₋ \cosh(η₋)]
\]

\[
Dₓ = \frac{1}{2} \left( \pm 1 + \frac{\psi + 2Δθ^2}{(ψ^2 + Δθ^2)^{1/2}} \right)
\]

\[
ηₓ = \frac{τ_p}{\sqrt{2}} \left( (ψ^2 + Δθ^2)^{1/2} \right)^{1/2}
\]

\[
ψ = k_{ex}^2 - Δθ^2, \quad Δθ = -2∆θ(p_A - p_B)
\]

where \(p_A\) and \(p_B\) are the populations of the two states, \(Δθ\) is the chemical shift difference of the two states, \(τ_p\) is the time between the \(p\) pulses, \(k_{ex}\) is the sum of the rates of the exchange process, and \(R_2^0\) is the intrinsic transverse relaxation rate. The fitting was performed by minimizing the function \(χ^2\) as previously described (45). The group fit of selected residues was performed if the \(χ^2_{\text{Group}}/χ^2_{\text{Local}}\) was less than 2.0 (44).

An estimate of the \(R_{ex}\) can also be approximated by measuring the \(R_2\) for a high and low \(τ_p\) value. The approximate \(R_{ex}\) was found with the same CPMG setup but with two values (with \(τ_p\) values of 1 and 20 ms). The approximate \(R_{ex}\) for the ATPγN/PLN₁₋₁₉ bound form of PKA-C was estimated by

\[
R_{ex} \approx \frac{1}{T} \ln \frac{I(τ_p = 1 \text{ ms})}{I(τ_p = 20 \text{ ms})}
\]

where \(I\) is the intensity of the residue with the corresponding spin-echo period and \(T\) is the constant time period (40 ms).
Western blot analysis
QuikChange I and II site-directed mutagenesis kits (Agilent Technologies) were used to introduce mutations. Wild-type and mutants of PKA-C were tested for their autophosphorylation ability using the Western blot-based assay as previously described (6). Briefly, recombinant forms of murine PKA-C were transformed in BL21 (DE3) cells and grown in 5-ml batch cultures at 37°C until an optical density at 600 nm (OD_600) of 0.6 was reached. At this time, protein overexpression was induced with 0.2 mM IPTG and allowed to grow overnight. Induced cultures were lysed in a buffer containing 50 mM phosphate buffer (pH 7.0), 250 mM NaCl, and 2 mM DTT. Lysates were run on an SDS–polyacrylamide gel electrophoresis gel and tested for phosphorylation using site-specific antibodies. trans-Phosphorylation at the activation loop was probed by pThr\(^{373}\)-specific polyclonal antibody from Invitrogen. cis-Phosphorylation at the C-tail was probed by pSer\(^{338}\) antibodies (BD Biosciences), and expression levels of the recombinant PKA-C were probed by monoclonal antibodies from BD Biosciences.

ConSurf analysis
The sequence conservation was plotted onto the protein structure using ConSurf analysis (51). Initially, a CS-BLAST (49) search against the SWISS-PROT database was performed to obtain at least 50 unique hits. The list of collected homologs was subsequently filtered by coverage (minimum of 80%) and sequence identity (between 60% and 95%). The remaining homologs were filtered again using Cluster Database at High Identity with Tolerance algorithm with 90% sequence identity. The remaining homologs were filtered again using Cluster Database with a clustering threshold (at High Identity with Tolerance algorithm with 90% sequence identity). The remaining homologs were filtered again using Cluster Database (minimum of 80%) and sequence identity (between 60 and 95%). The list of collected homologs was subsequently filtered by coverage using the SWISS-PROT database was performed to obtain at least 50 unique hits. Ionotropic (2H,13C,15N,1H) 53% of 2H,13C-isoleucine, leucine, and valine (IVL) 19) forms of 2H,13C-isoleucine, leucine, and valine (IVL) were used to introduce mutations. Wild-type and mutants of PKA-C were probed by monoclonal antibodies from BD Biosciences.

Dynamic light scattering
The homogeneity of the PKA-C samples prepared for NMR relaxation experiments was analyzed with a Malvern Zetasizer as colors onto the Protein Data Bank (PDB) structure using PyMOL (Schrodinger Inc.). The homogeneity of the PKA-C samples prepared for NMR relaxation experiments was performed at 27°C where all NMR experiments were performed.

SUPPLEMENTARY MATERIALS
Supplementary material for this article is available at http://advances.sciencemag.org/cgi/content/full/3/4/e1600063/DC1
fig. S1. (H\(^{1+13}\)C) methyl-TROSY spectra of apo, nucleotide-bound (ATP\(_{C}\)), and ternary (ATP\(_{N}/\text{PKI}_{5-24}\) and ATP\(_{N}/\text{PLN}_{1-19}\)) forms of \(^{13}\)C-isoleucine, leucine, and valine (NL)–labeled PKA-C.
fig. S2. Plots of the IVL methyl group CSPs upon ligand binding.


A dynamic hydrophobic core orchestrates allostery in protein kinases

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