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Regulation of signaling directionality revealed by 3D snapshots of a kinase: regulator complex in action

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Abstract Two-component systems (TCS) are protein machineries that enable cells to respond to input signals. Histidine kinases (HK) are the sensory component, transferring information toward downstream response regulators (RR). HKs transfer phosphoryl groups to their specific RRs, but also dephosphorylate them, overall ensuring proper signaling. The mechanisms by which HKs discriminate between such disparate directions, are yet unknown. We now disclose crystal structures of the HK:RR complex DesK:DesR from *Bacillus subtilis*, comprising snapshots of the phosphotransfer and the dephosphorylation reactions. The HK dictates the reactional outcome through conformational rearrangements that include the reactive histidine. The phosphotransfer center is asymmetric, poised for dissociative nucleophilic substitution. The structural bases of HK phosphatase/phosphotransferase control are uncovered, and the unexpected discovery of a dissociative reactional center, sheds light on the evolution of TCS phosphotransfer reversibility. Our findings should be applicable to a broad range of signaling systems and instrumental in synthetic TCS rewiring.

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Introduction

Perception of environmental and intracellular cues is an essential feature of life. Signaling pathways enable cells to regulate genetic and biochemical programs for adaptation and survival. Two-component systems (TCSs) play a particularly important role among protein machineries that cells have evolved to carry out signaling, widely distributed in bacteria, fungi and plants. The simplest TCSs comprise a sensory histidine kinase (HK) and a response regulator (RR) component (*Gao and Stock, 2009*). The signal is transmitted downstream from HK to RR, through an orderly sequence of conformational rearrangements coupled to phosphoryl-transfer reactions. The HK is turned on or off after signal-dependent allosteric rearrangements, which control autophosphorylation on a conserved histidine residue. The phosphoryl group is then transferred from the phosphorylated kinase to a conserved aspartate on the RR receiver domain, activating it to effect a specific output response. When auto-kinase activity is turned off, HKs often act as a phosphatase of their cognate phosphorylated RR (P~RR) (*Goulian, 2010*), contributing to shutting down the pathway. Despite being paradoxical for a protein kinase, the phosphatase activity of HKs is physiologically relevant ensuring robust homeostatic responses (*Schramke et al., 2016*).

HKs are homodimeric proteins, comprising an N-terminal sensor domain and a more conserved catalytic core. Trans-membrane HKs display the catalytic region in the cytoplasm and the sensory domain within the lipid bilayer or toward the extracellular/periplasmic space. The catalytic core typically includes a helical Dimerization and Histidine phosphotransfer (DHP) domain, followed by a Catalytic and ATP-binding (CA) domain. HKs are classified according to DHP sequence signatures in HisKA, HisKA_2, HisKA_3, HWE_HK and His_kinase families. CheA-like HKs instead harbor the reactive His within a Histidine Phosphotransfer (HPt) domain that is not involved in dimerization. Overall sequence conservation among HKs can be very low, but the entire catalytic region is remarkably conserved at the structural level (Zschiedrich *et al.*, 2016), suggesting that functional mechanisms are shared. RRs comprise a receiver domain (REC), which may be adjacent to additional effector domains (DNA binding, enzymatic, etc.). REC phosphorylation stabilizes structural rearrangements associated with effector domain activation (Gao and Stock, 2010).

Phosphorelays are examples of more complex TCS pathways, involving additional intermediate phosphotransfer proteins, such as extra REC, HPt and/or modified DHP domains. Phosphorelays allow for more complex signaling circuits and sharper regulation (Goulian, 2010). HKs have evolved to catalyze irreversible phosphotransfer reactions (Porter *et al.*, 2008; Potter *et al.*, 2002) ensuring a unidirectional input→output information flow (Bourret and Stock, 2002). However, evolutionary pressure has also resulted in signaling pathways where reversible steps are simultaneously required, mostly in phosphorelays (Burbulys *et al.*, 1991; Janiak-Spens *et al.*, 2005), where dedicated HPt modules act as phospho-donors and -acceptors. The molecular bases underlying such differential reversibility patterns remain unknown, raising the question of how signaling pathway directionality is ensured, from input signal to adaptive response.

To address this question we have used the TCS DesK-DesR from *B. subtilis* as a model (de Mendoza, 2014). We had previously shown that the HK DesK, a member of the HisKA_3 family, undergoes important conformational changes to switch between phosphatase- and phosphotransfer-competent states (Albanesi *et al.*, 2009). The cytoplasmic portion of DesK (DesKC) including its entire catalytic region, displays a symmetric and rigid structure in the phosphatase state. DesK activation implies its rearrangement into an auto-kinase competent form with substantially higher flexibility. Upon auto-phosphorylation DesKC adopts a strongly asymmetric conformation (Albanesi *et al.*, 2009) able to transfer the phosphoryl group to DesR, its downstream RR partner. The DHP α -helices include a conserved membrane-proximal coiled-coil motif, upstream of the phosphorylatable His. Such DHP segment is critical for signal transmission, folding into a coiled-coil in the phosphatase state, whereas in the phosphotransferase form this coiled-coil breaks apart (Saita *et al.*, 2015). We now report the crystal structures of the DesKC:DesR complex, trapped in the phosphatase and the phosphotransferase functional states. Extensive conformational rearrangements reveal how the two reaction centers are remodeled. Our data indicate that the relative orientation and distance of the reactive histidine with respect to the receiver aspartate is a molecular determinant controlling signal directionality.

Results

Two crystal structures of the DesKC:DesR complex represent snapshots of the dephosphorylation and the phosphotransfer reactions

To grasp the molecular determinants of unidirectional TCS signaling, the crystal structures of DesKC in complex with DesR were determined in two functional states (Figure 1, Table 1). One using the phosphatase-constitutive mutant DesKC_{STAB}, and the other with DesKC_{H188E} carrying a phosphomimetic substitution (Albanesi *et al.*, 2009), both in complex with the REC domain of DesR. DesKC_{STAB} includes amino acid replacements Ser150Ile, Ser153Leu and Arg157Ile in the DHP domain (Figure 1—figure supplement 1A), designed to stabilize the coiled-coil region (Saita *et al.*, 2015).

The DesKC_{STAB}:DesR-REC complex

The asymmetric unit (ASU) of this crystal structure shows one full complex and half of a second one, which is completed through the crystallographic two-fold symmetry operator (Figure 1—figure supplement 1B). Several features indicate that this complex represents a snapshot of the phosphatase reaction, hereafter denominated the 'phosphatase complex'. Symmetric organizations are a hallmark

of HKs in the 'auto-kinase off / phosphatase on' states (Albanesi et al., 2009; Casino et al., 2009; Yamada et al., 2009). The phosphatase complex we have now crystallized indeed displays high symmetry, both in the way the REC domains associate to the HK dimer, as well as between HK monomers. The two independently refined phosphatase complexes show one dimer of DesKC_{STAB} bound to two molecules of DesR-REC (Figure 1A and Figure 1—figure supplement 1C), with the latter occupying equivalent positions on either side of the HK core. The 2:2 HK:RR stoichiometry was confirmed in solution by isothermal titration calorimetry (ITC) (Figure 1B, Table 2), revealing an entropy-driven, endothermic association reaction. Size exclusion chromatography-coupled small-angle X ray scattering (SEC-SAXS) further supported the 2:2 stoichiometry (Figure 1C; Tables 3 and 4). There is also a high internal symmetry between both HK monomers, with CA domains rigidly fixed onto the central DHP and a resulting butterfly-like shape of the whole molecule. A second feature consistent with a phosphatase-competent configuration, is that one AMP-PCP (ATP analogue) moiety is bound to each CA domain, far (>27 Å) from the His_{188(HK)} phosphorylation sites on the central DHP, precluding auto-kinase activity (subscripts HK and RR highlight the protein to which the indicated residues belong). Finally, the structural rearrangements triggered at high temperatures (Saita et al., 2015), driving DesK to its phosphatase state, have been linked to the stabilization of an N-terminal coiled-coil (Albanesi et al., 2009). The three point-mutations engineered within DesK helix α 1, are indeed observed to stabilize a coiled-coil structure toward the N-terminus in the phosphatase complex (Figure 1—figure supplement 1D). Coiled-coil stabilization was devised according to previous structures of free DesKC_{H188V} (Albanesi et al., 2009), a mutant version that maintains normal P~DesR-specific phosphatase activities in vitro and in vivo (Albanesi et al., 2004). Free DesKC_{H188V} structures superimpose extremely well onto the HK partner of the phosphatase complex (Figure 1—figure supplement 1E). SEC-SAXS data further suggest that the coiled-coil conformation in the phosphatase complex was the most abundant species in solution (Figure 1C and Figure 1—figure supplement 1B; Table 4), involving a large α 1: α 1 dimerization interface of \sim 2100 Å² (Figure 1A and Figure 1—figure supplement 1D and E). Regarding the RR component within the phosphatase complex, the three DesR-REC molecules in the ASU display identical 'active-like' structures (Trajtenberg et al., 2014). BeF₃⁻ moieties mimicking phosphoryl groups are observed bonded to each phosphorylatable Asp_{54(RR)}, and coordinated to Mg²⁺ cations, consistent with this crystal form being a snapshot of the phosphatase reaction at its pre-dephosphorylation step.

The DesKC_{H188E}:DesR-REC complex

Wild-type phosphorylated DesKC (wt P~DesKC) in complex with DesR variants resisted crystallization. The crystal structure was eventually solved by using the phosphomimetic DesKC_{H188E} mutant in complex with DesR-REC. Three X ray diffraction datasets were obtained from DesKC_{H188E}:DesR-REC crystals soaked with different concentrations of MgCl₂ and BeF₃⁻ (Table 1). The three structures are similar, showing two DesKC_{H188E}:DesR-REC complexes in the ASU (Figure 1D). A number of observations readily distinguish this complex from the phosphatase one, supporting it has captured a ground state of the P~HK→RR phosphoryl-transfer reaction, hereafter referred to as the 'phosphotransferase complex'. In the first place, the phosphotransferase complex is highly asymmetric, a typical feature of phosphorylated and kinase-active forms of HKs (Albanesi et al., 2009; Mechaly et al., 2014; Casino et al., 2014). One DesKC_{H188E} dimer was observed bound to one DesR-REC monomer in the crystal. This 2:1 stoichiometry was further supported by ITC (Figure 1E and Table 2) and SEC-SAXS (Figure 1F; Tables 3 and 4) in solution. The symmetry is also broken within the HK dimer, with one of the two CAs interacting with the DHP helix α 1 of the other protomer (Figure 1D). The second CA domain is free, visible in the structure due to fortuitous crystal packing contacts. Such arrangement leaves only one RR-binding site available on the HK in the phosphotransferase state, explaining the asymmetric stoichiometry. Secondly, structural superposition of the HK partner extracted from the phosphotransferase complex, with the available structure of free phosphorylated DesK (wt P~DesKC [PDB 3GIG]), reveals closely similar conformations (Figure 1—figure supplement 1F), confirming that the structure of DesKC_{H188E} in complex is a reliable model of the wt phosphorylated species. Finally, BeF₃⁻ was not observed bonded to Asp_{54(RR)} in any of the DesKC_{H188E}:DesR-REC complex structures, despite testing elevated soaking concentrations. Yet, the phosphorylation sites of the RR partners are very similar compared to the phosphatase complex, allowing for the Mg²⁺ cation to bind alike (the cation was observed bound only in phosphotransferase complex structures

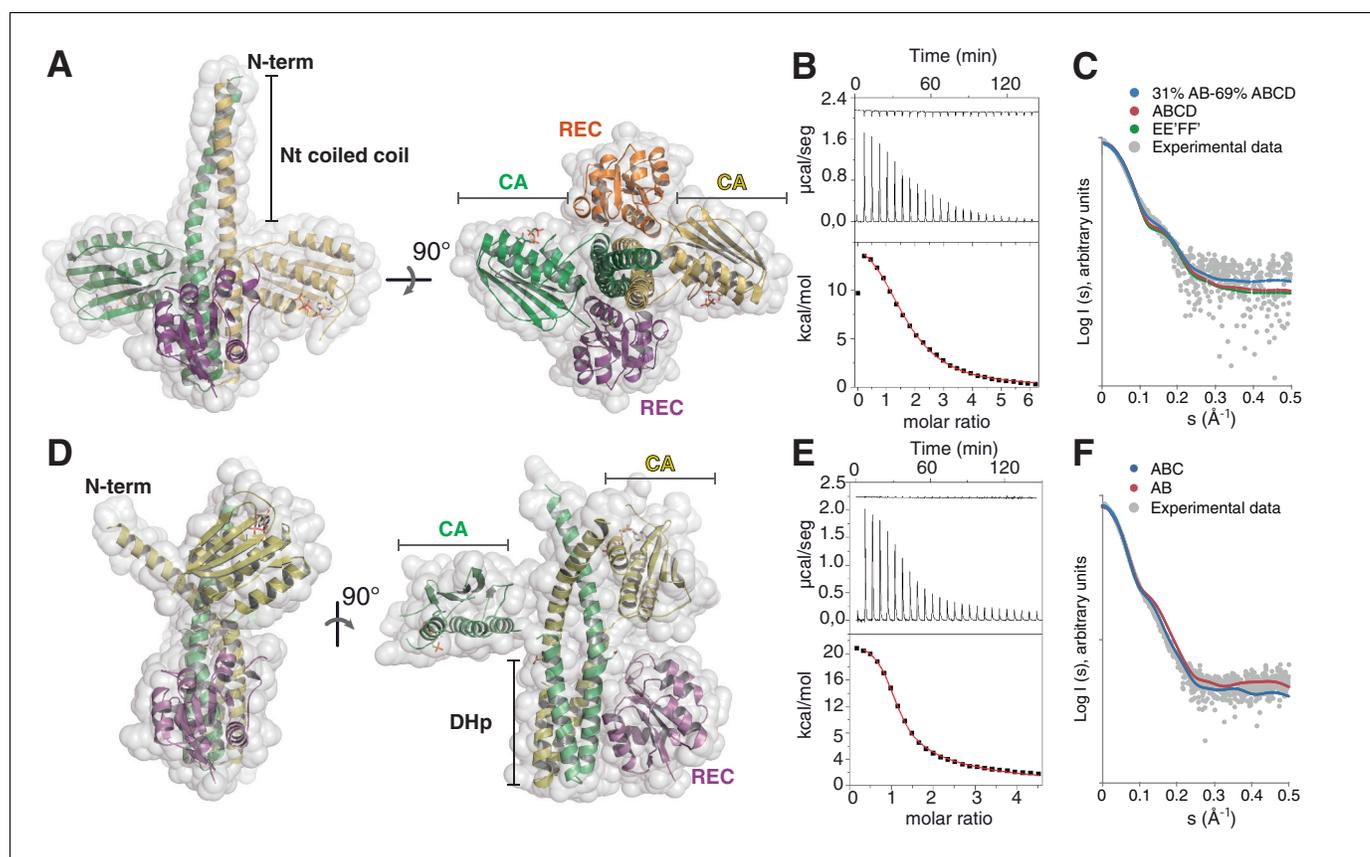


Figure 1. The phosphatase and the phosphotransferase complexes: crystal structures and stoichiometries in solution. (A) Cartoon representation of the DesKC_{STAB}:DesR-REC (phosphatase) complex, along two orthogonal views (left and right subpanels). The two chains within the DesKC_{STAB} dimer are depicted in green and yellow, and the two DesR-REC molecules, in magenta and orange. Solvent exposed surfaces are shown in transparent gray. (B) Isothermal titration calorimetry (ITC) of the phosphatase DesKC_{H188V}:DesR-REC binding reaction, top panel showing the raw heat flow data; in the bottom, integrated heat exchange as a function of RR(monomer):HK(dimer) molar ratio. ITC was performed with the mutant DesKC_{H188V}, equivalently trapped in the phosphatase state (Albanesi et al., 2009), because DesKC_{STAB} was insoluble when not co-expressed with DesR-REC. (C) X ray scattering curve for the DesKC_{STAB}:DesR-REC complex. Experimental data are plotted as gray dots, with theoretical curves overlaid as colored lines, revealing best fitting for the AB+ABCD mixture (see Table 4). Letters distinguish protomer chains: DesKC_{STAB} dimers AB and EE' (primed labels distinguish crystallographically related partners) and DesR-REC monomers C, D, F and F'. (D) Cartoon representation of the DesKC_{H188E}:DesR-REC (phosphotransferase) complex, along two orthogonal views (left and right subpanels). Coloring scheme and solvent exposed surface displayed as in (A). (E) ITC of the DesKC_{H188E}:DesR-REC binding reaction, details as in (B). (F) SAXS curve for the DesKC_{H188E}:DesR-REC complex, details as in (C). Note best fitting with a one dimer DesKC_{H188E} (chains AB) to one monomer DesR-REC (chain C) model (see Table 4).

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The following figure supplement is available for figure 1:

Figure supplement 1. Phosphatase and phosphotransferase complexes: structural details.

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soaked with high Mg²⁺ concentrations). The absence of BeF₃⁻ in the phosphotransferase state, even though Mg²⁺ was eventually bound, is consistent with phosphorylated HKs not interacting with phosphorylated RR species. The molecular basis for a decreased BeF₃⁻ affinity in the phosphotransferase became clear later by comparing both states' reaction centers (see below). Taken together, the evidence indicates the phosphotransferase complex represents a snapshot of the transfer reaction, prior to the P~His→Asp phosphoryl group migration.

DesK and DesR interact through a 'slippery' interface

Both phosphatase and phosphotransferase complexes show that the REC domain of DesR interacts with DesK through its α1_(RR) helix and the β5α5_(RR) loop, with a few additional contacts from the N-terminal portion of helix α5_(RR) and the beginning of loop β4α4_(RR). On the HK side, the main

Table 1. X ray diffraction data collection and refinement statistics.

	Phosphatase complex	Phosphotransferase complex I (low [Mg ²⁺])	Phosphotransferase complex II (high [Mg ²⁺])	Phosphotransferase complex III (high [Mg ²⁺] + BeF ₃ ⁻)	wt P-DesKC
Data collection					
Space group	P3 ₁ 21	P2 ₁	P2 ₁	P2 ₁	P3 ₁ 21
Cell dimensions					
a, b, c (Å)	94.3, 94.3, 239.9	87.8, 114.6, 91.6	87.9, 115.6, 91.6	88.2, 116.7, 91.9	94.4, 94.4, 161.8
α, β, γ (°)	90, 90, 120	90, 116.4, 90	90, 116.7, 90	90, 117.1, 90	90, 90, 120
Resolution (Å)	48.35–2.79 (2.94–2.79)*	66.70–3.2 (3.37–3.2)	66.78–2.9 (3.06–2.9)	38.59–3.15 (3.32–3.15)	36.5–3.16 (3.33–3.16)
Unique reflections	31510 (4385)	26369 (3906)	36132 (5279)	28033 (3944)	14829 (2109)
R _{meas}	0.1 (2.07)	0.14 (0.98)	0.13 (1.61)	0.1 (0.57)	0.07 (1.18)
R _{pim}	0.03 (0.63)	0.09 (0.62)	0.05 (0.65)	0.05 (0.29)	0.03 (0.43)
CC _{1/2}	1.00 (0.56)	0.99 (0.55)	0.99 (0.67)	1.00 (0.83)	1.00 (0.77)
I / σI	15.6 (1.5)	7.7 (1.7)	12.5 (2.4)	11.8 (2.7)	23.8 (1.9)
Completeness (%)	99.5 (96.7)	97.8 (98.8)	99.1 (99.2)	98 (95.6)	99.9 (99.6)
Redundancy	10.8 (10.3)	2.4 (2.4)	5.8 (6.0)	3.8 (3.8)	7.2 (7.2)
Refinement					
Resolution (Å)	48.35–2.79	66.70–3.2	39.27–2.9	38.59–3.15	36.5–3.16
Number of refls used (N in the free set)	31454 (1568)	26231 (1320)	35374 (1720)	28015 (1361)	14787 (791)
R _{work} / R _{free}	0.214/0.249	0.187/0.24	0.195/0.239	0.19/0.231	0.257/0.295
Number of atoms					
Protein	7972	8869	8702	8709	3151
Ligands + ions	93 (AMP-PCP)/6 (Mg ²⁺)/12 (MES)/12 (BeF ₃ ⁻)/5 (SO ₄ ²⁻)	124 (AMP-PCP)/4 (Mg ²⁺)/2 (K ⁺)	124 (AMP-PCP)/5 (Mg ²⁺)/2 (K ⁺)	124 (AMP-PCP)/6 (Mg ²⁺)/2 (K ⁺)	62 (AMP-PCP)/2 (Mg ²⁺)/6 (glycerol)
Water	12	15	9	18	4
B-factors (Å ²)					
Wilson plot	98.3	91.4	95.9	84.1	128.9
Mean (overall)	118.7 ‡	93.1 ‡	102.5 ‡	89.6 ‡	123.8 ‡
R.m.s. deviations					
Bond lengths (Å)	0.01	0.01	0.01	0.01	0.01
Bond angles (°)	1.2	1.3	1.25	1.25	1.3
Number of residues in Ramachandran plot § (favored / outliers)	1025/1	1099/3	1074/2	1073/5	382/4
PDB ID	5IUN	5IUJ	5IUK	5IUL	5IUM

*Values in parentheses correspond to the highest-resolution shell.

‡ Including TLS contribution.

§ Calculated with Molprobity ([Chen et al., 2010](#))

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element engaged in the interface is helix α1_(HK) on the Dhp domain from one DesK protomer, and minor contacts with helix α2'_(HK) from the second protomer. This interface broadly resembles the one found in other TCS complexes ([Casino et al., 2009](#); [Willett et al., 2015](#); [Yamada et al., 2009](#); [Zapf et al., 2000](#)) ([Figure 2A](#)).

The Dhp:REC interface engages van der Waals contacts almost exclusively, including several hydrophobic residues ([Figure 2B](#)), which clustered among the few HK:RR covariant residue pairs ([Figure 2—figure supplement 1A](#)) found to be conserved in other HK and RR families

Table 2. Isothermal titration calorimetry parameters. For details on the two different titration procedures, see Materials and methods.

HK:RR complexes	K_a (M^{-1})	ΔG ($\times 10^3$ kcal.mol $^{-1}$)	ΔH ($\times 10^4$ kcal.mol $^{-1}$)	$T\Delta S$ ($\times 10^4$ kcal.mol $^{-1}$)
DesKC _{H188V} :DesR-REC	7.7×10^5	-7.8	1.5	2.3
	9.5×10^4	-6.5	1.4	2.1
DesKC _{H188E} :DesR-REC	1.7×10^6	-8.2	2.2	3.0
	3.0×10^4	-5.9	2.1	2.7

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(Skerker et al., 2008). Leu_{200(HK)} plays a central role by inserting its side chain into a hydrophobic pocket within DesR, delimited by the side chains of Leu_{13(RR)}, Ala_{16(RR)}, Leu_{17(RR)} and Leu_{20(RR)} on one side, and the main chain atoms of the loop spanning residues 102–106_(RR), on the other (Figure 2—figure supplement 1B). Ser_{196(HK)} and Gln_{193(HK)}, located on the same face of $\alpha 1_{(HK)}$ as Leu_{200(HK)}, establish additional van der Waals contacts within a groove on the RR's surface, extending the Leu_{200(HK)}-lodging pocket all the way to the reaction center, which is rather open (Figure 2—figure supplement 1B). Very few polar contacts are observed surrounding the hydrophobic core, on both ends of the interface patch: Asp_{203(HK)}/Arg_{206(HK)} with Ser_{106(RR)}/Glu_{107(RR)}, and a salt bridge between Arg_{235(HK)} and Asp_{103(RR)} on the other end. The buried surface is $\sim 1400 \text{ \AA}^2$ in the phosphatase state, the DHP:REC interface being the major contributor ($\sim 1000 \text{ \AA}^2$) (Figure 2C and Figure 2—figure supplement 1C), although the CA domain adds a buried area of $>400 \text{ \AA}^2$. On the other hand, the phosphotransferase complex shows a slightly smaller interface ($\sim 1000 \text{ \AA}^2$), dominated by the DHP:REC interaction ($\sim 900 \text{ \AA}^2$) (Figure 2D and Figure 2—figure supplement 1C).

Superposing the C α atoms of the DHP region involved in DesR-binding (DesK residues 190–235) among the nine independently refined DesKC:DesR interfaces, reveals large variability in the relative positioning of the REC domains, with rmsd as high as 2.4 \AA for the RECs' 129 C α atoms (Figure 2E, Table 5). These significant movements engage corresponding shifts in the positions of interacting residues, albeit preserving the contacts, reminiscent of a 'slippery' interface. The slippery nature of the interface is consistent with its planarity, a low number of mainly hydrophobic contacts, leading to low shape complementarity and unfilled cavities within the RR partner (Figure 2F).

A single response regulator conformation is selected by the two functional states of the kinase

Comparing the phosphatase vs phosphotransferase complexes, the HK partner reveals substantial rearrangements. Concerning the DHP domain, a large rotational shift is observed on the membrane-proximal side of the phosphorylation site (Figure 3—figure supplement 1A), in contrast to the distal RR-binding site, which remains essentially invariant. The relative orientation of the CA domains is also dramatically changed (Figure 3—figure supplement 1B). On the other hand, the RR REC domain adopts a unique conformation in both complexes, with an average rmsd of 0.3 \AA aligning all C α atoms, and no significant contrast clustering phosphatase vs phosphotransferase structures (Table 5). Interestingly, this conformation adopted by the REC domain in complex with the kinase, combines structural features of both active and inactive states as seen in the free forms of RRs (Gao and Stock, 2009; Trajtenberg et al., 2014) (Figure 3). Namely, the $\beta 4\alpha 4$ loop and $\beta 5$ strand

Table 3. Small angle X ray scattering and derived molecular size parameters. Data derived from experiments performed by SEC-coupled SAXS. Figures for the phosphatase-trapped species DesK_{H188V}, are to be compared with those corresponding to HK:RR species. Full details in Materials and methods.

Protein species	$I_{(0)}$	$I_{(0)}$ real	Rg (Guinier)	Rg (real)	Vc	MM _{Vc} (Da)	MM _{seq} (Da)	Dmax (\AA)
DesKC _{H188V}	4.47×10^{-2}	4.26×10^{-2}	31.25	31.21	442.9	46400	44780	100
DesKC _{STAB} :DesR-REC	1.16×10^{-1}	1.11×10^{-1}	31.48	31.42	494.8	58400	77110	103
DesKC _{H188E} :DesR-REC	1.05×10^{-1}	9.76×10^{-2}	32.84	32.18	486.2	51800	61790	103

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Table 4. Fitting figures (χ^2) comparing alternative theoretical SAXS curves to the experimentally collected scattering data.

Experimental Data	Structural models	χ^2
DesKC _{STAB} :DesR-REC	AB+ABCD mixture	6.1
	ABCD (folded coiled-coil)	8.9
	EE'FF' (disrupted coiled-coil)	11.6
DesKC _{H188E} :DesR-REC	ABC	8
	AB	24.9

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emulate the inactive RR, while the rest of the protein resembles the free phosphorylated species. The $\beta 4\alpha 4$ loop is closed in phosphorylated RRs (e.g. PDB 4LE0), coupled to a fully wound first turn of helix $\alpha 4$ (Pro_{85(RR)} displays a dihedral ψ angle near -40° , allowing for α -helicity). The REC molecules in both phosphatase and phosphotransferase complexes, show instead this $\beta 4\alpha 4$ loop open (with Pro_{85(RR)} ψ angles of respectively 126° and 130° , and the first turn of $\alpha 4$ unwound, as observed in the inactive state of the Mg²⁺-free protein [PDB 4LE1]). Thus acting as a 'phosphate lid', the $\beta 4\alpha 4$ loop opens and closes to control phosphoryl-transfer chemistry on Asp_{54(RR)} (**Figure 3**).

Regarding the positions of the phosphorylatable Asp, as well those of the $\beta 1\alpha 1$ loop and contiguous helix $\alpha 1$, HK-binding triggers RR 'activation', as we had previously predicted based on functional readouts and *in silico* modeling (**Trajtenberg et al., 2014**). The shift of loop $\beta 1\alpha 1$ locates key acidic residues in place to coordinate the essential Mg²⁺ cation within the RR's reaction center.

Taken together, the RR adopts a single intermediate pre-activated conformation, instrumental for both dephosphorylation and phosphotransfer reactions, due to increased exposure of the phosphorylation site and stabilized configuration of the pocket poised for reaction.

The phosphatase to phosphotransferase transition: reconfiguring the reaction center for catalysis

The crystal structure of wt P~DesKC was determined (**Table 1**), improving previous resolution limits (**Albanesi et al., 2009**). The reactive His_{188(HK)} was observed to be phosphorylated on its N ϵ 2 atom (**Figure 4—figure supplement 1A**), and superposition of this model onto the HK partner of the phosphotransferase complex (PDB 5IUK) revealed very similar structures, with 1.2 Å rmsd aligning all 1994 atoms that comprise the DHP domain and the fixed CA (**Figure 4—figure supplement 1B**). Substituting the DesKC_{H188E} component in the crystal complex by the superposed wt P~DesKC, revealed surprisingly good geometry between P~DesKC and DesR-REC (**Figure 4—figure supplement 1C**). Energy minimization easily fixed minor interatomic bumping, rendering a model with P~His_{188(HK)} well oriented toward Asp_{54(RR)}, poised for phosphotransfer (**Figure 4A**). The P~His_{188(HK)}N ϵ 2, phosphorus and Asp_{54(RR)}O δ 1 atoms describe a $\sim 180^\circ$ angle, with the phosphoryl group at H-bonding distance to the side chain oxygens of Thr_{80(RR)} and Thr_{81(RR)}. The model proved to be stable in molecular dynamics simulations (**Figure 4B**), overall making chemical and biological sense. We shall thus use the minimized wt P~DesKC:DesR-REC model to perform further structural analyses of the phosphotransferase complex reaction center.

The phosphatase→phosphotransferase switch engages major rearrangements. An $\sim 80^\circ$ rotation of helix $\alpha 1$ is coupled to a cogwheel-like shift of helix $\alpha 2$, at the level of the phosphorylation site within the DHP domain (**Albanesi et al., 2009**). This rotation includes the phosphorylatable His_{188(HK)} (**Figure 4C**), which had not been previously observed. Due to this shift, the side chain of His_{188(HK)} ends up buried in the phosphatase state, establishing a H-bond with the same residue of the other protomer (**Figure 4C** and **Figure 4—figure supplement 1D**), displaced away from the reaction center as compared to the phosphotransferase (**Figure 4D**). Asp_{189(HK)} right next to the phosphorylation site, follows a similarly dramatic rearrangement, well positioned only in the phosphatase complex to

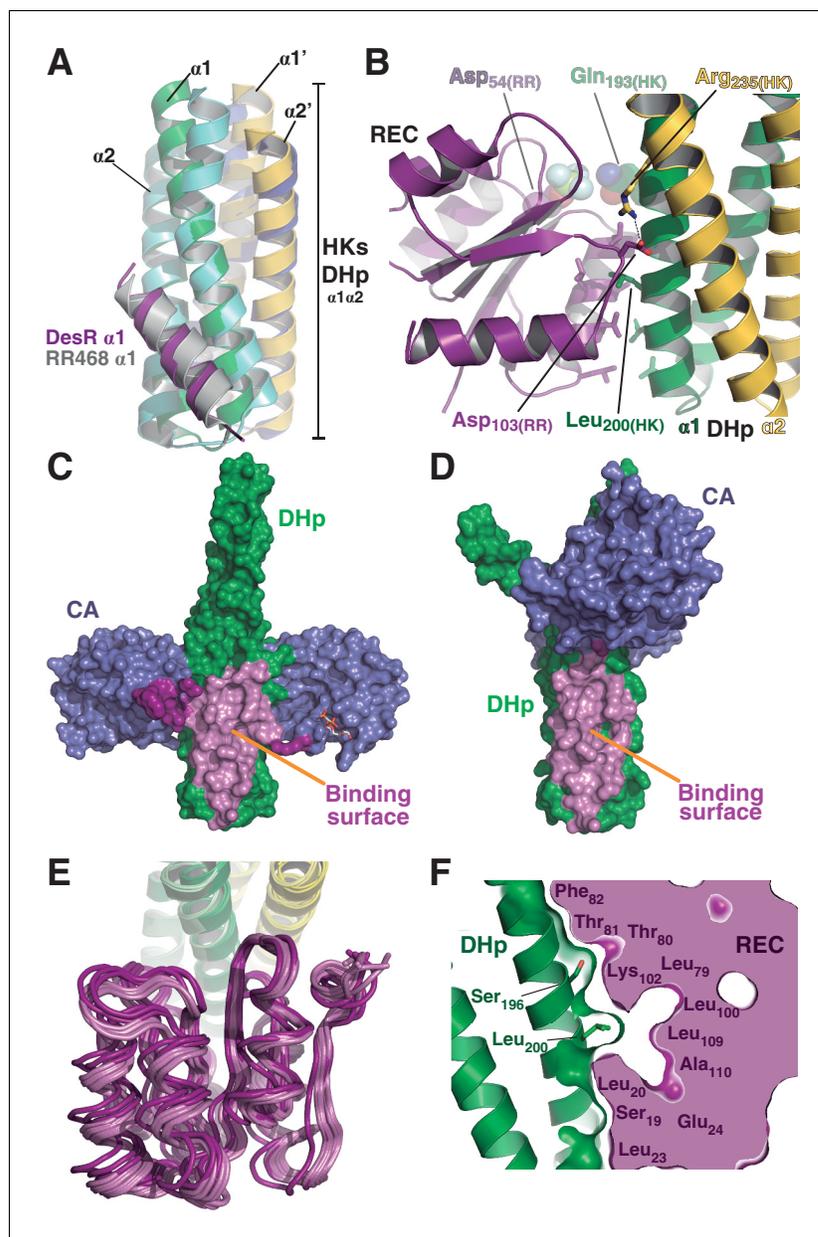


Figure 2. The HK:RR interface and structural variability. (A) Structural superposition of the DesK:DesR phosphatase complex (this study, PDB 5IUN) with HK853:RR468 from *Thermotoga maritima* (PDB 3DGE). Only selected structural elements from both partners are shown as cartoons for clarity. The DesK:DesR phosphotransferase complex is overall similar (not shown here). DesK and DesR are illustrated with the same color code as in **Figure 1**. Superposed *T. maritima* RR468 helix $\alpha 1$ is shown in gray, and HK853 protomers in cyan and blue. Primed labels distinguish DHp helices from each HK protomer. (B) Close-up of the phosphatase complex DesK:DesR interface showing only a few selected interactions for clarity (see text for detailed description). Coloring scheme as in (A). Note Leu_{200(HK)} inserted into a hydrophobic pocket of DesR, and several other hydrophobic residues completing the interface (not labeled, shown as sticks). Among the polar contacts surrounding the hydrophobic core, Arg_{235(HK)}:Asp_{103(RR)} and Gln_{193(HK)}:BeF₃⁻-modified Asp_{54(RR)}, are highlighted. (C) Solvent accessible surface of DesKC_{STAB} indicating the interaction footprint (in magenta) with DesR in the phosphatase complex. DesK domains are highlighted in green (DHp) and blue (CAs); these domains participate differently in DesR interaction, depicted in light magenta (DHp) and dark magenta (CA). Note the ATP analogue (AMP-PCP, in sticks colored by atom) bound to the CA domains, visible in this view on the rightmost one. (D) Same as (C), but for the phosphotransferase complex, illustrating the DesR-binding surface of DesKC_{H188E}. (E) The variable position of the REC_(RR) domain with respect to the DHp_(HK) is illustrated, superimposing the structurally invariant region of the DHp (including the nine

Figure 2 continued on next page

Figure 2 continued

independently refined DesK:DesR complexes). The coloring scheme is the same as in (A), with light and dark colors distinguishing phosphotransferase and phosphatase complexes, respectively. (F) The solid volume corresponds to DesR-REC, which is shown sliced to highlight its outline, revealing unfilled cavities at the protein: protein interface. The relative position of interfacing amino acids is roughly indicated with residue labels. DesK DHP helices are shown in cartoon representation, with its superposed molecular surface on top. Two key residues on the HK partner are shown as sticks.

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The following figure supplements are available for figure 2:

Figure supplement 1. Structural details of the DesK:DesR interface.

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Figure supplement 2. Molecular dynamics (MD) simulation of the phosphotransferase complex.

DOI: [10.7554/eLife.21422.010](https://doi.org/10.7554/eLife.21422.010)

make a salt bridge with Arg_{84(RR)}. Neighboring Phe_{82(RR)} thus inserts its aromatic side chain into a hydrophobic pocket formed between DHP α 1 of one HK protomer and α 2' of the other (**Figure 4D**). That the reactive histidine moves away from the catalytic site in the phosphatase state is consistent with its marginal role in P~RR dephosphorylation catalysis in HisKA_3 HKs. Indeed, the phosphorylatable His was proved nonessential for NarX-mediated P~NarL dephosphorylation (**Huynh et al., 2010**). His has been implicated in EnvZ-mediated phosphatase catalysis (**Zhu et al., 2000**) in which case its imidazole would act as a base assisting the attack of water on the P~aspartyl group (**Bhate et al., 2015**). But contradictory results cast doubts on such a direct role (**Hsing and Silhavy, 1997; Skarphol et al., 1997**). Instead, a Gln in HisKA_3 (**Huynh et al., 2010**), corresponding to Gln_{193(HK)} in DesK, or a Thr/Asn in HisKA HKs (**Willett and Kirby, 2012**), equivalently positioned one helical turn C-terminal to the phosphorylatable His, play a key role in the phosphatase reaction, positioning a catalytic water molecule. Such water or hydroxide molecule is well located to perform the nucleophilic attack on the phosphoryl group, as observed in the P~CheY3 phosphatase CheX (**Pazy et al., 2010**), a shared geometry among phosphatases and HKs as we now observe in the DesKC:DesR-REC phosphatase complex (**Figure 4—figure supplement 2**).

Table 5. Square matrix of all possible pair-wise structural superpositions. Superpositions were calculated using the nine independently refined DesKC:DesR-REC pairs, taken from the 1.5 complexes in the phosphatase asymmetric unit (named STAB1, STAB2 and STAB3) and the 2 complexes present in the ASU of each of the three different phosphotransferase structures (named as E₁₈₈[1–6]). Below the main diagonal the structurally invariant region of DesK DHP domains (residues 190–234_(HK)) were superposed, whereas above the diagonal, the DesR REC domains (residues 1–129_(RR)) were superposed instead. Resulting root mean squared deviations (in Å) were calculated, for both matrix halves, between all C α atoms of the REC domains (residues 1–129_(RR)) after superposition, as indicated in each individual matrix cell. Font colors highlight the two different alignment procedures performed. The significant differences between corresponding blue- and red-colored values, indicate that the REC domains do not change within, but rather 'slip' with respect to the kinase DHP domain.

	STAB1	STAB2	STAB3	E ₁₈₈ 1	E ₁₈₈ 2	E ₁₈₈ 3	E ₁₈₈ 4	E ₁₈₈ 5	E ₁₈₈ 6
STAB1		0.229	0.158	0.414	0.349	0.442	0.361	0.344	0.343
STAB2	0.826		0.188	0.496	0.459	0.512	0.334	0.343	0.333
STAB3	1.362	1.917		0.393	0.343	0.414	0.292	0.287	0.277
E ₁₈₈ 1	1.168	1.133	1.682		0.275	0.146	0.305	0.301	0.301
E ₁₈₈ 2	0.909	1.140	1.224	0.584		0.301	0.351	0.309	0.341
E ₁₈₈ 3	1.085	1.110	1.530	0.233	0.459		0.346	0.315	0.313
E ₁₈₈ 4	1.548	1.140	2.453	0.893	1.316	1.040		0.175	0.117
E ₁₈₈ 5	1.295	0.978	2.096	0.593	0.969	0.716	0.449		0.140
E ₁₈₈ 6	1.424	1.058	2.300	0.770	1.182	0.905	0.193	0.306	

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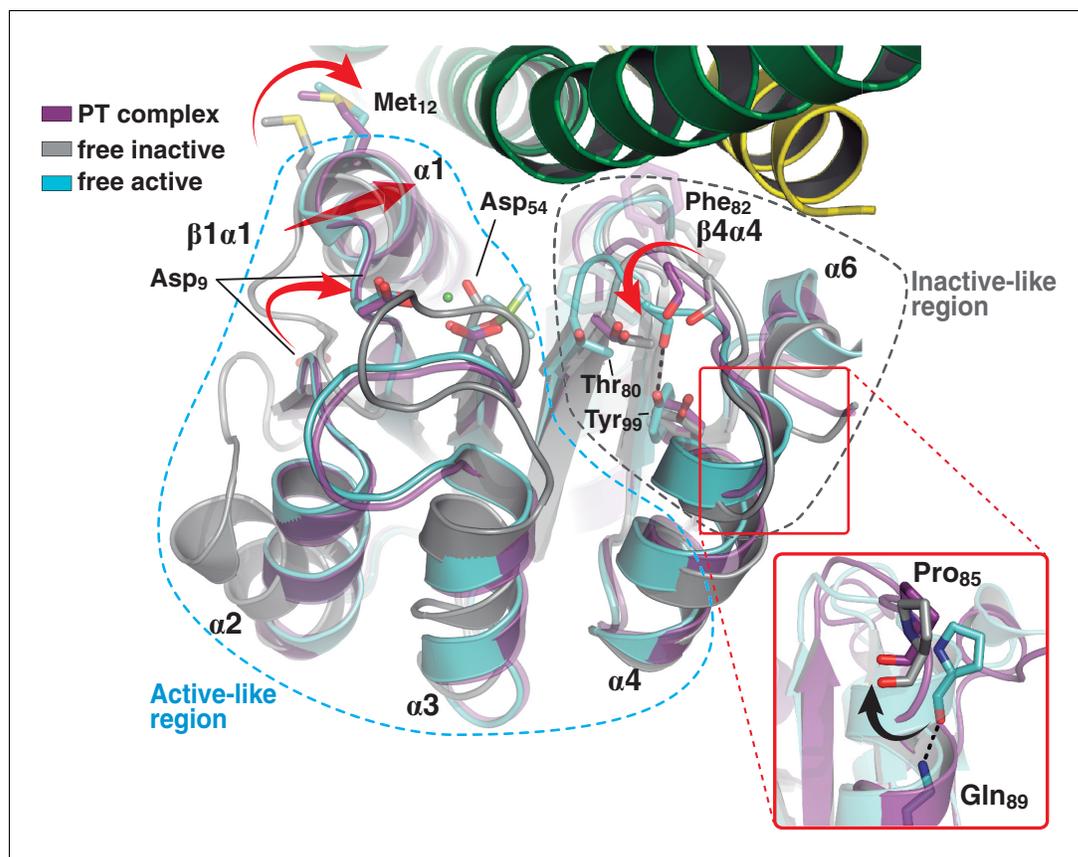


Figure 3. A shared RR REC domain conformation in the phosphatase and phosphotransferase complexes, has active- and inactive-like features. The phosphotransferase complex (this study, PDB 5IUK) is shown in cartoon representation, following the same coloring scheme as in **Figure 2**. REC domains from reported free DesR species are shown superposed, in gray the inactive Mg^{2+} -free RR (PDB 4LE1) and the active species in cyan (PDB 4LE0). Note that the complexed form of the regulator (in magenta) is mainly similar to the active species, including in the $\beta 1\alpha 1$ loop and $\alpha 1$ helix (that play key roles in the structuring of the RR's active site), even though the RR is not phosphorylated on the reactive $Asp_{54(RR)}$. Dashed outlines highlight the regions that most resemble each functional state. Note that the inactive-like region essentially involves the end of $\beta 4$ and the $\beta 4\alpha 4$ loop, which closes in as a phosphate lid in P-DesR. $Phe_{82(RR)}$ (shown in transparent sticks for clarity) interacts with the HK, playing a role in phosphate lid opening in both phosphatase and phosphotransferase complexes. $Tyr_{99(RR)}$ on strand $\beta 5$ (the strand is not visible below $\alpha 4$), typically forming a H-bond with the main chain O of $Phe_{82(RR)}$ in activated RRs, is observed farther away in both DesK:DesR complexes, yet another feature of inactive states. Inset: a close-up of the N-terminal tip of helix $\alpha 4$ is shown. The dihedral ψ angle of $Pro_{85(RR)}$ is linked to the active/inactive shift of the $\beta 4\alpha 4$ loop.

DOI: [10.7554/eLife.21422.012](https://doi.org/10.7554/eLife.21422.012)

The following figure supplement is available for figure 3:

Figure supplement 1. Major structural differences of the HK partner, comparing the phosphatase and phosphotransferase complexes.

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The rotational phosphatase→phosphotransferase rearrangement disrupts the $Asp_{189(HK)}:Arg_{84(RR)}$ ionic bond, and coupled to the $\alpha 1_{(HK)}$ and $\alpha 2'_{(HK)}$ movements, the position of $Phe_{82(RR)}$ shifts, propagating along the rest of the phosphate lid $\beta 4\alpha 4$ loop (**Figure 4D**). This shift accounts for the capacity of $Thr_{80(RR)}$ to be at H-bonding distance from the P-His $_{188(HK)}$ phosphoryl moiety, accompanying its transfer, and maintaining the H-bond to the phosphate on P- $Asp_{54(RR)}$ (**Figure 4D**). The side chain of $Phe_{82(RR)}$ is also contacting P-His $_{188(HK)}$, which could increase the pKa of its imidazole N $\delta 1$ atom due to cation- π interactions (Loewenthal et al., 1992). Indeed, P-DesKC appeared to catalyze slower phosphoryl-transfer to DesR-REC $_{F82A}$ compared to wt DesR-REC (**Figure 4E**), consistent with a

poorer N δ 1 protonation, and hence a stronger P-N phosphoramidate bond (Attwood *et al.*, 2007). Systematic mutagenesis at RR position 82 and functional assays (catalytic activities and HK:RR affinities) will be needed to reach a definitive conclusion.

Thus, the phosphatase and phosphotransferase catalytic activities depend upon modifications in the position of a subset of residues in both partners. Both complexes share a common RR intermediate conformation, and the reaction outcome is dictated by the HK switch of helices α 1 and α 2.

Dissociative phosphotransfer is linked to unidirectional signaling

DesK:DesR-catalyzed phosphotransfer occurred essentially in the forward, P~DesK \rightarrow DesR direction (Figure 5A), extending earlier suggestions (Albanesi *et al.*, 2004) into quantitative figures, and indicating that back-transfer is minimal. How is immediate P~RR dephosphorylation prevented once the phosphate has been transferred? The side chain of Thr_{80(RR)} at the end of β 4 strand, appears to drive the whole phosphate lid to a closed configuration, escorting the phosphoryl group during P~His \rightarrow Asp migration (Figures 3 and 4D). To test whether a closed phosphate lid indeed plays a role in minimizing P~RR dephosphorylation, as a means to ensuring phosphotransfer unidirectionality, Phe_{82(RR)} was substituted by alanine. Phe_{82(RR)} covers the P~Asp_{54(RR)} in active free DesR, while exposing it to bulk solvent through phosphate lid-opening when bound to DesK (Figure 4—figure supplement 2). The half-life of P~DesR_{F82A} was significantly reduced when compared to wt P~DesR (Figure 5B), confirming that the phosphate lid is a key structural element to avoid otherwise futile P~RR dephosphorylation after phosphotransfer.

Then how is P~RR \rightarrow HK back-transfer inhibited? The phosphate lid could also play a role. Yet, comparing wt P~DesR with point-mutants P~DesR_{F82A} or P~DesR_{R84A}, no change was detected in their marginal abilities to transfer the phosphoryl group back to DesK (data not shown). There is however a parameter that was found to discriminate irreversible phosphotransfer systems from others that reveal appreciable reversibility: the reactional His-Asp distance. The distance between P~His_{188(HK)}N ϵ 2 and Asp_{54(RR)}O δ 1 in the phosphotransferase complex is \sim 7.6 Å (Figures 4A and 5C), strongly suggesting that the phosphotransfer reaction occurs through a loose (or predominantly dissociative) nucleophilic substitution mechanism. This is in contrast to what has been observed in reversible complexes (Figure 5C), which efficiently catalyze phosphotransfer also in the opposite, P~Asp \rightarrow His direction. The latter display significantly shorter N ϵ 2-to-O δ 1 distances, consistent with an associative nucleophilic mechanism (Table 6). Examples include several phosphorelay protein pairs (Bauer *et al.*, 2013; Zapf *et al.*, 2000; Zhao *et al.*, 2008) as well as *Escherichia coli* CheY variants in complex with imidazole (Page *et al.*, 2015), or yet two autophosphorylating HKs where the phosphoryl group is poised to migrate from ATP to the reactive histidine (Casino *et al.*, 2014; Mechaly *et al.*, 2014). In contrast, HK:RR TCSs with unidirectional P~His \rightarrow Asp transfer, such as *B. subtilis* DesK:DesR (this study), *Thermotoga maritima* HK853:RR469 (Casino *et al.*, 2009) and *Rhodobacter sphaeroides* CheA3:CheY6 (Bell *et al.*, 2010), consistently show $>$ 6.6 Å HisN ϵ 2-AspO δ 1 distances and correlated large reaction coordinate distances when these could be measured (Table 6). It will be interesting to test whether the phosphorelay pair ChpT:CtrA from *Brucella abortus* (Willett *et al.*, 2015) displays reversible transfer, as anticipated on the basis of the reported 5.9 Å HisN ϵ 2-AspO δ 1 distance.

In a loose reaction center like that of DesK:DesR, a largely dissociated meta-phosphate intermediate would be stabilized migrating toward the positive charges of the Mg²⁺ cation and the conserved Lys_{102(RR)} in the RR partner. The position of these positive charges, next to the RR phosphorylation site, introduces an intrinsic asymmetry within the reaction center, which must translate into unequal likelihood for phosphoryl-transfer directions. As the Mg²⁺ position becomes more asymmetric, the P~Asp \rightarrow His back-transfer would be less favorable, explaining unidirectionality. The ratio between the Mg²⁺-Asp(O δ 1) and the Mg²⁺-His(N ϵ 2) distances, can be used to quantitate such asymmetry within reaction centers. This ratio was calculated for different TCS complexes (Table 6), indeed clustering those complexes that catalyze reversible P~Asp \rightarrow His/P~His \rightarrow Asp reactions, apart from the ones that carry out P~His \rightarrow Asp unidirectional transfer (Figure 5D). Signaling pathways appear to have evolved associative or dissociative mechanisms, corresponding respectively to more reversible or irreversible reactions, such that they are fit to drive a defined directionality in the flow of information.

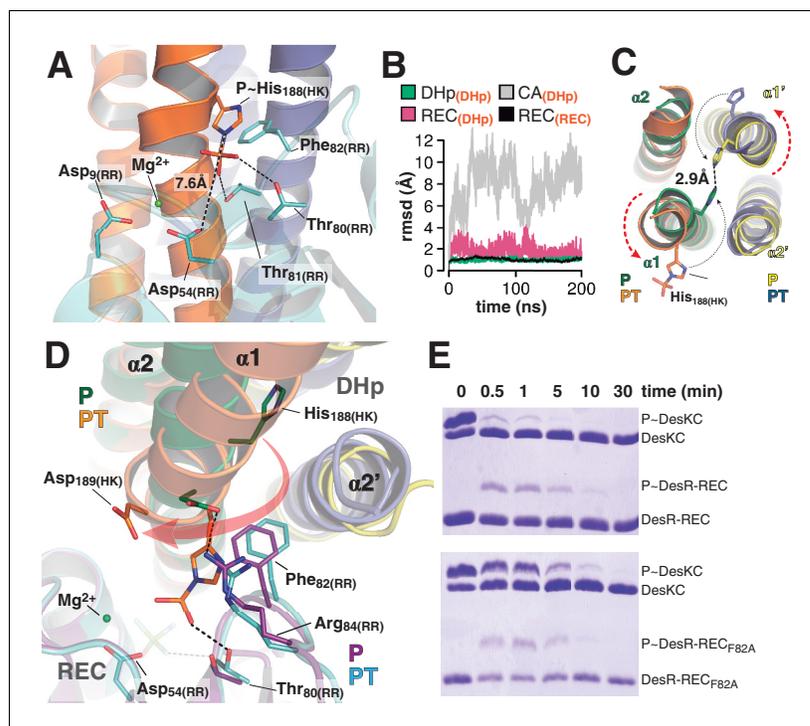


Figure 4. At the reaction center: the phosphatase to phosphotransferase transition. (A) Cartoon model of the phosphotransferase reaction center prior to DesR phosphorylation (see model construction details the Materials and methods section). DesK protomers (orange and blue) and DesR (cyan, transparent for clarity) are displayed with selected key residues in sticks colored by atom. Mg^{2+} (green sphere) is already in place, coordinated by the two shown Asp residues and water molecules (not included). Phosphoryl moiety interactions with DesR, and the reactive Asp-His distance are indicated. (B) Evolution of the atomic coordinates of the phosphotransferase complex along molecular dynamics (MD) calculations. Selected HK or RR domains were structurally aligned (marked with orange subscripts on each curve's label on top), to thereafter compare the evolving MD model with the initial experimental structure (calculating rmsds of chosen domains as marked in black fonts on each curve's label on top). Resulting rmsds for all C α atoms of chosen domains are plotted (colored curves) as a function of time. Note that the time lapse is enough to detect large CA mobility (gray curve) whereas the DHp:REC complex remains attached and stable (pink curve). (C) Cartoon illustration of the HK DHp domains of the phosphatase complex (P), with its two HK protomers in green and yellow, superposed onto the phosphotransferase (PT) in orange and blue. Residue His_{188(HK)} (in sticks) reveals the rotational rearrangement between both states. (D) Similar phosphatase vs phosphotransferase illustration as in (C), along a different view. The RR partner is now shown (magenta for the phosphatase complex [P], cyan for the phosphotransferase [PT]). The Arg_{84(RR)}:Asp_{189(HK)} salt bridge is disrupted in the PT complex due to the DHp rotational shift (red arrow). Note the shift of the RR $\beta 4\alpha$ loop, including Phe_{82(RR)}, propagating toward Thr_{80(RR)}. The latter is positioned at H-bonding distance to the phosphoryl group either on the P-His or on the P-Asp residue (the BeF₃⁻ moiety in the P complex is transparent), with a shift of 1.5 Å of its side chain O atom. (E) Phosphotransfer kinetics comparing wt DesR-REC (top panel) with phosphate lid mutant DesR_{F82A}-REC (bottom panel), revealed by Phos-tag SDS-PAGE. DOI: 10.7554/eLife.21422.014

The following figure supplements are available for figure 4:

Figure supplement 1. Electron density maps and structural similarity of wt P-DesKC compared to DesKC_{H188E} in the phosphotransferase complex.

DOI: 10.7554/eLife.21422.015

Figure supplement 2. Structural details of the reaction centers.

DOI: 10.7554/eLife.21422.016

Figure supplement 3. Phosphate lid opening for RR dephosphorylation.

DOI: 10.7554/eLife.21422.017

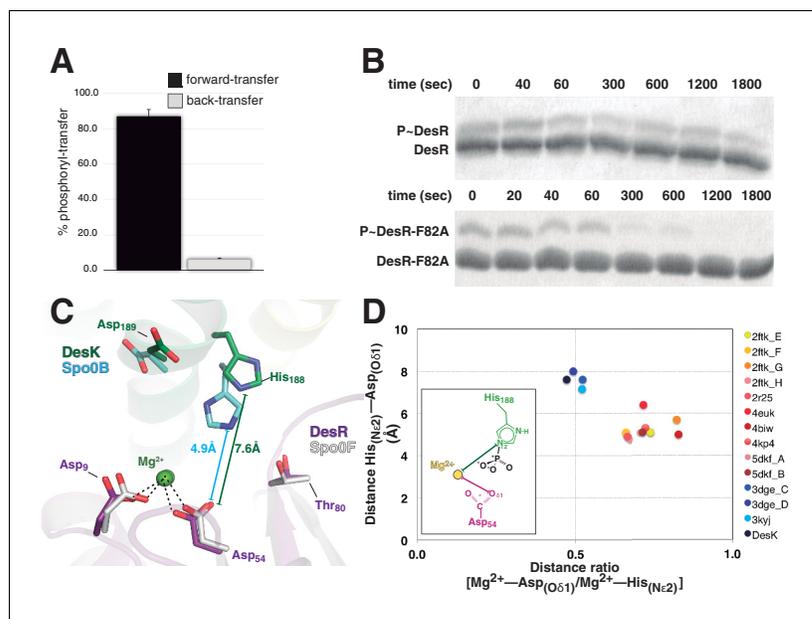


Figure 5. Dissociative phosphotransfer and asymmetric position of the divalent cation. (A) Phosphoryl-transfer reactions were analyzed in vitro using Phos-tag SDS-PAGE. Band intensities corresponding to unphosphorylated and phosphorylated species were quantified by densitometry. Forward- and back-transfer directions were compared, with bars representing the amount of phosphotransfer in each direction, plotted as the percentage of phosphorylated DesKC with respect to its total initial amount (100%). Forward phosphotransfer (black bar) was measured by incubating pre-phosphorylated P-DesKC and DesR-REC. Back-transfer (gray bar) was assayed by incubating pre-phosphorylated P-DesR and DesKC. The standard deviation for triplicate replicas are shown as error bars. (B) Stabilization of the phosphorylated DesR species by the phosphorylation lid. The intrinsic dephosphorylation velocities of P-DesR-REC and P-DesR_{F82A}-REC were compared using Phos-tag SDS-PAGE. One of three similar replicas is shown. (C) Structural superposition of the DesK:DesR phosphotransferase complex (colored in green:magenta) onto the *B. subtilis* Spo0B:Spo0F complex (PDB 2FTK, depicted in cyan:white), aligning the DesR and Spo0F REC domains. Selected key residues are labeled and numbered according to DesR's sequence. Note the strictly conserved position of the divalent cation (green sphere). The relative shift in the position of the phosphorylatable histidine (His₁₈₈ in DesK corresponds to His₃₀ in Spo0B), is highlighted by depicting the His_(Nε2)-Asp_(Oδ1) distances, not including the phosphoryl group for clarity. Note that besides the His shift, the overall DHP:REC binding interface is otherwise preserved. (D) Plot of His_(Nε2)-Asp_(Oδ1) distance vs degree of asymmetry in the position of the Mg²⁺ cation, comparing independently refined TCS complexes (color-labeled). The inset shows the two interatomic distances used to calculate the ratio plotted in the x-axis. Note the clustering in two groups, which discriminate complexes catalyzing reversible (yellow-to-brown range of points) vs irreversible (blue range of points) phosphotransfer reactions.

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Discussion

Signaling requires *specificity*, to ensure that a given stimulus is linked to a defined adaptive response while minimizing cross-talk; *efficiency*, to avoid wasting cellular energy through futile cycles; and *directionality*, to guarantee the information is directed from the stimulus to the output response. Two component systems constitute a particularly interesting biological model to understand signaling, considering that so many histidine kinases catalyze both kinase and phosphatase reactions according to their functional status, implying exquisite regulation mechanisms at play.

By solving the crystal structures of a *bona fide* HK in complex with its cognate RR in two distinct functional states, such molecular mechanisms of TCS unidirectional signal transmission are being uncovered. These structures correspond to snapshots of the phosphotransferase reaction just prior of actual phosphoryl migration, and of the phosphatase reaction in a pre-dephosphorylation complex.

Table 6. Summary of reactive His_(HK)-to-Asp_(RR) distances in reported TCS protein complexes.

	Group (based on distance discriminants*)	Complex partners (N° complexes per ASU†)	Distance his(Nε2)-Asp(Oδ1) (Å)‡	Reaction coordinate distance (Å)§	Distance ratio [Mg ²⁺ -Asp(Oδ1)/Mg ²⁺ -His(Nε2)]	Pdb (ID)	Phosphotransfer reaction direction	Remarks
Phosphorelays	I	Spo0F: Spo0B ^{¶¶} (4)	5.7; 5.3; 5.1; 5.1	4.1; 3.8; 3.6; 3.5	0.82; 0.74; 0.65; 0.67	2FTK	P~Asp _(Spo0F) →His _(Spo0B)	Spo0B bears a DHp domain
		SLN1: YPD1 ^{**} (1)	4.9	3.2	0.67	2R25	P~Asp _(SLN1) →His _(YPD1)	YPD1 bears an HPt domain
		AHK5: AHP1 ^{††} (1)	6.4	ND[8]	0.72	4EUK	P~Asp _(AHK5-REC) →His _(AHP1)	AHP1 bears an HPt domain
		ChpT: CtrA ^{§§} (2)	5.9 ^{¶¶¶}	ND ^{‡‡}	0.64 ^{***}	4QPJ	P~His _(ChpT) →Asp _(CtrA)	ChpT bears a DHp domain P~Asp _(CtrA) →His _(ChpT) remains to be confirmed
HKs undergoing auto-phosphorylation	I	CpxA: ATP ^{†††} (1)	5.0 ^{‡‡‡}	3.4	0.82	4BIW	ATP→His _(CpxA)	AMP-PCP was used as non-hydrolyzable ATP analogs (between phosphates β and γ there is a C atom substituting the O)
		HK853/EnvZ chimera: ATP ¹⁴ (1)	5.3 ^{‡‡‡}	3.6	0.72	4KP4	ATP→His _(HK853)	
RR-mediated phosphorylation of imidazole	I	Imidazole: CheY ^{¶¶¶¶} (2)	5.1; 5.1	3.3; 3.3	0.71; 0.73	5DKF	P~Asp _(CheY) →imidazole	In vitro engineering of a phosphorelaying system from CheY to PhoR, using imidazole as a rudimentary HPt
HK:RR complexes	II	HK853: RR468 ^{****} (2)	7.6; 8.0 ^{††††}	5.7; 6.1 ^{††††}	0.51; 0.48 ^{††††}	3DGE	P~His _(HK853) →Asp _(RR468)	HK:RR complex, snapshot of the phosphatase state according to the authors.
		DesK: DesR ^{††††} (2)	7.6 ^{§§§§}	5.8 ^{§§§§}	0.47 ^{§§§§}	5IUK	P~His _(DesK) →Asp _(DesR)	HK:RR complex in the phosphotransferase state
		CheA3: CheY6 ^{¶¶¶¶¶} (1)	7.3	ND ^{‡‡}	0.52 ^{*****}	3KYJ	P~His _(CheA3) →Asp _(CheY6)	3KYJ is not phosphorylated on either partner; 3KYI with P~His on CheA3 displays an unproductive P~His rotamer, otherwise confirming 3KYJ's 3D organization

* Distance between reactive His(Nε2)-Asp(Oδ1) less than or greater than 6.5Å, and distance ratio [Mg²⁺-Asp(Oδ1) / Mg²⁺-His(Nε2)] less than or greater than 0.6

† ASU = asymmetric unit

‡ Each distance corresponds to the one measured in each one of the independently refined complexes in the ASU.

§ Distances are reported in the same order as in the previous column, with correspondence among same individual complexes.

¶ Bacillus subtilis (Varughese et al., 2006)

** Saccharomyces cerevisiae (Zhao et al., 2008)

†† Arabidopsis thaliana (Bauer et al., 2013)

‡‡ Not determined : no phosphoryl group or phosphoryl-mimetic present in the structure.

§§ Brucella abortus (Willett et al., 2015)

¶¶ Only one of the two complexes in the ASU (ChpTchainA: CtrAchainC, display the reactive His and Asp properly oriented poised for reacting. Only one distance is thus recorded.

*** No Mg²⁺ cation was actually bound on CtrA. A metal atom was modeled by superimposing the structure of RR468 with bound BeF₃ (PDB 3GL9), one of the top ranking structures in multiple structural alignments with CtrA (DALI Z score 12.9, rmsd 0.8Å superimposing 98 αCs).

††† Escherichia coli (Mechaly et al., 2014)

‡‡‡ In the autophosphorylation complexes the distance is recorded between His(Nε2) and the position of the O between phosphates β and γ.

§§§ Thermotoga maritima (HK853) / E. coli (EnvZ) (Casino et al., 2014)

¶¶¶¶ E. coli (Page et al., 2015)

***** Thermotoga maritima (Casino et al., 2009)

†††† These distances have been calculated by superimposing the phosphorylation-mimetic structure of RR468 alone with bound BeF₃ (PDB 3GL9), onto 3DGE, in order to use better estimations of the receiver Asp position as well as of the phosphorus atom.

†††† B. subtilis, this report.

§§§§ Glu188 present in the crystal structure was substituted by wt P~His following superposition of wt phosphorylated DesKC (PDB 5IUM) onto one of the phosphotransferase DesKCH188E:DesR-REC complexes in the ASU (chains A-B:E). Distances are reported after energy minimization. See Materials and methods.

¶¶¶¶ Rhodobacter sphaeroides (Bell et al., 2010)

***** The position of Mg²⁺ cation was modeled using PDB 4TMY as template.

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Specificity determinants within a loose and slippery interface

The small number of interactions and low surface complementarity that we observe in the several DesKC:DesR complexes (**Figure 2F**) are consistent with the observed 'slippery' nature of the protein:protein interface, displaying significant shifts in the relative positions of both partners. Further supporting that this ensemble of conformations is functionally relevant, molecular dynamics simulations of the phosphotransferase complex (**Figure 2—figure supplement 2**) recapitulated the gliding shifts of one partner with respect to the other, reaching the different arrangements observed in the crystal structures at different moments of the simulations (**Figure 2—figure supplement 2B**). Loose HK:RR association architectures can actually be observed in other TCS-related complexes (Zapf et al., 2000; Casino et al., 2009; Willett et al., 2015), suggesting it is a universal feature. This explains the considerable promiscuity reported in different TCSs in vitro assays, if long enough incubation times are allowed for a given HK to phosphotransfer to different RRs, including surrogate partners (Skerker et al., 2005). HK:RR interactions thus ensure high specificity, but also avoiding exceedingly high stability, which would hinder proper functional modulation and, ultimately, signal transmission altogether. Evolutionarily selected to be loose, slippery interfaces are inherently tolerant for substantial sequence variation of protein:protein contacting residues. Such tolerance is consistent with the extensive degeneracy found in TCS pairs (Podgornaia and Laub, 2015), likely a general feature of HK:RR interactions. However, a permissive tendency for substitutions would appear to be contradictory with the high protein:protein specificity requirements of these systems. The DesK:DesR structures now provide with molecular details explaining why epistasis is found as a key property of HK:RR interfaces (Podgornaia and Laub, 2015). Epistatic amino acid changes are those that depend on the presence of other substitutions, leading to different effects in combination than individually. While interfacing residue replacements on any one of the TCS components can easily be accommodated due to the loose character of the protein:protein association, epistatic substitutions in the same component and/or in the other partner, are called to play essential stabilizing or destabilizing compensatory roles. Epistatic substitutions are thus selected, such that partner recognition and non-cognate discrimination are maintained, but without leading to overstable, non-functional complexes.

A novel conformation of the response regulator when complexed with the histidine kinase

Our data indicate that the HK is largely unable to discriminate binding to RR or P~RR on the basis of structural determinants. The structures show the RR REC domain adopting identical conformations in both phosphatase and phosphotransferase complexes, presenting active- and inactive-like features in different regions of the protein, arranging the RR's active site for phosphoryl group migration (**Figure 3**). The two complexes also show equivalent HK:RR interfacing surfaces, with all major contacts preserved. Among the few differential contacts, they mainly implicate the HK's CA domain, but unlikely to grant discrimination ability given the highly similar conformation of the RR in those regions.

Calorimetry data are consistent with the crystallographic evidence, revealing only modest differences between DesR-REC:DesK_{H188V} and DesR-REC:DesK_{H188E} binding affinities (**Figure 1** and **Table 2**), further supporting that both HK functional states interact with DesR-REC with similar strengths. Collecting calorimetric data by titrating DesKC with P~DesR-REC would be interesting to directly compare with the unphosphorylated form, but binding confounds with simultaneous P~RR dephosphorylation, and the addition of BeF₃⁻/Mg²⁺ to mimic RR phosphorylation resulted in protein

aggregation. We did succeed in engineering a different phosphatase-trapped DesKC mutant, by insertion of additional heptad repeats in the coiled-coil segment (to be published elsewhere), indeed displaying equivalent DesR-REC-binding K_D figures compared to DesKC_{H188E}. HK:RR binding data in other TCSs are scarce, particularly exploring differential affinities according to the proteins' phosphorylation status. The few cases dealing with HKs that switch between phosphatase/phosphotransferase-competent states, do provide further evidence in support of non-discrimination. EnvZ binds OmpR and P~OmpR with equivalent affinities (Yoshida *et al.*, 2002), as well as PhoQ binding to PhoP or P~PhoP (Castelli *et al.*, 2003). Future studies shall uncover the relevance of additional mechanisms at the cell level that might favor particular HK:RR pairs to form, such as absolute and relative TCS protein concentrations, stability of non-phosphorylated HK:RR species in complex, sub-cellular spatial dynamics of each partner, and precise time-courses for HK and RR phosphorylated species to appear/disappear.

A HK:RR concerted switch controls efficiency along the signaling pathway

Considering all the evidence, we can now propose a conceptual model explaining the molecular workings of the DesK/DesR signaling pathway (Figure 6). When the pathway is turned off, HK auto-phosphorylation is inhibited by: 1) hindering CA domains mobility, holding ATP substrates far from the His_{188(HK)} phosphorylation site; and, 2) DHP helical rotation, making the His_{188(HK)} side chains inaccessible (Figures 4D and 6A, Video 1). This 'open' autokinase-off configuration is linked to a folded N-terminal coiled-coil, in which DesK can bind P~DesR and trigger phosphatase catalysis. Linked to P~DesR phosphate lid opening, Asp_{189(HK)} binds to Arg_{84(RR)}, and Phe_{82(RR)} becomes inserted into a DHP pocket (Figure 6A, Video 1). Phe_{82(RR)} is thus removed from the closed position, which would otherwise clash with an incoming nucleophilic water molecule (Figure 4—figure supplements 2 and 3) that brings about dephosphorylation (Pazy *et al.*, 2010). DesK Gln_{193(HK)} is well located to position the catalytic water to perform dephosphorylation, consistent with the essential role of the homologous Gln residues in the phosphatase reactions mediated by the phosphatase CheZ (Zhao *et al.*, 2002) or by the HK NarX (Huynh *et al.*, 2010). The rotational movement of His_{188(HK)}, placing it at >18 Å away from the reactive Asp_{54(RR)}, makes phosphoryl back-transfer P~DesR→DesK impossible (Figure 4E). Although sequence divergence makes the DHP rotational shift unlikely in HisKA HKs (Marina *et al.*, 2005), DHP kinking by means of a conserved proline (Casino *et al.*, 2014; Mechaly *et al.*, 2014) could regulate the separation of the phosphorylatable His from the reaction center, analogously to the HisKA_3 rotational switch. Additional mechanisms that stabilize an 'open' state of HisKA HKs via extensive CA-DHP interfaces, have been reported to favor the phosphatase reaction (Dubey *et al.*, 2016) further supporting our hypotheses.

This DHP helical switch is thus coupled to signal sensing, such that sensory domain rearrangements lead to intra-cytoplasmic HK coiled-coil disruption. Possibly implying trans-membrane helix tilting, or other signal-triggered helical reconfigurations, a direct connection of sensory to intracytoplasmic α -helices can be posited. This switch can be readily understood as an order-disorder transition, which has been observed in HKs and other modular proteins, to work as an intraprotein signaling mechanism (Schultz and Natarajan, 2013). Helical rotational shifts have also been uncovered in structural elements within chemotaxis and TCSs, such as in HAMP domains, likely revealing common underlying transduction mechanisms eventually controlling their physiologic directionality (Airola *et al.*, 2013).

HK activation leads to autophosphorylation and subsequent switching to its phosphotransferase state (Figure 6B). Comparing the conformation of the phosphorylatable histidine between HKs performing auto-phosphorylation (Casino *et al.*, 2014; Mechaly *et al.*, 2014), and the phosphotransferase complex (this study), the His residue shows ~180° flipping of its imidazole side chain (Figure 6B). During auto-phosphorylation, His_{N δ 1} has been observed to H-bond with the ultra-conserved acidic His + 1 residue, which corresponds to Asp_{189(HK)} in DesK. Such H-bond enhances His N ϵ 2 nucleophilicity (Quezada *et al.*, 2005), and indeed a carboxylate at His + 1 is critical for autophosphorylation in a number of HKs (Willett and Kirby, 2012). Once phosphorylated, the His disrupts its contact with this neighbor carboxylate group, stabilizing the P~HK species and leading to His χ -2 dihedral angle flipping (Figure 6C). Subsequent re-protonation of His_{188N δ 1}, triggered by RR binding, is consistent with P-N bond weakening, as a first step towards phosphotransfer (Figure 4E). This

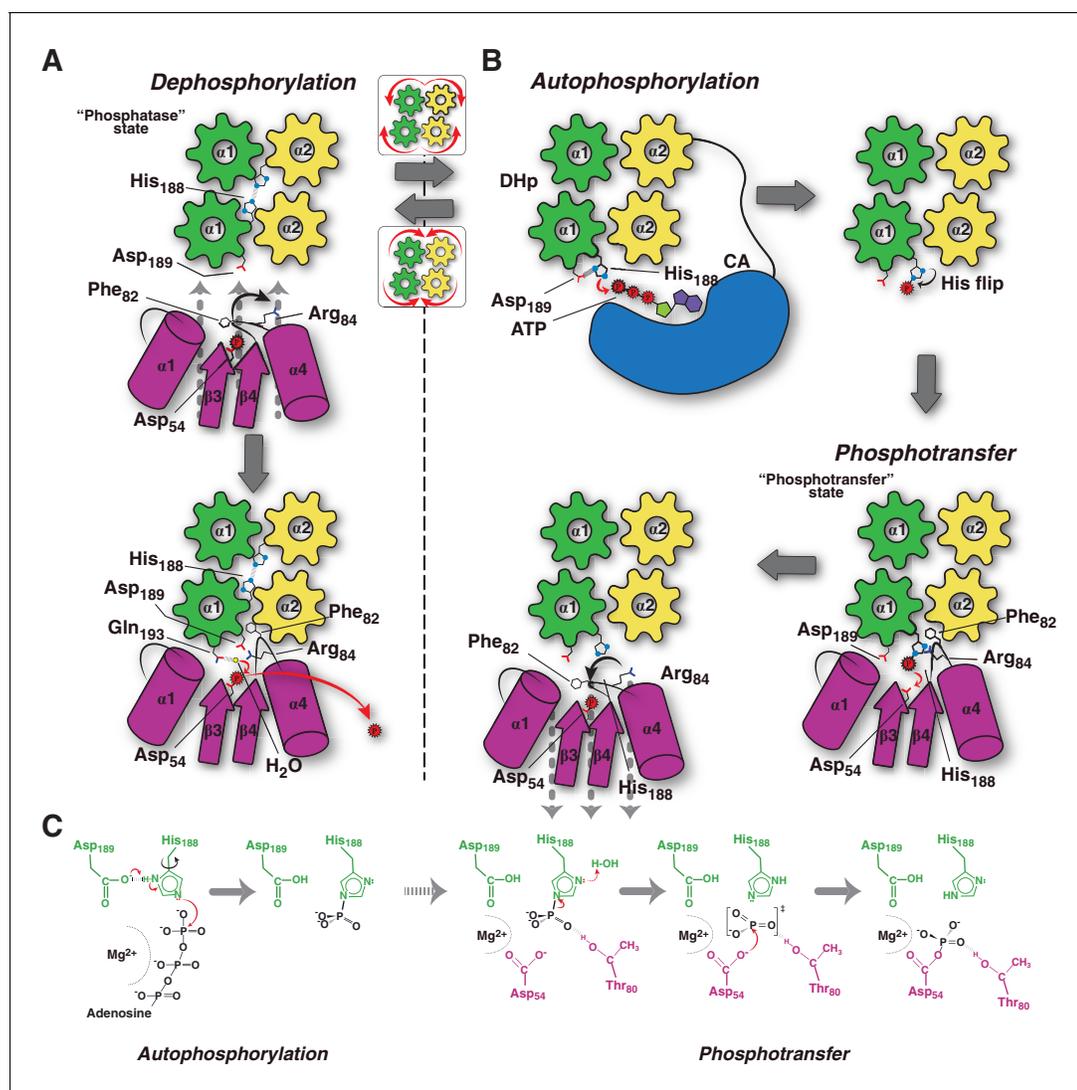
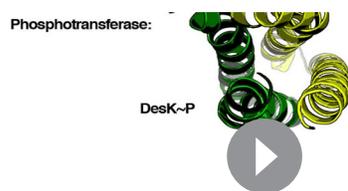


Figure 6. Conceptual model of TCS unidirectional signaling. (A) P~DesR dephosphorylation catalysis is favored by the opening of the RR phosphorylation lid, via Phe₈₂(RR) insertion into an HK hydrophobic pocket, and an ionic interaction between Asp₁₈₉(HK) and Arg₈₄(RR). The Phe₈₂(RR) movement allows for entry of a nucleophilic water (yellow sphere) positioned by Gln₁₉₃(HK). In the phosphatase state the phosphorylatable His₁₈₈(HK) is buried within the DHp domain core, avoiding autophosphorylation and phosphoryl back-transfer reactions. (B) Signal-mediated pathway activation triggers the cogwheel rotation of the HK DHp α -helices, via coiled-coil disruption. The His₁₈₈(HK) becomes exposed to the solvent, CA domains are released, and autophosphorylation thus enabled. Binding to unphosphorylated RR positions the P~His₁₈₈(HK) into a suitable orientation for phosphoryl-transfer to occur through a loose (dissociative) nucleophilic substitution. Movement of Thr₈₀(RR) is coupled with phosphoryl group migration, triggering RR's phosphate lid closure and P~RR release. (C) Reaction scheme summarizing the autophosphorylation and phosphotransfer reactions. Stabilization of a protonated N δ 1 tautomer of the reactive histidine, is required in autophosphorylation. In keeping with the imidazole aromaticity, the nucleophilicity of N ϵ 2 is thus finely regulated, allowing for phospho-acceptor/donor roles of the His.

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mechanism might be critical in dissociative P~HK→RR reactions, which lack the simultaneous nucleophilic attack that the receiver Asp_(RR) carries out in associative ones.

Our model predicts an important role of the highly conserved Thr₈₀(RR) in stabilizing an intermediary metaphosphate group, ideally positioned to accompany the migrating phosphoryl moiety. This threonine has been substituted in CheY (Ganguli *et al.*, 1995; Appleby and Bourret, 1998) and DosR (Gautam *et al.*, 2011), suggesting it is not directly engaged in catalysis, but does play important roles in phosphoryl-transfer in vitro and proper in vivo signaling. Systematic mutagenesis studies substituting this threonine in different TCS pairs are needed to be conclusive about its role in phosphotransfer. Once the phosphotransferase reaction is complete, immediate dephosphorylation of



Video 1. Animation illustrating the phosphotransferase to phosphatase transition, within the model of TCS-mediated phosphorylation and dephosphorylation of the RR component. As an animated support to main **Figure 6**, the fixed ground states correspond to crystal structures, and the moving intermediates in between obtained by linearly interpolated morphing, as implemented in Pymol v 1.8.2

(Schrodinger, LLC, 2015). This animation does not include the kinase's auto-phosphorylation step, shown at the beginning of **Figure 6B**. Structures of phosphorylated ('active') and non-phosphorylated ('inactive') DesR, when shown separate from the HK DesK, correspond respectively to PDB structures 4LE0 and 4LE1 (Trajtenberg et al., 2014). On the other hand, the fixed positions of P~DesK in complex with DesR (phosphotransferase state), and of DesK in complex with P~DesR (phosphatase state), correspond to the structures reported in this work (PDBs 5IUK and 5IUN, respectively). The reactive His_{188(HK)} on DesK is depicted as spheres, as well as the active Asp_{54(RR)} and Phe_{82(RR)} on DesR. The rest of the proteins are rendered as cartoons. Chains A and B of DesK are colored in green and yellow, respectively. DesR is blue when not phosphorylated, and magenta once activated by phosphorylation. The P~His was modeled into DesK as explained in the text and Materials and methods. P~DesR corresponds to the phosphomimetic BeF₃⁻ bonded to Asp_{54(RR)}.

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the P~Asp_{54(RR)}, as well as P~DesR→DesK back-transfer, are minimized. The β4α4 loop in DesR plays a key role in these controls, remodeling its configuration in concert with DesK-association. This loop, acting as a phosphate lid, is open in the phosphotransferase complex, enabling the transfer reaction (**Video 1**). Yet, the phosphate lid is expected to close in, following the phosphoryl migration during transfer, due to the strong phosphoryl-Thr_{80(RR)} interaction. Phosphotransfer thus results in a fully activated RR configuration, with a closed phosphate lid that ensures the release of a stable P~DesR species (**Figure 6B** and **Video 1**), as observed in active free RR structures (Trajtenberg et al., 2014).

The β4α4 loop amino acid sequence is not well conserved among different RRs and RR families (Page et al., 2016), despite its direct implication in forming the HK:RR interface. Conspicuous examples of variable residues can be mapped to Thr_{81(RR)}, well positioned to interact with the phosphoryl group of P~DesK, or yet Phe_{82(RR)} and Arg_{84(RR)}, key players in the open/closure switch of DesR's phosphate lid. Some of these positions had already been pinpointed as highly variable, yet functionally relevant (Thomas et al., 2013; Immormino et al., 2016; Page et al., 2016), recognizing positions T + 1 and T + 2 (starting from T, the conserved position Thr_{80(RR)}) among others, as important modulators of RRs' catalytic activities. Particular sequence signatures of these variable motifs, have been found to correlate with RR families, in turn connected to the distinct physiological constraints of each family's signaling functions (Page et al., 2016). Our data now illustrate an additional selective pressure dimension to such variable positions, linked to the evolution of the HK:RR interface. Beyond sequence variability, the RR β4α4 loop is bound to play functional roles, as it is always located at the HK:RR interface in available structures. Muta-

genesis studies with functional readouts including HK-mediated activities are needed, together with a larger number of 3D structures of HK:RR complexes from different TCS families, in order to delimit universal TCS mechanisms and system-dependent variations.

Associative vs dissociative phosphotransfer: evolution of signaling directionality

Regulating the direction of information flow is central to signaling performance. The energies of P-N and P-O bonds engaged in phosphoryl-transfer reactions among histidine and aspartate, appear to be equivalent, explaining a few known reversible systems (Burbulys et al., 1991). However, and except for such systems where P~Asp→His phosphorylation has been selected (such as in phosphorelays), phosphoryl back-transfer (P~RR→HK) is typically minimized. Alternative sources of reactional asymmetry, other than bonding energetics, must be at play. Taking into account that phosphotransfer is a nucleophilic substitution at phosphorus, the reaction coordinate distance (i.e. between phosphorus and the entering atom, before the transfer reaction has begun) strongly suggests that P~DesK→DesR phosphotransfer occurs through a largely dissociative, or 'loose' mechanism

(Lassila *et al.*, 2011): a reaction coordinate distance of ≥ 4.9 Å provides room for a fully dissociated metaphosphate intermediate (Mildvan, 1997). This distance is directly correlated with the one separating side chain N and O atoms, respectively on the reactive His and Asp residues. Considering van der Waals radii, full dissociation of the phosphate intermediate is predicted for N-O distances of ≥ 6.6 Å. This threshold distance classifying phosphotransfer reactions into loose vs tight nucleophilic substitutions, nicely discriminates signaling complexes involved in, respectively, phosphorylations from the histidine imidazole group, or towards it (Figure 5D, Table 6). The His-Asp distance thus correlates with associative vs dissociative phosphotransfer, but also with the symmetry of the Mg^{2+} cation with respect to the entering and leaving atoms (Figure 5C and D). Increased back-transfer has been observed when the cation is removed (Albanesi *et al.*, 2004; Lukat *et al.*, 1990; Shi *et al.*, 1999), further supporting that Mg^{2+} asymmetry favors irreversible P~His→Asp transfer reactions. Furthermore, mutations that likely increase the distance between TCS partners, without impeding their association, have been reported to affect differentially the forward- and back-transfer directions in otherwise reversible phosphoryl-transfer steps (Janiak-Spens *et al.*, 2005).

The DesK:DesR structures disclose the catalytic centers for TCS-mediated dephosphorylation and phosphotransfer reactions, built with contributing residues from both partners. Moreover, the HK serves as the driving component to modulate the construction of a given active site, selecting a conformation of the RR that is prone for transfer reactions on its catalytic aspartate. Except for the phosphate lid, the REC domain bound to the HK largely resembles the RR's active form, shifting the position of the $\beta 1\alpha 1$ loop and helix $\alpha 1$, now able to recruit the Mg^{2+} cation essential for phosphoryl-transfer catalysis. The phosphatase/phosphotransferase transition, switches the distance and orientation of the HK histidine side chain with respect to the RR aspartate, enabling for the distinct reaction mechanisms to take place, and ultimately ensuring the unidirectional flow of information.

Materials and methods

Cloning and mutagenesis

For protein crystallization purposes, recombinant protein expression plasmid pACYC-DesK_{STAB}:DesR-REC was generated by sub-cloning DesK_{STAB} from pHPKS/Pxyl-desKSTA (Saita *et al.*, 2015) into pACYC-DesK_{H188E}:DesR-REC (Trajtenberg *et al.*, 2014) using RF cloning (Unger *et al.*, 2010) with primers STAB_F (5'-CCTGTATTTTCAGGGATCC GGTATTATAAAAACCTCGCAAG-3') and STAB_R (5'-GTCAGACACTGTAATCACAACCTCCTCCAG-3'). The two expression cassettes present in the pACYC-DesK_{STAB}:DesR-REC encode the two recombinant proteins each fused to a His-tag and a TEV protease cleavage site.

For phosphotransfer assays, two DesR point mutants Phe82Ala and Arg84Ala were designed by mutagenesis of pQE80-DesR-REC (Trajtenberg *et al.*, 2014), using mutagenic primers DesR_Fw (5'-GATTAGTATATTTATTGCAGAAGATCAGCAAATGC-3'), F82A_Rev (5'-CGGGTCTGGCGGCGGTGG TTAAG ATGATAATTTG-3') and DesR_R84A_Rev (5'-GAAAGTAACCGGGTGCGGCGAAGGTGG TTAAG-3'), resulting in plasmids pQE80-DesR-REC_{F82A} and pQE80-DesR-REC_{R84A}. Cloning procedures, as well as protein expression, crystallization and structure solution methods, are described in more detail at Bio-protocol (Imelio *et al.*, 2017).

Protein expression, purification and crystallization

Recombinant proteins to be used in crystallization were co-expressed from pACYC-DesK_{STAB}:DesR-REC and pACYC-DesK_{H188E}:DesR-REC plasmids, transformed in *Escherichia coli* strain BL21 (DE3) (Novagen). For functional assays proteins were expressed from plasmids pQE32-DesK (Albanesi *et al.*, 2009), pQE80-DesR-REC (Trajtenberg *et al.*, 2014), pQE80-DesR-REC_{F82A} and pQE80-DesR-REC_{R84A}, in *E. coli* TOP10F' (Invitrogen). Proteins were expressed and purified as previously described (Albanesi *et al.*, 2009; Trajtenberg *et al.*, 2014). For the DesK-DesR complexes the last size exclusion chromatography step (HiLoad 16/60 Superdex 75 preparation grade column; GE Healthcare) was used to select those fractions corresponding to the complex only.

Hanging-drop crystallizations were performed at 20°C in Linbro plates. The DesK_{STAB}:DesR-REC complex (10 mg/mL protein prepared in 5 mM AMP-PCP, 10 mM $MgCl_2$) crystallized in 1 mL mother liquor [30% (w/v) PEG 4000, 0.1 M Tris.HCl pH 8.5, 0.2 M Li_2SO_4]. Protein drops were set up by mixing 0.8 μ L mother liquor + 2 μ L protein solution + 1.2 μ L additive solution [27% (v/v) PEG 600, 0.1 M

MES pH 6.5, 0.15 M MgSO_4 , 5% (v/v) glycerol]. Initial crystals were optimized by microseeding. Cryo-protection was achieved by slowly adding 4 μL of cryoprotection solution [32% (w/v) PEG 4000, 0.1 M Tris.HCl pH 8, 0.2 M Li_2SO_4 , 20 mM MgCl_2 , 18 mM BeF_3^- , 5 mM AMP-PCP, 15% (v/v) glycerol], then soaked in 100% cryoprotection solution and frozen in liquid N_2 . The DesKC_{H188E}:DesR-REC complex (8.3 mg/mL protein prepared in 5 mM AMP-PCP, 20 mM MgCl_2) crystallized in mother liquor 18% (w/v) PEG 3350, 0.3 M tri-potassium citrate. Protein drops were set up by mixing 2 μL protein + 2 μL mother liquor. Cryo-protection was achieved by quick soaking in 20% (w/v) PEG 3350, 0.3 M tri-potassium citrate, 5 mM AMP-PCP, 25% (v/v) glycerol, and 20 to 150 mM MgCl_2 + 0 to 5 mM BeF_3^- . wt P~DesKC was crystallized as described (Albanesi *et al.*, 2009), except that 5 mM ATP was used throughout instead of AMP-PCP.

X ray diffraction data processing, crystal structure solution, refinement and analysis

Single crystal X ray diffraction was performed with a copper rotating anode home-source (Protein Crystallography Facility, Institut Pasteur de Montevideo) or synchrotron radiation (Soleil, France). Data processing was performed with autoPROC (Vonrhein *et al.*, 2011). Structures were solved by molecular replacement (McCoy *et al.*, 2007), using an in silico-generated model of a DesKC:DesR complex (Trajtenberg *et al.*, 2014) as search probe. The other domains were then manually located in the electron density maps, and complete models were rebuilt using Coot (Emsley *et al.*, 2010) and refined with Buster (Bricogne *et al.*, 2009). Validation was done throughout and towards the end of refinement, using MolProbity tools (Chen *et al.*, 2010). DesKC:DesR-REC surface complementarity was calculated according to (Lawrence and Colman, 1993), resulting in figures of ~0.6. Visualization of protein models and structural analyses, figure rendering and morphing for animation were performed with Pymol (Schrodinger, LLC, 2015). Software for data processing, structure determination and analysis was provided by the SBGrid Consortium (Morin *et al.*, 2013).

Small angle X-ray scattering data acquisition and analysis

SEC-SAXS experiments were carried out at beamline SWING (Soleil synchrotron, France). Purified DesK-DesR complexes samples were brought to 10 mg/mL in 50 mM Tris.HCl pH 8.0, 300 mM NaCl, and injected into a Superdex 75 5/150 GL column (GE Healthcare Biosciences) equilibrated in the same buffer. Samples were eluted at constant flow (0.15 mL/min) and loaded into capillary cell for X-ray exposure. Recorded frames were processed with Foxtrot (David and Pérez, 2009) following standard procedures. Subsequent analyses of the scattering data were performed with tools from the ATSAS (Konarev *et al.*, 2006). Forward scattering $I_{(0)}$ and the particle's radius of gyration (R_g) were estimated using the Guinier approximation. $I_{(0)}$ real was calculated as the extrapolated intensity at zero scattering angle. Pair-distance distribution functions were calculated from the scattering patterns with GNOM, which also provides maximum particle dimension (D_{max}) and R_g (real) values. Molecular mass (MM) of particles was estimated from the volume-of-correlation (V_c) values (Rambo and Tainer, 2013). Theoretical scattering patterns from atomic models were calculated and fitted to experimental curves using CRY SOL. Mixture analysis was performed with OLIGOMER.

Isothermal titration calorimetry

Isothermal titration calorimetry (ITC) assays were performed on a VP-ITC (MicroCal Inc., Northampton, MA). Titrations consisted of an initial injection (1 μL), followed by 25–30 injections of 10 μL of DesR-REC (ligand) on the cell containing the different DesKC variants (partner). Assays were carried out at 15°C in a buffer containing 20 mM Tris.HCl pH 8, 0.3 M NaCl, 10 mM MgCl_2 and 0.5 mM AMP-PNP. The heat released by the dilution of the ligand was determined injecting the ligand, on the cell containing the working solution without partner, using the same sequence of injections. The concentrations used for the experiments were between 25–30 μM for DesKC and 350–400 μM for DesR-REC and each titration was done in duplicate. The data were analyzed with MicroCal Origin version 7 software (MicroCal Software Inc.), after manual baseline correction, and subtraction of heat due to ligand dilution. Binding isotherms were fitted to a two independent sequential site model.

Sequence-based direct coupling analysis of the HK:RR complex

Sequences of HK and RR pairs, with the same architecture as DesK and DesR (families HisKA₃ and NarL, respectively), belonging to the same operon, were selected from the Uniprot database using hmmssearch (Eddy, 1998). Hidden Markov model-profiles for each domain were obtained from Pfam (Finn et al., 2016). Redundancy was filtered using a cutoff of 90% (Li and Godzik, 2006) resulting in a total of 3318 sequences including HKs (DHP+CA) and RRs (REC+HTH). The concatenated HK:RR sequences were aligned with hmalign (Eddy, 1998) and manually curated, removing sequences with large insertions. DI (direct information) of each pair of residues of the alignment was calculated using mfDCA matlab script (Morcos et al., 2011) and we only take in consideration coevolving pairs of residues farther than five positions apart in sequence. The same procedure was followed for the HisKA and PhoB families for comparison purposes. A similar trend in equivalent 3D positions for covariant residue pairs among different HK and RR families was thus confirmed (Podgornaia et al., 2013; Skerker et al., 2008; Weigt et al., 2009).

In silico modeling and molecular dynamics simulations

The phosphotransferase complex was built by replacing the DesK_{H188E} dimer of 5IUK with wt P-DesKC (PDB 5IUM). This replacement resulted in model with barely no clashes, except for P-His_{188(HK)} slightly bumping into Phe_{82(RR)} and Arg_{84(RR)}. Energy minimization was then performed on such constructed complex with all-atom constraints in Rosetta (Das and Baker, 2008), strongly restraining shifts to maintain the experimental coordinates. A final wt P-DesKC:DesR-REC model was thus obtained, with optimal stereochemical geometry and no clashes (0.09 rmsd between the energy minimized model and the pdb 5IUM experimental structure, aligning all 1930 DesK atoms). To further evaluate the minimized wt P-DesKC:DesR-REC model (see Figure 4A), molecular dynamics simulations were performed, observing that its conformation is stable throughout the trajectory (see Figure 4B), especially considering the positions of His_{188(HK)}, Asp_{54(RR)} and Thr_{80(RR)} in contact with the phosphate.

For molecular dynamics (MD) calculations, the missing ATP lid loop within the CA domain in chain A (residues 328 to 335) were reconstructed by using a previously reported high resolution structure (PDB 3EHG) as template. The other CA domain (in chain B, residues 241–367) was deleted. The missing N-terminal portion of chain B helix α 1, was reconstructed and extended up to residue Lys_{155(HK)} using 5IUK as a template.

For computational efficiency only DesK residues Lys_{155(HK)} to Asn_{368(HK)} in chain A and Lys_{155(HK)} to Ser_{239(HK)} in chain B were considered. Only His_{188(HK)} in chain B was set to be phosphorylated. His_{335(HK)} was protonated at position N δ 1, while other histidines were protonated at N ϵ 2, to preserve the interaction network. The ATP moiety and Mg²⁺ cation present within the CA domain were both kept in the model. The structure of DesR-REC spanned residues Ser_{0(RR)} to Leu_{131(RR)}. The Mg²⁺ cation in the active site of DesR-REC and the three crystallographic water molecules of the metal's coordination sphere, were included in the model.

Acetyl and N-methylamide capping groups were added to N- and C- terminal residues, which did not correspond to the real terminals in the full protein sequences. The protein complex was solvated within an octahedral box of 15 Å from the solute, and sodium counterions were added to preserve the electroneutrality of the system. The AMBER force field ff14SB (Maier et al., 2015) was used to represent the amino acids. The parameters for His_{188(HK)} phosphorylated at NE2 with an unprotonated phosphate group were taken from (Homeyer et al., 2006). The ATP molecule was described with reported parameters (Meagher et al., 2003). The TIP3P model (Jorgensen et al., 1983) was used for water molecules. Monovalent ions were treated with previously described parameters (Joung and Cheatham, 2008), while Mg²⁺ ions were modeled using the compromise set of parameters for the 12–6 non-bonded potential in TIP3P (Li et al., 2013).

All calculations were performed with the GPU version of AMBER14 (Salomon-Ferrer et al., 2013). Initially, the whole system was relaxed by energy minimization, then it was equilibrated for 0.2 ns in the NVT ensemble imposing harmonic positional restrains of 10 kcal mol⁻¹ Å⁻² on the protein atoms. A reference temperature of 300 K was set by coupling the system to the Langevin thermostat (Pastor et al., 1988; Wu and Brooks, 2003) with a friction constant of 50 ps⁻¹, which approximates the physical collision frequency for liquid water (Izaguirre et al., 2001). A 10 Å cut-off was used for non-bonded interactions, while long-range electrostatics were evaluated using Particle

Mesh Ewald (PME) (Darden *et al.*, 1993; Essmann *et al.*, 1995). A time step of 2 fs was used and all bonds involving hydrogen atoms were restrained using the SHAKE algorithm (Miyamoto and Kollman, 1992; Ryckaert *et al.*, 1977). Production simulations were performed in the NPT ensemble. The pressure was kept at 1 atm by means of the Berendsen barostat (Berendsen *et al.*, 1984). Snapshots were recorded every 5 ps for analysis.

Autodephosphorylation and phosphotransfer assays

To purify the phosphorylated species of DesR-REC and DesR-REC_{F82A}, 600 μ M of both recombinant proteins were auto-phosphorylated using 50 mM acetyl phosphate in a buffer containing 25 mM Tris.HCl pH 8, 300 mM NaCl and 30 mM MgCl₂, at room temperature. Reactions were stopped by adding EDTA to a final concentration of 50 mM, and buffer exchanged by using a PD Minitrap G-25 (GE Healthcare) desalting column. Auto-dephosphorylation assays of P-DesR-REC variants were performed at 30 μ M of protein concentration and incubated in the presence of 30 mM MgCl₂. At different time points the reactions were stopped by adding SDS-PAGE sample buffer. For each time point DTT was added to a final concentration of 25 mM and incubated for 5 min. DTT in excess was blocked with 40 mM iodo-acetamide and loaded in a Phos-tag acrylamide SDS-PAGE, as described before (Trajtenberg *et al.*, 2014), Coomassie blue-stained gels were scanned and quantification of all reactions was done by densitometry using ImageJ (Schneider *et al.*, 2012).

To perform phosphotransfer assays (Figure 4E), DesKC was phosphorylated by incubation with 10 mM ATP and 20 mM MgCl₂ at 100 μ M of protein, for 1 hr at room temperature. P-DesKC was then purified by size exclusion chromatography S75 10/300 (GE Healthcare), equilibrated in 20 mM Tris.HCl pH 8, 300 mM NaCl. Phosphotransfer reactions were performed in triplicate by combining 30 μ M P-DesKC with equimolar concentrations of DesR-REC (both wild type and DesR-REC_{F82A}), at 25°C in 20 mM Tris.HCl pH 8, 300 mM NaCl, 30 mM MgCl₂. Reactions were stopped at different time points as described above. Samples were then separated by Phos-tag SDS-PAGE.

In order to assess unidirectionality (Figure 5A), the reverse and forward reactions were analyzed. The forward reaction was prepared as described above using P-DesKC as phosphodonor for 1 min. On the other hand, the reverse of the reaction was measured by incubating 30 μ M of P-DesR-REC (wild type or point mutants DesR-REC_{F82A} or DesR-REC_{R84A}) with 30 μ M DesKC, 50 mM acetyl phosphate and 30 mM MgCl₂, for 1 min at room temperature. Reactions were stopped as described above and samples analyzed by Phos-tag SDS-PAGE and quantified by densitometry.

Accession numbers

The X ray structures presented have been deposited in the wwPDB with accession codes 5IUJ (DesK-DesR complex in the phosphotransfer state with low Mg²⁺ [20 mM]), 5IUK (DesK-DesR complex in the phosphotransfer state with high Mg²⁺ [150 mM]), 5IUL (DesK-DesR complex in the phosphotransfer state with high Mg²⁺ [150 mM] and BeF₃⁻), 5IUM (phosphorylated wild type DesKC) and 5IUN (DesK-DesR complex in the phosphatase state).

Raw X ray diffraction data corresponding to each one of these structures are publicly available at SBCGrid Data Bank (<http://data.sbgrid.org>) as dataset entries 399, 400, 401, 407 and 408.

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Author contributions

FT, AB, Conception and design, Acquisition of data, Analysis and interpretation of data, Drafting or revising the article; JAI, GO, Acquisition of data, Analysis and interpretation of data; MRM, Acquisition of data, Analysis and interpretation of data, Contributed unpublished essential data or reagents; NL, Acquisition of data, Contributed unpublished essential data or reagents; MAM, Analysis and interpretation of data, Drafting or revising the article, Contributed unpublished essential data or reagents; AEM, Acquisition of data, Analysis and interpretation of data, Drafting or revising the article

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Additional files

Major datasets

The following datasets were generated:

Author(s)	Year	Dataset title	Dataset URL	Database, license, and accessibility information
Trajtenberg F, Buschiazzo A	2016	Crystal structure of phosphorylated DesKC	http://www.rcsb.org/pdb/search/structid-Search.do?structureId=5IUM	Publicly available at the RCSB Protein Data Bank (accession no: 5IUM)
Trajtenberg F, Imelio JA, Larrieux N, Buschiazzo A	2016	Crystal structure of the DesK-DesR complex in the phosphatase state	http://www.rcsb.org/pdb/search/structid-Search.do?structureId=5IUN	Publicly available at the RCSB Protein Data Bank (accession no: 5IUN)
Trajtenberg F, Imelio JA, Larrieux N, Buschiazzo A	2016	Crystal structure of the DesK-DesR complex in the phosphotransfer state with low Mg ²⁺ (20 mM)	http://www.rcsb.org/pdb/search/structid-Search.do?structureId=5IUJ	Publicly available at the RCSB Protein Data Bank (accession no: 5IUJ)
Trajtenberg F, Imelio JA, Larrieux N, Buschiazzo A	2016	Crystal structure of the DesK-DesR complex in the phosphotransfer state with high Mg ²⁺ (150 mM)	http://www.rcsb.org/pdb/search/structid-Search.do?structureId=5IUK	Publicly available at the RCSB Protein Data Bank (accession no: 5IUK)
Trajtenberg F, Imelio JA, Larrieux N, Buschiazzo A	2016	Crystal structure of the DesK-DesR complex in the phosphotransfer state with high Mg ²⁺ (150 mM) and BeF ₃	http://www.rcsb.org/pdb/search/structid-Search.do?structureId=5IUL	Publicly available at the RCSB Protein Data Bank (accession no: 5IUL)
Trajtenberg F,	2016	Complex DesKC:DesR-REC (B.	http://dx.doi.org/10.	Publicly available at

Buschiazio A		subtilis) - phosphotransferase state, low Mg ²⁺	15785/SBGRID/399	the SBGrid database (dataset no. 399; X-ray diffraction raw data for PDB 51UJ)
Trajtenberg F, Buschiazio A	2016	Complex DesKC:DesR-REC (B. subtilis) - phosphatase state	http://dx.doi.org/10.15785/SBGRID/400	Publicly available at the SBGrid database (dataset no. 400; X-ray diffraction raw data for PDB 51UN)
Trajtenberg F, Buschiazio A	2016	Complex DesKC:DesR-REC (B. subtilis) - phosphotransferase state, high Mg ²⁺	http://dx.doi.org/10.15785/SBGRID/401	Publicly available at the SBGrid database (dataset no. 401; X-ray diffraction raw data for PDB 51UK)
Buschiazio A	2016	Phosphorylated DesKC (B. subtilis)	http://dx.doi.org/10.15785/SBGRID/407	Publicly available at the SBGrid database (dataset no. 407; X-ray diffraction raw data for PDB 51UM)
Trajtenberg F, Buschiazio A	2016	Complex DesKC:DesR-REC (B. subtilis) - phosphotransferase state, high Mg ²⁺ and Bef3-	http://dx.doi.org/10.15785/SBGRID/408	Publicly available at the SBGrid database (dataset no. 408; X-ray diffraction raw data for PDB 51UL)

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