

A conserved protonation-dependent switch controls drug binding in the Abl kinase

Yibing Shan^a, Markus A. Seeliger^b, Michael P. Eastwood^a, Filip Frank^b, Huafeng Xu^a, Morten Ø. Jensen^a, Ron O. Dror^a, John Kuriyan^{b,1}, and David E. Shaw^{a,c,1}

^aD. E. Shaw Research, New York, NY 10036; ^bHoward Hughes Medical Institute, Department of Molecular and Cell Biology and Department of Chemistry, University of California, Berkeley, CA 94720; and ^cCenter for Computational Biology and Bioinformatics, Columbia University, New York, NY 10032

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In many protein kinases, a characteristic conformational change (the “DFG flip”) connects catalytically active and inactive conformations. Many kinase inhibitors—including the cancer drug imatinib—selectively target a specific DFG conformation, but the function and mechanism of the flip remain unclear. Using long molecular dynamics simulations of the Abl kinase, we visualized the DFG flip in atomic-level detail and formulated an energetic model predicting that protonation of the DFG aspartate controls the flip. Consistent with our model’s predictions, we demonstrated experimentally that the kinetics of imatinib binding to Abl kinase have a pH dependence that disappears when the DFG aspartate is mutated. Our model suggests a possible explanation for the high degree of conservation of the DFG motif: that the flip, modulated by electrostatic changes inherent to the catalytic cycle, allows the kinase to access flexible conformations facilitating nucleotide binding and release.

conformational change | DFG motif | imatinib | molecular dynamics simulation | pH dependence

Nearly 2% of human genes encode protein kinases (1), enzymes involved in almost all aspects of cellular regulation, which use ATP to phosphorylate protein substrates and release ADP as a byproduct. Given the importance of protein kinases to cell growth, movement, and death, it is not surprising that the protein kinase catalytic domain is the protein domain most commonly encoded by known cancer genes (2), and that protein kinases constitute one of the largest families of drug targets (3).

Lying within nearly every eukaryotic protein kinase domain, adjacent to the ATP-binding site, is a sequence of 3 residues, Asp-Phe-Gly (DFG). The reasons for the conservation of these residues are not fully understood, but the motif is known to be highly important, as was recently highlighted by the identification of potential “driver” mutations in human cancers at the DFG motif in 8 different kinases (4). The DFG aspartate is believed to be important for catalysis, and indeed typically points into the ATP-binding site (the DFG-in conformation), where it can coordinate an ATP-bound magnesium ion (5, 6). In a diverse subset of protein kinases, however, including c-Abl, IRK, c-Kit, Flt3, Csk, RAF, and P38, the DFG motif can also adopt a dramatically different conformation in which the DFG aspartate and phenylalanine side chains swap positions, leaving the aspartate pointing away from the binding site (the DFG-out conformation). The conformational change connecting the DFG-in and DFG-out conformations (a “DFG flip”; see Fig. 1) involves a ≈ 10 -Å change in the positions of the 2 side chains.

The DFG flip (see [Movie S1](#)) is a central feature of the activation of the nonreceptor tyrosine kinase c-Abl, which before activation is held in an inactive DFG-out conformation by an autoinhibitory interaction with its regulatory domains (7). The DFG flip underlies the success of the cancer drug imatinib, which binds to the DFG-out conformation of the aberrant gene fusion product BCR-Abl (8) and has proven remarkably efficacious in the clinical treatment of chronic myeloid leukemia and certain other cancers (9, 10). The ability of imatinib to specifically recognize the DFG-out conformation of BCR-Abl has been credited for imatinib’s selectivity, and

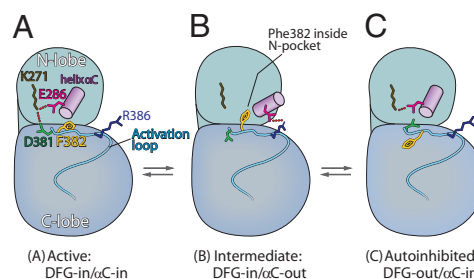


Fig. 1. A proposed mechanism for the DFG flip based on crystal structures and our Abl simulations. Key structural features are highlighted, with salt bridges represented by red dotted lines. The ATP-binding site is located between the N-lobe and the C-lobe. The active structure (A) and the autoinhibited structure (C) differ in their DFG conformation, and interconvert via a DFG flip. Nevertheless, these structures both adopt conformations referred to as α C-in, in which helix α C is close to the ATP-binding site so that the Lys-271–Glu-286 salt bridge may form. (B) The structure shown is taken from the simulations, and is a proposed intermediate in the DFG flip. This structure differs from A and C in 2 significant ways. First, helix α C is positioned away from the ATP-binding site so that Glu-286 may form a salt bridge with Arg-386 (an α C-out conformation). Second, the outward displacement of helix α C creates a pocket at the base of the N-lobe, which we refer to as the N-pocket, and which in B is occupied by Phe-382. The PDB ID codes of the structures on which the figure is based are 2F4J (A) and 1OPK (C). (1OPK contains regulatory domains, but only the kinase domain is shown here.)

intensive searches are now under way for other kinase inhibitors that selectively bind to DFG-out conformations (11, 12).

Knowledge of the detailed mechanism and driving forces of DFG conformational change would deepen our understanding of kinase activation and open new directions in drug discovery. However, it is very challenging to directly probe the dynamics of this motif experimentally (13). Molecular dynamics (MD) simulation provides an alternative means to study kinase dynamics (14–16), but the time scale accessible to such studies has typically been restricted to between 100 ps and 10 ns. Observation of the DFG flip at such short time scales has required the introduction of artificial biasing forces (17), which can generate nonphysical trajectories and obscure the forces that actually drive the flip. In the work reported here, we used recently developed algorithms for the high-speed, parallel execution of MD simulations (18–21) to perform unbiased MD simulations, each hundreds of nanoseconds in length, of a construct

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¹To whom correspondence may be addressed. E-mail: David.Shaw@DEShawResearch.com or kuriyan@berkeley.edu.

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$$P_{\text{out}} = \frac{1}{1 + \exp(\Delta G_{2 \rightarrow 3}/RT) + \exp(\Delta G_{2 \rightarrow 3}/RT + [\text{pH} - \text{p}K_{\text{a, in}}] \times \log_e 10)}, \quad [1]$$

where $\Delta G_{2 \rightarrow 3}$ is the free-energy cost of the DFG flip from DFG-in to DFG-out with Asp-381 protonated, R is the gas constant, and $\text{p}K_{\text{a, in}}$ is the $\text{p}K_{\text{a}}$ of Asp-381 in a DFG-in conformation (i.e., it is the hypothetical $\text{p}K_{\text{a}}$ when DFG-out states are excluded, and thus differs from the actual $\text{p}K_{\text{a}}$).

At equilibrium, one might anticipate DFG-in conformations to dominate, because the aspartate and phenylalanine residues are favorably placed in environments matching their polarity. Three other factors combined, however, may tip the thermodynamic balance in favor of DFG-out conformations. First, in the DFG-out conformation of Abl, which we model on the kinase domain of c-Abl PDB ID code 1OPK (7), Asp-381 is well positioned to form a hydrogen bond with the exposed backbone carbonyl of Val-299. Second, contrary to what one might expect given the preponderance of reported kinase structures that appear in the DFG-in conformation, we find that the ϕ - ψ angles of the DFG aspartate in active Abl, and in most other DFG-in kinase structures, fall within the unusual left-handed helix region of Ramachandran space (Fig. S4B), indicating a more strained main-chain configuration for DFG-in than DFG-out. Third, standard electrostatics calculations (30) suggest that Asp-381 has an elevated value of $\text{p}K_{\text{a, in}}$ in the absence of the ATP-bound magnesium ion, due in part to the influence of the conserved residues Glu-286, Asp-363, and Asn-368. Analysis of kinase crystal structures as a function of crystallization buffer pH (Fig. S4C) also suggests that an elevated $\text{p}K_{\text{a, in}}$ may occur in other kinases.

Combining standard literature estimates (35) and simulation results (see *SI Text*), we have estimated the contributions to $\Delta G_{2 \rightarrow 3}$ and we find $\Delta G_{2 \rightarrow 3} \approx -1.1$ kcal/mol. The electrostatics calculations yield $\text{p}K_{\text{a, in}} \approx 6.6$. With these estimates, Eq. 1 gives a large relative population of DFG-out conformations at $\text{pH} \approx 6$ that decreases rapidly around $\text{pH} \approx 7$ (Fig. 4B). Although the estimated parameters are, of course, only semiquantitative, the basic conclusion of a near-balance in the free energies of DFG-in and DFG-out conformations close to physiological pH is robust to moderate changes in the parameters. Importantly, Eq. 1 shows that with a small $\Delta G_{2 \rightarrow 3}$, the DFG conformation becomes sensitive to changes in $\text{p}K_{\text{a, in}}$ and can thus be controlled by electrostatic changes in the ATP-binding site (we discuss the potential significance of this observation to kinase catalysis below). Similarly, our model predicts the effect of mutating Asp-381 or Phe-382 on DFG conformation; replacing Phe-382 with a less hydrophobic residue, for example, increases the relative population of DFG-out (Fig. 4B).

Our energetic model should also be applicable to other kinases, most directly if their activation loops adopt the common open conformation (22, 36) in DFG-in and DFG-out states (as do Abl and c-Abl in PDB ID codes 2F4J (22) and 1OPK (7), respectively). Other activation-loop conformations—particularly those occluding substrate binding, as in Abl–imatinib complexes such as PDB ID code 1IEP (8)—can lead to solvent exposure of both Asp-381 and Phe-382. Such structures are found to adopt DFG-out conformations. This observation is consistent with our analysis, because the key factor stabilizing DFG-in conformations (the hydrophobic packing of Phe-382) is missing from these structures.

Probing the DFG-out Conformation by Using Abl–Imatinib Binding.

With imatinib bound, Abl adopts a DFG-out conformation (8), suggesting that imatinib binding kinetics may provide a probe of DFG conformation. We derived the on-rate constant, k_{on} , of imatinib binding to Abl as a function of pH by using stopped-flow fluorescence assays. As shown in Fig. 4A, k_{on} for wild-type Abl decreases by nearly an order of magnitude as pH increases from 5.5 to 7.5, in which range imatinib is predominantly neutral (37). We

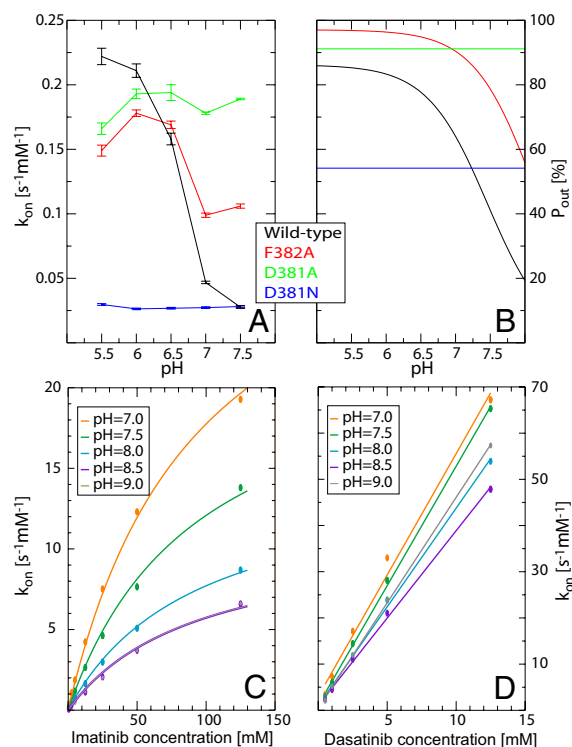


Fig. 4. pH dependence of Abl–imatinib binding. (A) The on-rate constant of imatinib binding to wild-type Abl and to 3 mutants as a function of pH, as measured by stopped-flow tryptophan fluorescence assays. The error bars indicate the uncertainty of the linear regression fit to the observed rates from which the on-rate constants are obtained. (B) The relative population of DFG-out estimated according to Eq. 1. The similarity between the model results for P_{out} and the experimental k_{on} values supports our interpretation of the experimental data and the factors underlying DFG flips. (C) The observed fluorescence-decay rates for Abl–imatinib binding rates at 5 different pH values (virtually identical at pH 8.5 and pH 9). The solid lines are obtained from independent 2-parameter fits using the proposed kinetic scheme (see *SI Text* for details). (D) The dasatinib binding rate constants for wild-type Abl measured at 5 different pH values show only weak pH dependence. The lines are linear fits to the data.

observed a very similar pH dependence for a c-Abl construct containing its kinase and SH2-SH3 domains (Fig. S2A). However, control experiments showed the imatinib-binding kinetics for Abl mutants D381A and D381N to have essentially no dependence on pH over the same range (Fig. 4A). The results for the mutants strongly suggest that the pH dependence observed for the wild type was a result of protonation at Asp-381.

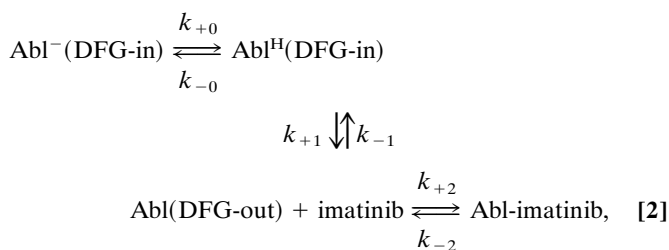
Independent structural evidence supporting protonation of Asp-381 also exists. Most convincingly, the DFG-out c-Abl structure (PDB ID code 1OPK, resolution 1.8 Å, crystallized at pH 7.0) (7) shows 1 Asp-381 side-chain oxygen to be only 2.8 Å away from the Val-299 backbone carbonyl oxygen in a position that would be hard to rationalize if Asp-381 were not protonated in this structure. In addition, our analysis of the representative kinase structures (restricted to DFG-in conformations without bound ligands) shows that the distance of the DFG aspartate from the conserved lysine (Lys-271 in Abl) generally increases with decreasing pH of the crystallization buffer (Fig. S4C). This suggests that protonation of the DFG aspartate may be common in protein kinases.

The above results provide strong evidence that the $\text{p}K_{\text{a}}$ of Asp-381 is significantly raised above typical values for aspartate residues. Because imatinib is a DFG-out binder (8), another natural conclusion from the data is that protonation of Asp-381 favors DFG-out conformations. As a negative control, we have also performed fluorescence experiments using the DFG-in binder

dasatinib (38, 39). It is thought that so-called DFG-in binders typically bind to both DFG-in and DFG-out conformations (13). We find only a very weak pH dependence for dasatinib binding (Fig. 4D), consistent with the interpretation that the pH dependence observed for imatinib binding is a result of a pH-dependent DFG conformation.

More evidence supporting a link between protonation and DFG conformation comes from the pH dependence of imatinib binding to the Abl mutant F382A shown in Fig. 4A. This mutant shows a decrease in the imatinib on-rate constant with increasing pH, although the decrease is less pronounced and shifted to higher pH compared with the wild type. This implies a higher pK_a for Asp-381 in the mutant, despite the fact that both phenylalanine and alanine are nonpolar and are positioned away from Asp-381, indicating that the mutation causes a conformational change and hence the elevated pK_a . As explained above, an F382A mutation is predicted to lead to a larger population of DFG-out conformations, thus raising the pK_a of Asp-381 in agreement with our experimental observations.

Kinetic Scheme for Imatinib Binding. In contrast to its kinetics, the thermodynamics of Abl–imatinib binding determined from isothermal titration calorimetry (ITC) do not show a clear pH dependence (see Table S2). This is consistent with the observation made above that in the Abl–imatinib complex—unlike in the apo DFG-out conformation—Asp-381 is solvent-exposed (because of rearrangement of the activation loop on binding; see Fig. S5A). To provide a quantitative explanation for the pH-dependent imatinib-binding kinetics, we propose the following kinetic scheme:



where Abl^{H} and Abl^- denote Abl with and without protonation of Asp-381. For the imatinib concentrations ($<12.5 \mu\text{M}$) used in our experiments (discussed above), the fluorescence relaxation data fits a single-exponential decay, and the resultant relaxation rates are found to increase linearly with imatinib concentration (Fig. S6). Under these conditions, we thus assume that the Asp-381-protonation and DFG-flip steps of the kinetic scheme (the leftmost and middle steps in Eq. 2, respectively) are fast compared with imatinib binding to the DFG-out conformation (rightmost step in Eq. 2). This leads to effective 1-step kinetics with an on-rate constant proportional to the population of the DFG-out state: $k_{\text{on}} = P_{\text{out}}k_{+2}$. In Fig. 4B, we show our estimates for P_{out} (Eq. 1) as a function of pH for Abl and 3 mutants. These estimates are clearly consistent with our experimental results for k_{on} , supporting our interpretation that the protonation of Asp-381 increases the population of DFG-out.

At sufficiently high imatinib concentrations, the rate of imatinib binding to the DFG-out conformation should become comparable with that of the DFG flip. In this case, the kinetic scheme leads to effective 2-step kinetics. This is consistent with the observed fluorescence decay rates, k_{obs} , which are found to deviate from a linear dependence on imatinib concentration (see Fig. 4C). In contrast, similar experiments for dasatinib binding showed no clear deviation from linearity (Fig. 4D). These results clearly demonstrate an intermediate step in imatinib binding, and support the interpretation that the pH dependence of imatinib binding arises from the DFG flip. Detailed analysis of the kinetic data of imatinib binding also revealed the DFG-flip time scale to be in the tens of millise-

conds (see *SI Text*), which is consistent with NMR studies of DFG dynamics (13). This relatively long time scale may reflect the fact that the DFG flip involves conformational changes of the activation loop beyond the DFG motif itself (33).

Conservation of the DFG Motif and Its Role in Kinase Catalysis.

Although the DFG aspartate is clearly important for catalysis, the roles of the phenylalanine and glycine residues and the reason for their conservation are much less certain (40). Our results provide evidence that both of these residues may be essential to DFG conformational change. We find that the phenylalanine plays a key role in maintaining an energetic balance between the DFG-in and DFG-out states by compensating for the unfavorable DFG aspartate backbone torsion angles adopted in the DFG-in conformation. Our simulations also show that the DFG flip involves extensive backbone-torsion motions at the glycine residue, suggesting that the unique flexibility of glycine may serve to lower the kinetic barrier for the flip. This is consistent with work (41) suggesting that the DFG motif is optimized for functional dynamics rather than for structural stability, as is our finding that the DFG-out conformation can be populated under physiological conditions even in the absence of DFG-out binders.

If the DFG flip is to explain the conservation of the DFG motif, its role in kinase function must extend beyond the specific activation mechanism of c-Abl. The DFG flip has been suggested to promote ADP release (27), the rate-limiting step in kinase catalysis (42–44). Intriguingly, our results could be interpreted as hinting at a mechanistic explanation for such a proposed role. In ATP-bound kinases, the DFG aspartate interacts with a catalytically essential magnesium ion (Mg^{2+}) 2.3 Å away, which in turn coordinates the β and γ phosphates of ATP (5). Encouraged by observations on small G proteins (45, 46) showing that Mg^{2+} binding is weakened by ≈ 3 orders of magnitude after transfer of the γ phosphate, we speculate that in kinases, this Mg^{2+} may likewise be released after phosphate transfer, thus reducing the positive charge in the immediate vicinity of the DFG aspartate. If this is the case, we predict that this change in the electrostatic environment of the DFG aspartate would lead to its protonation—either from the solvent or through proton transfer from the catalytic base Asp-363—causing the motif to flip to a DFG-out conformation and thus facilitating ADP release. In such a scenario, nucleotide release could, in principle, be regulated by magnesium, consistent with the experimental observation that the ADP-release rate decreases with increasing Mg^{2+} concentration (44).

The DFG flip could facilitate ADP release, and potentially binding of the ATP- Mg^{2+} complex, by allowing the kinase to adopt a DFG-out conformation, which we find to be significantly more flexible in terms of interlobe motions than the DFG-in conformation (Fig. S7). This increased flexibility is consistent with the finding (32) that, in the active conformation, the DFG phenylalanine is part of the “hydrophobic spine” that lends stability to the DFG-in conformation. Although speculative, our suggestions are also consistent with kinetic data showing that at physiological Mg^{2+} concentration, a 10-ms time scale conformational change follows ATP binding and precedes ADP release (44), as well as with recent NMR titration experiments showing that upon the binding of ATP to PKA kinase, the DFG motif and nucleotide-binding loop of PKA exhibit microsecond to millisecond dynamics (31).

Conclusion

Using Abl as a model system, we have simulated the DFG flip, a conformational change known to be significant in a diverse set of kinases and, in particular, to be an important switch for c-Abl activation (7). The simulations highlight the role of large-scale conformational changes in facilitating the DFG flip and point to the importance of the protonation of the DFG aspartate residue. This conclusion is supported experimentally by our Abl–imatinib binding assays, which reveal markedly pH-dependent binding

kinetics. Our analysis shows that the DFG conformation is determined by a number of opposing factors that approximately balance, resulting in a DFG conformation sensitive to small changes in the electrostatics of the ATP-binding site. Combined with existing structural and kinetic data, this leads us to suggest that the conservation of the DFG motif stems from the DFG flip, which switches the kinase from the less-flexible DFG-in form required for catalysis to the more-flexible DFG-out form that facilitates nucleotide binding and release. The set of intermediate conformations observed in our simulations of the DFG flip may present new opportunities for inhibitor design. Furthermore, the pH sensitivity of the DFG conformation raises the possibility that local pH variations may play a role in kinase regulation in vivo.

Materials and Methods

Additional details are provided in [SI Text](#).

Simulation Details. All simulations used the simple point charge (SPC) model for water (47), the standard OPLS-AA parameters for ions (48, 49), and the

OPLS-AA/L force field for proteins (28, 29) as provided by the program IMPACT (50) (note that the relevant force-field option, referred to as OPLS.2003 in the IMPACT paper, has subsequently been renamed as OPLS.2005 in the actual program). All molecular dynamics simulations were performed by using the parallel MD program Desmond (19) on 64 or 128 dual-processor Opteron nodes connected by a high-speed Infiniband network (Topspin, San Jose, CA). Each simulation is represented by a code number that is given, along with other key simulation details, in [Table S1](#).

Kinetic Measurements of Drug Binding. Wild-type human c-Abl kinase domain (residues 248–532) and mutant proteins were generated and purified as described previously (51). The drug-binding kinetics is measured by monitoring the decrease of protein fluorescence at 350 nm upon excitation at 290 nm on a HORIBA Jobin Yvon FluoroMax-3 spectrofluorimeter for 10–20 half-lives of the transient, recorded by 1,000–2,000 data points.

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