

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

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Contributed by John Kuriyan, November 13, 2008 (sent for review September 29, 2008)

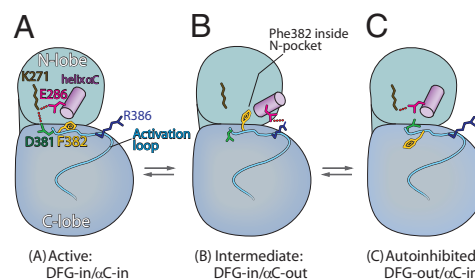
In many protein kinases, a characteristic conformational change (the “DFG flip”) connects catalytically active and inactive conformations. Many kinase inhibitors—including the cancer drug imatinib—selectively target a specific DFG conformation, but the function and mechanism of the flip remain unclear. 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  $\approx 10$ -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  $\alpha$ C-in, in which helix  $\alpha$ 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  $\alpha$ C is positioned away from the ATP-binding site so that Glu-286 may form a salt bridge with Arg-386 (an  $\alpha$ C-out conformation). Second, the outward displacement of helix  $\alpha$ 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

Author contributions: Y.S., J.K., and D.E.S. designed research; Y.S., M.A.S., and F.F. performed research; Y.S., M.A.S., M.P.E., and H.X. analyzed data; and Y.S., M.A.S., M.P.E., M.O.J., R.O.D., J.K., and D.E.S. wrote the paper.

The authors declare no conflict of interest.

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This article contains supporting information online at [www.pnas.org/cgi/content/full/0811223106/DCSupplemental](http://www.pnas.org/cgi/content/full/0811223106/DCSupplemental).

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of human c-Abl containing only the kinase domain. (Because the kinase domains of c-Abl and BCR-Abl are identical in sequence, we refer to this common domain below as simply “Abl.”) Our analysis of these simulations allowed us to formulate a model of key dynamic, energetic, and structural factors involved in DFG flips. To validate our model experimentally, we used fluorescence assays of Abl–imatinib binding to probe DFG conformation, the results of which confirmed our central computational findings.

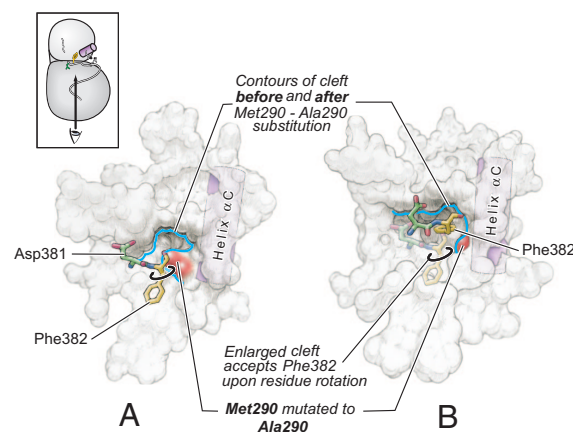
Like other eukaryotic protein kinase domains, Abl consists of a smaller N-terminal lobe and a larger C-terminal lobe (the N-lobe and the C-lobe, respectively), with the ATP-binding site located between them. The N-lobe is composed of 5  $\beta$ -strands and a helix referred to as  $\alpha$ C, whereas the C-lobe is predominantly helical. The DFG motif borders the ATP-binding site and is located between the 2 lobes, at the N terminus of a flexible segment known as the activation loop. Fig. 1*A* and *C* shows Abl adopting DFG-in (22) and DFG-out (7) conformations, respectively, whereas Fig. 1*B* shows an important intermediate conformation observed in our simulations of the DFG flip. This conformation differs from the other 2 in that helix  $\alpha$ C is displaced away from the binding site (the  $\alpha$ C-out conformation, by contrast with the  $\alpha$ C-in conformation shown in Fig. 1*A* and *C*). A recently solved crystal structure of Abl also adopts an  $\alpha$ C-out conformation, which has been suggested as a potential intermediate in the DFG flip (17). Although kinases in their active states adopt  $\alpha$ C-in conformations, many kinases in their inactive states are known to adopt  $\alpha$ C-out conformations, first defined in Cdk (23) and Src (24, 25) kinases, but now seen in many others—including EGFR (26), whose  $\alpha$ C-out conformation is recognized by the cancer drug lapatinib. Thus, just as many kinases can access alternative conformations through a DFG flip, many kinases can access alternative conformations through a transition between  $\alpha$ C-in and  $\alpha$ C-out conformations.

To study the conformational changes associated with the DFG flip without introducing any biasing forces in our MD simulations, we “loosened” helix  $\alpha$ C by mutating to alanine a methionine residue that previous studies suggested might be involved in stabilizing hydrophobic interactions between helix  $\alpha$ C and the remainder of the protein. In our simulations of the resulting protein, we were able to observe the DFG flip and its coupling with helix- $\alpha$ C motion at an atomic level of detail. Analysis of these simulations led us to propose an energetic model in which the protonation state of Asp-381, which is influenced by local electrostatic interactions, serves as the switch that controls the flip. On the basis of our simulation results, we made certain predictions regarding the pH dependence of Abl–imatinib binding, then performed fluorescence quenching experiments to test these predictions. The results of these experiments were consistent with our model (and also allowed us to estimate the time scale of the DFG flip to be tens of milliseconds). Our findings could be interpreted as providing mechanistic support for the suggestion (27) that the DFG flip plays a significant role in the catalytic cycle of protein kinases by facilitating nucleotide binding and release.

## Results and Discussion

**Simulations of Abl.** We performed a total of  $>2 \mu$ s of all-atom MD simulations of Abl in explicit water, using the OPLS-AA/L force field (28, 29) for proteins, with most individual simulations between 100 and 300 ns in length. All simulations were performed at constant temperature (300 K) and pressure (1 bar); see *SI Text*.

**N-Pocket as Part of the DFG-Flip Pathway.** Standard electrostatics calculations (30) (see *SI Text*) suggest that in a DFG-in structure of Abl [PDB ID code 2F4J; Young *et al.* (22)], the DFG aspartate has an elevated  $pK_a$ . In a DFG-out structure of Abl [PDB ID code 1OPK (7)] the DFG aspartate appears to be protonated and hydrogen bonded to a backbone carbonyl group. These observations led us to hypothesize that the protonation of the DFG aspartate might be an important determinant of DFG conforma-



**Fig. 2.** The DFG motif and the N-pocket. Both *A* and *B* show the base of the N-lobe, with the C-lobe removed to make the N-pocket and DFG motif fully visible (see *Inset*). (*A*) An  $\alpha$ C-out conformation illustrating the N-pocket. In this conformation the close interactions between Phe-382 and Met-290 have been transiently broken, but Met-290 still presents a steric barrier to the entry of Phe-382 into the N-pocket. This structure was taken from simulation 1a after 250 ns. (*B*) Phe-382 occupying the N-pocket, as enlarged by the M290A mutation. The motion through which Phe-382 enters the N-pocket (to yield a structure like that shown in Fig. 1*B*) principally involves a change of its  $\chi_1$  angle from approximately  $-60^\circ$  to  $60^\circ$ . This structure is taken from simulation 4a after 2 ns. The conformation of Asp-381 and Phe-382 before Phe-382 enters the N-pocket is shown in a transparent rendering.

tion, and we therefore performed simulations with the residue in different protonation states. We started 4 simulations in the active DFG-in conformation, including 2 (simulations 1a and 1b in *Table S1*) with Asp-381 protonated and 2 with Asp-381 deprotonated (simulations 2a and 2b). No DFG flip was observed in these simulations, suggesting that its time scale is much longer than the several hundred nanoseconds we simulated. This result is consistent with NMR studies of the p38 (13) and PKA (31) kinases, which are suggestive of microsecond- to millisecond-scale DFG motions.

In all 4 simulations, however, substantial motion of helix  $\alpha$ C was observed, despite the fact that no mutation was introduced to promote such motion. In the simulations with Asp-381 protonated, this motion was particularly pronounced, leading to the formation of the Glu-286–Arg-386 salt bridge characteristic of the  $\alpha$ C-out conformation and emergence of a hydrophobic pocket (the N-pocket; see Fig. 1*B*) at the base of the N-lobe near Phe-382 (Fig. 2*A*). The N-pocket, which is contiguous with a region exploited by many drugs (the so-called “kinase specificity pocket”), has been suggested to play a role in facilitating the DFG flip (17). Although simulations 1a and 1b showed the transition from  $\alpha$ C-in to  $\alpha$ C-out (Fig. 1), we did not observe Phe-382 entering into the N-pocket, apparently because of interactions between this residue and the helix- $\alpha$ C residue Met-290. Likewise, we did not observe Phe-382 and Met-290 to adopt the exchanged positions they assume in crystal structures of Abl and Src kinases in  $\alpha$ C-out conformations. These residues remained, for the most part, closely packed during the simulations (Fig. S1), and when they were briefly driven apart by thermal fluctuations, Met-290 remained a steric barrier at the entrance to the N-pocket (Fig. 2*A*).

To lower this barrier, we attempted to further increase the extent of motion of helix  $\alpha$ C by introducing an M290A mutation. Met-290 is not a conserved residue in protein kinases, and the c-Src mutant corresponding to M290A maintains substantial activity (Fig. S2*B*). In Abl, Met-290 is part of a “hydrophobic spine” that stabilizes the active conformation (32), and we anticipated that the M290A mutation might therefore facilitate entry of Phe-382 into the N-pocket by allowing a larger range of motion of helix  $\alpha$ C, as well as by directly reducing the size of the obstructing residue. We



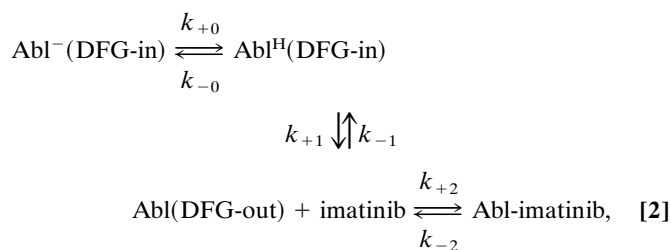




dasatinib (38, 39). It is thought that so-called DFG-in binders typically bind to both DFG-in and DFG-out conformations (13). We find only a very weak pH dependence for dasatinib binding (Fig. 4D), consistent with the interpretation that the pH dependence observed for imatinib binding is a result of 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  $\text{Abl}^{\text{H}}$  and  $\text{Abl}^-$  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  $P_{\text{out}}$  (Eq. 1) as a function of pH for Abl and 3 mutants. These estimates are clearly consistent with our experimental results for  $k_{\text{on}}$ , 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,  $k_{\text{obs}}$ , 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 ( $\text{Mg}^{2+}$ ) 2.3 Å away, which in turn coordinates the  $\beta$  and  $\gamma$  phosphates of ATP (5). Encouraged by observations on small G proteins (45, 46) showing that  $\text{Mg}^{2+}$  binding is weakened by  $\approx 3$  orders of magnitude after transfer of the  $\gamma$  phosphate, we speculate that in kinases, this  $\text{Mg}^{2+}$  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  $\text{Mg}^{2+}$  concentration (44).

The DFG flip could facilitate ADP release, and potentially binding of the ATP- $\text{Mg}^{2+}$  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  $\text{Mg}^{2+}$  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.

**ACKNOWLEDGMENTS.** We thank I. Arkin, D. Borhani, B. Gregersen, M. Jacobson, J. Klepeis, K. Lindorff-Larsen, P. Maragakis, and A. Wang for helpful discussions and critical reading of the manuscript. We also thank S. Miller for use of the stopped-flow equipment. M.A.S. is supported by National Institutes of Health Grant 5K99GM080097.

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