
Copyright:
© 2014 American Chemical Society

This is an open access article published under a Creative Commons Attribution (CC-BY) License, which permits unrestricted use, distribution and reproduction in any medium, provided the author and source are cited.

DOI link to article:
http://dx.doi.org/10.1021/cb500135f

Date deposited:
07/07/2015

This work is licensed under a Creative Commons Attribution 4.0 International License
An Inhibitor’s-Eye View of the ATP-Binding Site of CDKs in Different Regulatory States

Aude Echalier,‡§ Alison J. Hole,‡∥ Graziano Lolli,‡ Jane A. Endicott,‡⊥ and Martin E. M. Noble*‡∥⊥

1Laboratory of Molecular Biophysics, Department of Biochemistry, Oxford University, South Parks Road, Oxford OX1 3QU, United Kingdom
2Department of Chemical Sciences, University of Padua, via Marzolo 1, 35131 Padova, Italy

Supporting Information

ABSTRACT: We have used a chemically diverse panel of kinase inhibitors to assess the chemical similarity of the ATP-binding sites of cyclin-dependent kinase (CDK) subfamily members in a range of activation states. Using this approach, we find that different activation states of a particular CDK may differ from each other as much as different CDKs in the same activation state. We also find that inhibitors discriminate more effectively among CDK family members in their monomeric state than in their cyclin-bound state, providing direct evidence for the belief that selective binding to inactive kinase states might be more readily achieved than selective binding to active states.

Protein kinases represent attractive targets for therapeutic intervention against a wide variety of human diseases, most notably cancer1,2 and chronic inflammatory diseases.3 A number of drugs that selectively target the protein kinase ATP binding site have been successfully introduced into the clinic, and many more are in clinical trials.2,4 However, a significant requirement for the further development of therapeutically useful compounds is a more complete understanding of the factors that dictate inhibitor selectivity across the protein kinase family.5,2,6

The eukaryotic serine/threonine and tyrosine protein kinase family is characterized by a conserved fold in which residues from both the N- and C-terminal lobes contribute to the active site.7,8 The identities of the residues that line the ATP binding pocket and the structural plasticity of the protein kinase fold constitute two key elements that together determine the inhibitor-binding profile of a protein kinase. Both of these elements have been successfully exploited to generate clinically useful drugs.1,2

The cyclin-dependent kinases (CDKs) constitute a subfamily of ca. 13 members in humans9 that play important roles in both the control of cell cycle progression (CDKs 1, 2, 3, 4, and 6) and in the regulation of transcription (CDKs 7, 8, 9, 12, and 13).10–12 CDK2 has provided a structural paradigm for the CDK family and has been widely exploited for structure-aided CDK inhibitor design.13,14 The prevailing structural model for CDK regulation by cyclin and CDK inhibitor (CKI) binding and by phosphorylation has been elaborated through a series of structures of CDK2/cyclin A complexes.15,16 Monomeric CDK2 is inactive as a result of the disposition of active site residues, in turn dependent on the pose of the C-helix, and the conformation of the activation segment.17 Cyclin A binding and Thr160 phosphorylation within the activation segment rearrange the CDK2 active site to orientate key ATP binding and catalytic residues and create the peptide substrate binding site.15,18 This model for the mechanism of regulation appears not to apply across the entire CDK subfamily. The determination of structures for CDK4/cyclin D319 and CDK4/cyclin D1 phosphorylated on Thr172 (pCDK4/D)20 revealed that CDK4 adopts an inactive C-helix out conformation despite being cyclin-bound. Two further examples are CDK521 and CDK822 that both adopt active conformations upon p25 and cyclin C binding, respectively, in the absence of activation loop phosphorylation.

Differential scanning fluorimetry (DSF) can be used to characterize inhibitor binding.23 Here we define ΔT_m as the difference between the apparent melting temperature (T_m) of the protein in the presence of the ligand and the T_m of the protein alone. The value is related to the concentration and binding affinity of the ligand for the protein and to thermodynamic properties of the ligand binding event.23 The profile of ΔT_m values measured for a protein in the presence of a sufficiently diverse range of inhibitors is characteristic of the inhibitor binding properties of that protein and therefore represents a fingerprint of the physicochemical properties of the active site.24 We have used such fingerprints to assess the chemical similarity of a number of CDK subfamily members, namely, CDK2, CDK4, CDK7, and CDK9. Additionally, we...
have explored the inhibitor binding profiles of CDKs in different activation states to assess the extent to which CDK activation affects inhibitor binding.

**RESULTS AND DISCUSSION**

The discriminative power of this fingerprinting approach relies on including a sufficiently diverse panel of inhibitors to sense the chemical diversity apparent in the family of kinase active sites. To validate the use of our selected inhibitor set (Figure 1a and Supplementary Table 1) for this purpose, we confirmed that our set is able to target representative kinases from across the kinome. This analysis was achieved by mapping each inhibitor onto those kinases in the kinome phylogenetic tree that they are best documented to target (Supplementary Methods and Figure 1b). The approximately uniform distribution obtained confirms ours to be a relatively unbiased set of inhibitors.

We included four members of the CDK subfamily in the study. CDK2 was prepared in four different conformational states: monomeric unphosphorylated (CDK2), monomeric phosphorylated on Thr160 (pCDK2), unphosphorylated in complex with human cyclin A175−432 (CDK2/A), and Thr160-phosphorylated in complex with human cyclin A175−432 (pCDK2/A). CDK4 was characterized both as an inactive, nonphosphorylated monomer (CDK4) and as the fully activated binary complex (CDK4/cyclin D3 phosphorylated on Thr172, pCDK4/D). CDK7 and CDK9 were prepared in either a monomeric unphosphorylated (CDK7) or a cyclin-bound phosphorylated (CDK9/cyclin T1 phosphorylated on T186, pCDK9/T) state, respectively (Figure 2a). The identities of the CDKs together with their activation states define a search
Figure 2. CDK space explored in this study. (a) Representative melting curves for cyclin-dependent kinase (CDK)/cyclin complexes used to derive $\Delta T_m$ values in the presence of an ATP-competitive inhibitor. The shift in $T_m$ upon staurosporine binding (red) is illustrated for each of the CDK/cyclin complexes in this study. Black curves, CDK/cyclin complexes in the absence of added inhibitor. Curves with closed triangle symbols correspond to pCDK9/T, closed diamonds to CDK7, closed circles to CDK4, closed squares to pCDK4/D, open squares to CDK2/A, open diamonds to CDK2, and open triangles to pCDK2. (b) The full set of different phosphorylated and/or cyclin-associated states of the four CDKs characterized in this study are shown, with those members for which measurements were made highlighted by shaded boxes. Color intensity for these protein species reflects activation state (more saturated, more active). (c) The ATP binding site of CDKs. The CDK2 ATP-binding site residues from a pCDK2/A structure (PDB code: 1QMZ) are colored according to the degree of sequence conservation between the 4 CDK sequences, calculated as described in the Supplementary Methods, from blue (highly conserved) to red (highly variable). The ATP molecule is represented in ball-and-stick mode for context. (d) The ATP-binding site pocket is highly conserved in CDK2, CDK7, CDK4, and CDK9. Residues that constitute the first shell of the CDK2 ATP-binding site (CDK2 numbering) were identified as defined in Supplementary Methods and used to construct a pseudosequence alignment. This alignment is displayed together with the associated consensus sequence, with X representing any residue, and $\phi$ a hydrophobic amino acid. Sequence motifs that are highly conserved across the protein kinase family are identified by numbered bars. Bar 1, the glycine-rich loop (residues 11−16 in CDK2) that defines the general consensus sequence GXGXXG that recognizes the phosphate moieties of ATP; bar 2, the “hinge” sequence composed of residues 80−84 that links the N- and C-terminal lobes and contributes to the ATP
space within the functional kinome that has been explored by this study and is summarized in Figure 2b. CDK2, CDK7, CDK4, and CDK9 are highly homologous across their entire sequence (Supplementary Figure 1), and this similarity is particularly marked at the catalytic site (Figure 2c,d).

We have adopted the linear correlation coefficient of the inhibitory fingerprints of two kinases as a measure of the chemical similarity of those kinases (Tm values tabulated in Supplementary Table 1, Supplementary Figure 2). By this criterion, we find that the set of kinases studied in the monomeric, unphosphorylated state (CDK2, CDK4, and CDK7) demonstrate significantly different inhibitory profiles, with a lowest correlation coefficient of 0.39 (CDK4 vs CDK7) and a highest correlation coefficient of 0.74 (CDK2 vs CDK7) (Figure 3a). That the inhibitor binding fingerprint discriminates so effectively between highly related CDK subfamily members confirms that it provides a sensitive measure of active site similarity.

One application of this measure is to assess the chemical similarity of the active sites of orthologous proteins from different species. For example, the correlation coefficients obtained for the comparison of human monomeric, unphosphorylated CDKs with the closest CDK orthologue from *Plasmodium falciparum*, protein kinase 5 (PfPK5), range from 0.42 (PfPK5 vs CDK4) to 0.69 (PfPK5 vs CDK2). This result demonstrates that the active sites of orthologous proteins from highly diverged species may be more similar to each other than are the active sites of close paralogues within a species.

For context, the equivalent comparison of each of the monomeric, unphosphorylated CDKs with an “unrelated” kinase, *P. falciparum* protein kinase 7 (PfPK7),25 yields correlation coefficients in the range 0.33 (PfPK7 vs CDK2) to 0.31 (PfPK7 vs CDK2/A).

Next we investigated the apparent active-site similarity of a set of fully activated CDK subfamily members (pCDK2/A, pCDK4/D, and pCDK9/T). Although this comparison also yielded qualitatively different inhibitor fingerprints, our results suggest that the inhibitor-binding properties of the set of fully activated CDKs are more similar to each other than are those of the set of inactive monomeric forms (Figure 3b). The lowest correlation coefficient of 0.73 and the highest correlation coefficient of 0.78 were measured for comparisons of pCDK2/A vs pCDK9/T and pCDK2/A vs pCDK4/D, respectively. This result has two implications: first, it demonstrates that the inhibitor binding properties of CDK subfamily members depend not only on their respective sequence but also on the conformational state in which they are found. Second, it demonstrates that, in adopting an active conformation, two different CDKs assume more similar inhibitor binding properties. This latter point is most directly demonstrated by comparison of the correlations coefficients CDK2 vs CDK4 (CC = 0.69) and pCDK2/A vs pCDK4/D (CC = 0.78).

The inhibitor fingerprints of four different CDK2 activation states were recorded, and the resulting comparison is shown in Figure 3c. As expected from comparative structural studies,26,27 phosphorilation of the activation segment has little effect on the inhibitor-binding fingerprint (correlation coefficient for the
comparison CDK2 vs pCDK2 is 0.94 and for the comparison of CDK2/A vs pCDK2/A is 0.96). Indeed, sequence and structural comparisons demonstrate that phosphorylation of CDK2 or CDK2/A on Thr160 introduces only minor changes to the identity and structure of the amino acids that line the extended inhibitor binding site (Supplementary Figure 3).

By contrast, when pCDK2 associates with cyclin A, the CDK2 inhibitor-binding fingerprint changes significantly (correlation coefficient for pCDK2 vs pCDK2/A is 0.79). Interestingly, the equivalent comparison for unphosphorylated CDK2 demonstrates somewhat less of a response: when cyclin A associates with the unphosphorylated form of CDK2, it perturbs its inhibitor binding profile to a lesser extent (correlation coefficient for CDK2 vs CDK2/A = 0.91). Comparison of the ATP binding site topology highlights a similar degree of conservation between the closest pairs considered, as summarized in Supplementary Figure 3.

For the CDKs analyzed in this study, the abundance of each conformational state within a cell will depend on a large number of factors that include CDK identity, cell type, cell environment, and cell cycle stage. Targeting a specific conformation of a protein kinase is a validated route to the identity and structure of the amino acids that line the extended inhibitor binding site. Cell cycle CDKs of differing phosphorylation status and in monomeric and cyclin-bound forms can be detected in cells, suggesting that such states are accessible for targeting.

In summary, by using DSF we have demonstrated that it is possible to discriminate between closely related CDKs by comparing their inhibitor-binding profiles. We have shown that activation state can have a significant effect on inhibitor-binding profile, with cyclin binding having more profound consequences than phosphorylation at least in the case of CDK2. Our results further suggest that the inhibitor binding properties of a set of fully activated, cyclin-bound CDKs are more similar to each other than are those of a set of inactive, monomeric forms. These data provide direct evidence that, as has been proposed elsewhere, protein kinases may be more amenable to selective inhibitor binding in their inactive states.

**METHODS**

**Inhibitor Set.** A full description of the inhibitor set is provided in the Supporting Information.

**CDK Expression and Purification.** Human CDK4 and cyclin D3, CDK9 and cyclin T1 were all expressed in insect cells and purified as described. CDK2 phosphorylated on Thr160 in association with cyclin A2 (pCDK2/A), unphosphorylated CDK2 in association with cyclin A2 (CDK2/A), monomeric Thr160pCDK2 (pCDK2), monomeric unphosphorylated CDK2 (CDK2), and CDK7 were prepared as described. Monomeric CDK4 was expressed in SF9 cells as a GST fusion and purified by affinity chromatography followed by 3C cleavage of the GST tag and subsequent size-exclusion chromatography. The proteins were concentrated to 2 mg mL−1 and flash frozen in liquid nitrogen prior to storage at −80 °C until further use.

**Differential Scanning Fluorimetry.** Thermal melting experiments were carried out using an Mx3005p Real Time PCR machine (Stratagene) essentially as described. A full description of the method is provided in the Supporting Information.

**Kinase Targets of the Molecules Contained in the Inhibitor Set.** The kinase targeted by each of the inhibitors was identified from either the supplier’s information or by literature analysis, mainly via the work reported in refs 34−36.

**Computation.** Residues corresponding to the first coordination shell of the ATP-binding sites were identified in ref 37. Sequence conservation scores were calculated using the ConSurf webserver (http://consurf.tau.ac.il).

**ASSOCIATED CONTENT**

**Supporting Information**

This material is available free of charge via the Internet at http://pubs.acs.org.

**AUTHOR INFORMATION**

**Corresponding Author**

E-mail: martin.noble@ncl.ac.uk.

**Present Addresses**

1Centre de Biochimie Structurale, INSERM-CNRS-UM1-UM2, 29 rue de Navacelles, 34090 Montpellier cedex, France.

2114 Milton Park, Milton, Abingdon, Oxfordshire OX14 4SA, U.K.

3Newcastle Cancer Centre, Northern Institute for Cancer Research, Newcastle University, Paul O’Gorman Building, Medical School, Framlington Place, Newcastle NE2 4HH, U.K.

**Notes**

The authors declare no competing financial interest.

**ACKNOWLEDGMENTS**

We thank I. Taylor for technical assistance, E. Lowe and D. Staunton for crystallography and biophysics support, and TTP LabTech for help with inhibitor dispensing. We also thank R. Griffin and colleagues in the Department of Chemistry, Newcastle University and S. Knapp (SGC, Oxford) for providing reagents. We are grateful to S. Baumli (Merck Serono) for the gift of purified pCDK9/T and help with thermal denaturation data collection. This research was supported by Framework Program 6 (FP6) of the European Commission (PROKINASE Project, A.E.), and the MRC (J.A.E., A.H., and M.E.M.N.). A.J.H. was supported by a Wellcome Trust studentship (Grant 083113/Z/07/A).

**ABBREVIATIONS**

CDK, cyclin-dependent kinase; DSF, differential scanning fluorimetry; pCDK4/D, Thr172-phosphorylated CDK4/cyclin D3; pCDK9/T, Thr186-phosphorylated CDK9/cyclin T1; pCDK2/A, Thr160-phosphorylated CDK2/cyclin A2

**REFERENCES**


