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Slow Inhibition and Conformation Selective Properties of Extracellular Signal-Regulated Kinase 1 and 2 Inhibitors

Johannes Rudolph^{†,‡}, Yao Xiao^{†,‡}, Arthur Pardi[†], and Natalie G. Ahn^{†,‡,§,*}

[†]Department of Chemistry and Biochemistry, University of Colorado, Boulder, Colorado 80309, United States

[‡]Howard Hughes Medical Institute, University of Colorado, Boulder, Colorado 80309, United States

[§]BioFrontiers Institute, University of Colorado, Boulder, Colorado 80309, United States

Abstract

The mitogen-activated protein (MAP) kinase pathway is a target for anticancer therapy, validated using inhibitors of B-Raf and MAP kinase kinase (MKK) 1 and 2. Clinical outcomes show a high frequency of acquired resistance in patient tumors, involving upregulation of activity of the MAP kinase, extracellular signal-regulated kinase (ERK) 1 and 2. Thus, inhibitors for ERK1/2 are potentially important for targeted therapeutics against cancer. The structures and potencies of different ERK inhibitors have been published, but their kinetic mechanisms have not been characterized. Here we perform enzyme kinetic studies on six representative ERK inhibitors, with potencies varying from 100 pM to 20 μ M. Compounds with significant biological activity ($IC_{50} < 100$ nM) that inhibit in the subnanomolar range (Vertex-11e and SCH772984) display slow-onset inhibition and represent the first inhibitors of ERK2 known to demonstrate slow dissociation rate constants (values of 0.2 and 1.1 h^{-1} , respectively). Furthermore, we demonstrate using kinetic competition assays that Vertex-11e binds with differing affinities to ERK2 in its inactive, unphosphorylated and active, phosphorylated forms. Finally, two-dimensional heteronuclear multiple-quantum correlation nuclear magnetic resonance experiments reveal that distinct

*Corresponding Author: Department of Chemistry and Biochemistry, University of Colorado, Boulder, CO 80309. E-mail: natalie.ahn@colorado.edu. Phone: (303) 492-4799.

Notes

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Supporting Information

Determinations of K_M values for ATP and peptide substrate (Figure S1), representative data for time-dependent assays of a rapid on/off inhibitor of ERK2, olomoucine (Figure S2), representative data for time-dependent assays of a rapid on/off inhibitor of ERK2, SB220025 (Figure S3), representative data for time-dependent assays of a rapid on/off inhibitor of ERK2, Vertex-1 (Figure S4), representative data for a time-dependent inhibitor of ERK2, SCH772984 (Figure S5), and chemical shift perturbations upon binding of Vertex-11e to ERK2 (Figure S6). This material is available free of charge via the Internet at <http://pubs.acs.org>.

conformational states are formed in complexes of Vertex-11e with inactive and active ERK2. Importantly, two conformers interconvert in equilibrium in the active ERK2 apoenzyme, but Vertex-11e strongly shifts the equilibrium completely to one conformer. Thus, a high-affinity, slow dissociation inhibitor stabilizes different enzyme conformations depending on the activity state of ERK2 and reveals properties of conformational selection toward the active kinase.

The mitogen-activated protein (MAP) kinase pathway is a compelling target for anticancer therapy given the prevalence of oncogenic RAS and B-RAF in many cancer types.¹ Small molecule inhibitors specifically targeting B-RAF and MAP kinase kinase 1 and 2 (MKK1/2) in the MAP kinase pathway have been recently approved for the treatment of patients with metastatic melanomas harboring B-RAF^{V600E} activating mutations.² However, while clinical outcomes show impressive disease free survival for a median of 6–7 months with single-line treatment and 13 months using combination therapy with inhibitors of BRAF (dabrafenib) and MKK1/2 (trametinib), nearly all patients eventually relapse.^{3–5} Because of the prevalence of acquired resistance and the ensuing high patient mortality, alternative strategies for targeted therapeutics are an imperative goal.

Many tumors with acquired resistance show an increased copy number or alternative splicing of B-RAF^{V600E}, activating mutations in K-RAS, N-RAS, or MKK1/2, or elevated levels of expression of MAP kinases or receptor tyrosine kinases.^{6–9} In these cases, resistance can be attributed to reactivation of the MAP kinase, extracellular signal-regulated kinase 1 and 2 (ERK1/2).^{8,9} For this reason, ERK is an increasingly attractive target for therapeutics. In recent publications, ERK inhibitors (SCH772984 and ERK-i) have shown efficacy in blocking growth of cancer cell lines with acquired resistance to B-RAF and MKK inhibitors.^{10,11} Both SCH772984 and Vertex-11e, a compound closely related to ERK-i, have been reported to be potent inhibitors *in vitro* (K_d values of 0.19 and 0.11 nM, respectively) and *in vivo* (IC₅₀ values of 60 and 48 nM, respectively).^{10,12} However, to date, the kinetic properties of these molecules toward active ERK2 have not been compared to those of other inhibitors of ERK, and thus, the basis for their potency remains unknown.

ERK1 and -2 are activated by dual phosphorylation at Thr and Tyr residues within the activation loop, both events catalyzed by MKK1/2. X-ray structures of unphosphorylated ERK2 (“0P-ERK2”) and dually phosphorylated ERK2 (“2P-ERK2”) show that phosphorylation rearranges the activation loop to organize residues in the active site and allow productive recognition of substrates containing the phosphorylation motif, Pro-Xxx-pSer/pThr-Pro.^{13,14} However, overall structural changes within the active site of ERK2 are relatively modest, and it is unclear what additional features may explain the >500000-fold increase in k_{cat}/K_m induced by phosphorylation.¹⁵ Using nuclear magnetic resonance (NMR) relaxation dispersion experiments, we have recently discovered that dual phosphorylation of ERK2 induces significant changes in the dynamics of the enzyme.¹⁶ Whereas 0P-ERK2 exists as one major conformer, 2P-ERK2 exists in two separate conformers that exchange on a millisecond time scale. The minor (20%) conformer in 2P-ERK2 corresponds to the conformer seen in 0P-ERK2, while the major (80%) conformer in 2P-ERK2 represents a new form that is energetically inaccessible in 0P-ERK2. Thus, a major effect of ERK2

phosphorylation is to alter the dynamics of the enzyme, lowering the thermodynamic barrier to allow formation of a new conformer, which is accessible only in the activated state.

Such allosteric behavior of ERK2 suggests its possible exploitation by inhibitors with conformation selective properties. Conformational selection is emerging as an important property of certain kinase inhibitors, termed “type II”, which work by trapping nonproductive active site conformers. This was first shown for BCR-ABL kinase inhibitors (e.g., imatinib and posatinib), which induce a “DFG flip” where Asp and Phe side chains in the conserved DFG motif swap positions to form an inactive “DFG-out” conformer.^{17,18} In contrast to “DFG-out”, the EGF receptor inhibitor lapatinib binds an inactive “helix α C-out” conformer, disrupting the Mg^{2+} -coordinating Lys-Glu ion pair.^{19,20} In each case, the allosteric effects of type II inhibitors reinforce their ATP competitive effects.

Here we characterize the kinetics of inhibition by small molecules that target ERK1/2. Importantly, we show that two inhibitors (Vertex-11e and SCH772984), which have the greatest reported cellular efficacy, show slow binding and slow dissociation from ERK2. Slow dissociation is an important property that predicts higher drug potency because of greater cellular retention.^{21,22} Using NMR, we show that Vertex-11e is able to trap ERK2 into different conformations depending on the kinase activity state. Importantly, the thermodynamic equilibrium between conformers seen in 2P-ERK2 is shifted completely to one form upon Vertex-11e binding. Our results indicate that phosphorylation-induced dynamics introduces an allosteric property into ERK2 that can be exploited for conformation selective inhibition.

MATERIALS AND METHODS

Materials

Rat ERK2 was expressed in *Escherichia coli* in its inactive, unphosphorylated form (0P-ERK2) and phosphorylated with the active mutant MKK1-G7B to produce the active, stoichiometrically dually phosphorylated form (2P-ERK2) as previously described.^{23,24} Vertex-11e was purchased from Chemie-Tek. SCH772984 was purchased from Cedarlane Laboratories. Vertex-1 and FR180204 were purchased from Key Organics. ATP, SB220025, and olomoucine were purchased from Sigma-Aldrich.

Enzyme Kinetics

Kinase activity was measured by ^{32}P phosphoryl transfer from [γ - ^{32}P]ATP (1.7–5 Ci/mmol, PerkinElmer) to the peptide substrate, sub-D (Anaspec). Reactions were conducted at 25 °C in 25 mM HEPES (pH 7.4), 10 mM $MgCl_2$, 100 mM NaCl, 1 mM dithiothreitol, and 0.02% Tween 20. ATP concentrations were varied from 3 to 500 μ M for measurements of inhibitor K_i and held fixed at 100 and 500 μ M for measurements of inhibitor association and dissociation kinetics, respectively. Concentrations of sub-D were varied from 0.06 to 20 μ M for measurements of K_m and held fixed at 20 μ M for all assays using inhibitors. Inhibitor concentrations were varied as follows: olomoucine, 1–250 μ M; SB220025, 2–150 μ M; FR180204, 16–8000 nM; Vertex-1, 0.04–40 μ M; Vertex-11e, 0.5–200 nM, and SCH772984, 0.5–100 nM. Reactions with volumes of 10–20 μ L were run in 96-well V-bottom

polystyrene plates (Costar 3897) and quenched with a 10–20-fold excess volume of 200 mM phosphoric acid. The phosphorylated peptide was separated from ATP in a 96-well filter binding apparatus (S&S Minifold I) using P81 ion exchange paper (Whatman). The filter paper was washed in the apparatus with $3 \times 150 \mu\text{L}$ of 200 mM phosphoric acid and then washed in a tray with $3 \times 20 \text{ mL}$ of phosphoric acid. The phosphorylated peptide was detected by exposure of the filter paper to a phosphorimager plate (GE Healthcare) and quantified using ImageQuant (GE Healthcare). Data fitting was performed by least-squares minimization using Microsoft Excel.

For all K_D experiments involving tight-binding inhibitors, activity determinations were performed as described above, adding 10 $\times$ concentrated solutions of ATP and sub-D to initiate the reaction, and using a short reaction time (15 s). In this way, the activity measurements best reflect the equilibrium established during the preincubation. For experiments determining the K_i values of Vertex-11e and SCH772984, the inhibitor concentrations were varied from 3 to 200 nM, preincubating each inhibitor for 30 min with 10–100 nM 2P-ERK2. For the measurements of tight-binding inhibitor interactions with 0P-ERK2, the concentrations of Vertex-11e and SCH772984 were held fixed at 1 and 8 nM, respectively, preincubating each for 30 min with mixtures of 0P-ERK2 (0.2–200 nM) and 2P-ERK2 (4 nM).

NMR Spectroscopy

[*methyl*- ^1H , ^{13}C]Ile-, Leu-, and Val-labeled 0P- and 2P-ERK2 were prepared as previously described¹⁶ in a buffer containing 50 mM Tris (pH 7.4), 150 mM NaCl, 5 mM MgSO_4 , 0.1 mM EDTA, 5 mM dithiothreitol, 100% D_2O , and 2.5% (v/v) glycerol. Vertex-11e was prepared in a 2.5 mM stock solution in d_6 -DMSO, because of its limited solubility in D_2O . Vertex-11e (d_6 -DMSO) was then added to isotopically labeled 2P-ERK2 (40 μM) or 0P-ERK2 (30 μM), to form complexes with a 50% binding stoichiometry ([11e]: [protein] = 0.5:1) and a 100% binding stoichiometry ([11e]: [protein] = 1.2:1). The d_6 -DMSO concentration in the final protein sample was ~1%.

Two-dimensional (2D) ^{13}C - ^1H HMQC spectra of 0P-ERK2 and 2P-ERK2 were collected on Varian VNMRs 800 and 900 MHz NMR spectrometers, respectively. Data were collected at 25 $^\circ\text{C}$ (8 h for 0P-ERK2 and 11 h for 2P-ERK2) and at 5 $^\circ\text{C}$ (12 h for 0P-ERK2 and 18 h for 2P-ERK2). Each spectrum was acquired with 144 (800 MHz) or 160 complex points (900 MHz) in the t_1 (^{13}C) dimension, corresponding to 28.8 ms at 800 MHz and 29.1 ms at 900 MHz, and 1024 complex points in the acquisition period. WURST40 ^{13}C decoupling was applied during the 85 ms acquisition period, and a 1.5 s delay period was used between each scan. The spectra were processed using the software package NMRPipe.²⁵ Time domain data in the ^1H dimension were apodized by a cosine-squared window function and zero-filled prior to Fourier transformation. The indirect dimension (^{13}C) was apodized by a cosine window function and zero-filled prior to Fourier transformation. Spectral visualization and analysis were achieved using CCPNMR Analysis software.²⁶ Chemical shift perturbations upon Vertex-11e binding were calculated by overlaying the 2D ^{13}C - ^1H NMR spectra of 11e-bound and free forms of ERK2. In crowded regions of the spectra, the methyl peaks corresponding to the ligand-bound enzyme were

assigned as the new peaks that appeared closest to the methyl peaks corresponding to the free enzyme. In this way, the chemical shift perturbation values represent minimum estimates. The combined ^{13}C and ^1H chemical shift perturbation (in parts per million) upon binding of Vertex-11e to 0P- or 2P-ERK2 was calculated by the equation $\delta_{\text{C,H}} = (\delta_{\text{H}}^2 + 0.25 \delta_{\text{C}}^2)^{1/2}$, where δ_{H} and δ_{C} are the chemical shift perturbations on the proton and carbon dimensions, respectively.

RESULTS

We performed enzyme assays to examine the kinetic characteristics of six small molecule inhibitors of ERK2 (Table 1). The structures of all of these compounds have been determined crystallographically in complex with 0P-ERK2 and shown to occupy the ATP binding site between N- and C-terminal lobes (Figure 1A–G).^{27–30} A medium throughput assay for 2P-ERK2 activity was developed in 96-well plates using [γ - ^{32}P]ATP and the peptide substrate sub-D. Sub-D has a docking site derived from yeast STE7, which is linked to a C-terminal MAPK phosphorylation consensus sequence by a flexible hydrophobic linker.^{32,33} It has been previously shown to be an excellent substrate for 2P-ERK2, with k_{cat} values of $3\text{--}15 \text{ s}^{-1}$ and a $k_{\text{cat}}/K_{\text{m}}$ of $\sim 5 \mu\text{M}^{-1} \text{ s}^{-1}$, comparable to those of the Ets-1 protein substrate.³² Using 2P-ERK2, we measured a k_{cat} of $4.3 \pm 1.7 \text{ s}^{-1}$ and a $k_{\text{cat}}/K_{\text{m,sub-D}}$ of $4.5 \pm 0.9 \mu\text{M}^{-1} \text{ s}^{-1}$, in good agreement with literature values (Figure S1A of the Supporting Information). We also measured a $K_{\text{m,ATP}}$ of $78 \pm 30 \mu\text{M}$, in agreement with previous reports ($64 \mu\text{M}^{15}$) (Figure S1B of the Supporting Information). Using ATP and sub-D, assays were performed within the linear range of product formation over time courses of 0–8 h with 5 pM kinase and 0–60 s with 5 nM kinase.

We examined the time dependence of inhibition by olomoucine, SB220225, FR180204, and Vertex-1, initiating the reactions by addition of enzyme to a mixture of ATP, sub-D, and each inhibitor. Each experiment showed linear accumulation of the product over 10 min at varying inhibitor concentrations, as expected for rapidly binding inhibitors (Figure 2A and Figures S2A–S4A of the Supporting Information). IC_{50} values were derived using eq 1:

$$\frac{v_i}{v_0} = \frac{1}{1 + \frac{[\text{I}]}{\text{IC}_{50}}} \quad (1)$$

where v_i and v_0 are the observed rates measured in the presence and absence of each inhibitor, respectively, and $[\text{I}]$ is the inhibitor concentration. The resulting IC_{50} measurements for these compounds were in agreement with literature values (Table 1).

We next examined reversibility of inhibition for olomoucine, SB220225, FR180204, and Vertex-1 by preincubating the enzyme with each inhibitor followed by 300-fold dilution into assay buffer containing ATP and sub-D substrate. Rapid recovery of activity was observed upon dilution with each inhibitor, as evidenced by linear accumulation of product over time, with rates identical to those from the control experiment without inhibitor (Figure 2B and Figures S2B–S4B of the Supporting Information). This indicates that compounds that bind rapidly to the enzyme also show rapid release. With the knowledge that olomoucine, SB220225, FR180204, and Vertex-1 are rapid on/off inhibitors, we could measure the

affinity ($K_i = K_d$) of these compounds for 2P-ERK2, using a fixed time point assay at varying inhibitor concentrations and varying ATP. Lineweaver–Burk replots showed that each inhibitor was competitive versus ATP (Figure 2C and Figures S2C–S4C of the Supporting Information), consistent with their occupancy of the ATP site in each crystal structure (Figure 1B–E). Replots of calculated K_m values as a function of inhibitor concentration (Figure 2C, inset, and Figures S2C–S4C of the Supporting Information, inset) yielded K_i values varying from 63 nM to 19 μ M (Table 1).

Unexpectedly, when enzyme was added to Vertex-11e or SCH772984, the time dependence of product accumulation showed strong curvature, revealing a striking hysteric effect with each inhibitor (Figure 3A and Figure S5 of the Supporting Information). This behavior is diagnostic of slow-onset inhibition, wherein the equilibrium between free and enzyme-bound inhibitor is established slowly compared to the time scale of enzymatic turnover.³⁴ Product accumulation (P) during slow-onset inhibition can be fit by eq 2:³⁵

$$P = \frac{v_0}{k_{\text{obs}}} (1 - e^{-k_{\text{obs}}t}) + C \quad (2)$$

where v_0 is the initial velocity observed upon initiation of the reaction with enzyme, k_{obs} is the rate constant for the approach to complete inhibition, and C is a constant to account for the background signal. Replots of k_{obs} (from eq 2) versus inhibitor concentration (Figure 3A, inset, and Figure S5 of the Supporting Information, inset) were linear, supporting a onestep binding model. The slope of this line yielded estimates of k_{on} , the apparent rate constant for association of inhibitor with enzyme (Table 1). Vertex-11e and SCH772984 showed apparent k_{on} values of 0.21 and 2.8 $\mu\text{M}^{-1} \text{s}^{-1}$, respectively, both of which were slower than the rate of free diffusion. Thus, Vertex-11e and SCH772984 could be distinguished from the other compounds by their slow-onset inhibition. Such hysteresis is common for tight-binding inhibitors, although its physical origins are poorly understood.³⁶

Preincubation/dilution experiments were then performed to measure the release of Vertex-11e and SCH772984 from the enzyme (k_{off}). Both inhibitors showed slow recovery of product formation on a time scale of hours (Figure 3B). Product accumulation curves were fit by eq 2, where v_r , the final observed velocity, is used in place of v_0 . Under the conditions of the experiment, k_{off} is essentially equal to k_{obs} , yielding dissociation rate constants for Vertex-11e and SCH772984 of 0.21 and 1.1 h^{-1} , respectively (Table 1). In this manner, the experimentally determined values of k_{on} and k_{off} could be used to calculate binding affinity constants ($K_d = k_{\text{off}}/k_{\text{on}}$) of 0.28 and 0.11 nM for Vertex-11e and SCH772984, respectively (Table 1).

To confirm the dissociation constants for Vertex-11e and SCH772984 derived by the kinetic experiments, we used an alternative approach involving a “Morrison-titration” experiment, wherein a fixed concentration of enzyme and substrate are pre-equilibrated with varying concentrations of inhibitors, and the ratio of inhibitor bound to unbound is determined by an enzymatic assay (Figure 4A).³⁷ The data were fit using eq 3:

$$\frac{v_i}{v_0} = 1 - \frac{[E] + [I] + K_i - \sqrt{([E] + [I] + K_i)^2 - 4[E][I]}}{2[E]} \quad (3)$$

wherein [E] and [I] are the total concentrations of enzyme and inhibitor, respectively, and K_i is the affinity of the inhibitor for the enzyme. The K_i values from these experiments were 0.34 and 0.12 nM for Vertex-11e and SCH772984, respectively, in good agreement with the K_d values measured independently in the kinetic experiments (Table 1).

We next asked whether the affinities of Vertex-11e or SCH772984 differ between 0P-ERK2 and 2P-ERK2. Such differences might influence the responses of the cell to limiting concentrations of inhibitor, given that the abundance of ERK1/2 in cells is $\sim 1 \mu\text{M}$, and dual phosphorylation in response to cell stimulation rarely reaches 100%. However, it is experimentally challenging to determine affinities for tight-binding compounds from kinetic measurements for enzymes like 0P-ERK2, which have little or no activity. This is true because to measure significant activity, assays must be conducted at high enzyme concentrations, well above the expected binding constant. Therefore, we developed an alternative strategy involving competitive titration, wherein varying concentrations of 0P-ERK2 and fixed concentrations of 2P-ERK2 were mixed and preincubated with nanomolar concentrations of inhibitor, followed by addition of ATP and sub-D to initiate reactions and measurement of phosphoryl transfer by 2P-ERK2. When 0P-ERK2 binds inhibitor, it protects 2P-ERK2 from inhibition without contributing to phosphorylation of substrate; therefore, the initial rate of product formation reflects the relative amount of inhibitor-bound versus free 2P-ERK2. The data can then be fit to two instances of eq 3, the first describing binding of the inhibitor to 0P-ERK2 and the second to 2P-ERK2. The fitting yielded a binding affinity of Vertex-11 for 0P-ERK2 ($K_i = 2.5 \pm 0.5 \text{ nM}$) that was 1 order of magnitude weaker than that for 2P-ERK2 (0.34 nM) (Figure 4B and Table 1). This suggests that Vertex-11e either adopts distinct modes of binding to 0P-ERK2 versus 2P-ERK2 or binds by conformational selection in which the binding competent species has very different populations in 0P-ERK2 and 2P-ERK2. In contrast, the binding affinity of SCH772984 for 0P-ERK2 ($K_i = 0.12 \pm 0.04 \text{ nM}$) equaled that of 2P-ERK2 (0.12 nM). Thus, SCH772984 recognizes each enzyme form with comparable affinity, which validates the structural integrity of 0P-ERK2 in this assay.

To investigate the basis for the differential binding affinities of Vertex-11e for 0P-ERK2 versus 2P-ERK2, we examined responses to Vertex-11e binding by NMR. 2D ^{13}C - ^1H HMQC spectra were collected by titrating Vertex-11e into [*methyl*- ^1H , ^{13}C]Ile-, Leu-, and Val-labeled 0P- and 2P-ERK2 and monitoring the chemical shift changes of 70 labeled residues.¹⁶ The titration ended when peaks reflecting unbound ERK2 were undetectable. The chemical shift perturbations upon binding Vertex-11e occurred at similar residues in 0P-ERK2 and 2P-ERK2 (Figure S6 of the Supporting Information). As summarized in Figure 5, significant chemical shift perturbations were observed throughout the N-terminal lobe as well as the upper C-terminal lobe facing the active site, consistent with Vertex-11e occupancy in the ATP binding pocket. In detail, however, differences were observable between 0P- and 2P-ERK2 with respect to the magnitude of chemical shift perturbations. The largest differences occurred at residues L67, L73, I82, I87, I101, L105, and I138, which

were significantly perturbed in 0P-ERK2 but not 2P-ERK2, and residues I29, I70, I81, I84, L113, L154, and I209, which were significantly perturbed in 2P-ERK2 but not 0P-ERK2 (Figure S6 of the Supporting Information, with residue locations in Figure 5). This reveals different modes of binding of Vertex-11e to the inactive versus active forms of ERK2.

Our previous characterization of ERK2 by *methyl*-(^{13}C , ^1H) Carr–Purcell–Meiboom–Gill (CPMG) NMR relaxation dispersion experiments¹⁶ showed that 0P-ERK2 adopts one major conformer (which we will term the “T” conformer, 100%), while 2P-ERK2 undergoes exchange between two different conformers (“T” and “R” conformers, in a 20:80 ratio at 25 °C and a 50:50 ratio at 5 °C). Key residues were found where the chemical shift changes between the two states (in hertz) were larger than the observed chemical exchange rate constant (i.e., $\omega \gg k_{\text{ex}}$), therefore appearing as two peaks in the HMQC spectra of 2P-ERK2. Via comparison of the results of the CPMG to HMQC spectra, the relative intensities for each pair of peaks at these key residues were verified to directly report the relative populations of the T and R conformers.¹⁶

Examination of these key residues showed that different conformations were formed in the complexes of Vertex-11e with inactive versus active kinase (Figure 6A,B). Whereas binding of Vertex-11e to 0P-ERK2 formed the T conformer seen in the 0P-ERK2 apoenzyme, binding to 2P-ERK2 formed the R conformer. Thus, Vertex-11e favors different conformations in ERK2 depending on the kinase activity state, providing a structural basis for explaining the differential affinities of Vertex-11e for 0P-ERK2 and 2P-ERK2. Importantly, binding of the inhibitor to 2P-ERK2 resulted in a substantial shift in equilibrium between T and R conformers. In its apoenzyme form, 2P-ERK2 interconverts between the T and R conformers, whose equilibrium ratios are 20:80 at 25 °C and 50:50 at 5 °C. Upon ligand binding, the equilibrium shifted completely to the R conformer, at both temperatures. This reveals properties of conformational selection in the active kinase and the capability of inhibitor binding to modulate the thermodynamics of conformational exchange.

DISCUSSION

Our study reveals two significant insights into the behavior of inhibitors toward ERK2. First, we present a detailed kinetic analysis of inhibition to determine true binding constants as well as association and dissociation rate constants, greatly expanding previous studies that were restricted to measurements of relative potencies (IC_{50}) for these inhibitors. From this, we demonstrate that Vertex-11e and SCH772984 display the unexpected properties of slow onset and slow dissociation, distinguishing these two compounds from the other inhibitors. Second, we demonstrate that one of these inhibitors, Vertex-11e, binds with differential affinities to inactive, unphosphorylated (0P) and active, phosphorylated (2P) ERK2. Importantly, the inhibitor forms T-state and R-state conformers with the inactive and active enzyme, respectively. In 2P-ERK2, Vertex-11e binding strongly shifts the equilibrium between T and R conformers to favor the R form. Thus, the allosteric properties of ERK2 endow the active form of the kinase with a novel capability of being inhibited through mechanisms involving conformational selection.

Fast-on and fast-off kinetics were exhibited by four of the molecules examined, including the relatively potent inhibitor FR180204 (Figure 2 and Figures S2–S4 of the Supporting Information). Thus, most inhibitors could be described by classic inhibition kinetics, assessed by the measurement of initial rates. In contrast, both Vertex-11e and SCH772984 displayed hysteresis in the form of time-dependent onset of inhibition and slow dissociation (Figure 3 and Figure S5 of the Supporting Information). The K_i values determined for Vertex-11e and SCH772984 (0.34 and 0.12 nM, respectively) agreed well with K_d values determined previously by temperature-dependent fluorescence and increased thermal stability in the bound form (0.11 and 0.19 nM, respectively),¹⁰ although for unknown reasons they differed significantly from K_d values measured using Bio-Layer Interferometry (85 and 13 nM, respectively).³⁰ Although the two slow-binding compounds were comparable to each other with respect to their K_i and K_d values, differences were observed in their association and dissociation rate constants, which were 10- and 3-fold slower, respectively, for Vertex-11e than for SCH772984.

These kinetic measurements reveal fundamental differences between ERK inhibitors that are likely to impact cellular function. Characterization of inhibitor molecules during the drug discovery process typically involves IC_{50} determinations, selectivity screening, cell-based assays, and pharmacological absorption, distribution, metabolism, and excretion in animal models. However, a growing body of evidence suggests that residence time, defined as the inverse of k_{off} , can also benefit the success of a potential drug, by sustaining pharmacological effects *in vivo*, enhancing on-target occupancy with lower off-target effects, and reducing toxicity from degradation products.^{21,22} Our measurements of k_{off} predict long residence times of 4.8 and 0.9 h for Vertex-11e and SCH772984, respectively. As defined by Schramm,³⁸ an “ultimate physiological inhibitor” would maintain binding until the target is replaced by new protein synthesis. Our key finding that slow kinetics can be achieved for ERK1/2 inhibitors suggests that dissociation rates and residence times would be valuable characteristics to examine in structure–activity studies.

The properties of conformational selection in ERK2 can be linked to the regulation of dynamics induced by dual phosphorylation. Our previous work showed that upon phosphorylation, constraints to dynamics are released in ERK2, to allow equilibrium exchange between T and R conformers. In other words, phosphorylation lowers the thermodynamic barrier for forming the R state, allowing conformational exchange in 2P-ERK2 not seen in 0P-ERK2. Thus, the behavior of Vertex-11e resembles that of type II kinase inhibitors, which work by trapping nonproductive active site conformers. Most type II inhibitors described to date stabilize a kinase inactive conformation, involving a “DFG-out” conformer that disrupts Mg^{2+} -ATP binding by distorting the structure of the conserved DFG motif. However, in ERK2, the DFG-out conformation has been achieved only by incorporating multiple mutations, coupled with binding of a type II p38 MAP kinase inhibitor.³⁹ In its wild-type form, ERK2 shows no indication that the DFG-out conformation is allowed. The crystal structures of 0P-ERK2 bound to Vertex-11e and SCH772984 both clearly show a DFG-in conformation.³⁰ Thus, it is unlikely that the T and R conformers differ by a DFG flip. Instead, both Vertex-11e and SCH772984 access a novel binding pocket lined by Tyr36 and Tyr54 between helix αC and the phosphate binding loop.

Mutation of both of these tyrosines has been shown to increase the rates of dissociation of Vertex-11e and SCH772984 from 0P-ERK2 using Bio-Layer Interferometry.³⁰ Thus, it appears that Vertex-11e represents a new variant of type II kinase inhibitors, in that it binds tightly but differentially to 0P- and 2P-ERK2, both in the DFG-in conformation. Nevertheless, structural differences between T and R conformers with respect to inhibitor binding mode are clearly indicated by the differences in chemical shift perturbations between Vertex-11e bound to 0P-ERK2 (T) and 2P-ERK2 (R) (Figure 5A,B). Given that all cocrystal structures so far have been determined using 0P-ERK2, future X-ray structures of inhibitors in complex with 2P-ERK2 are needed to determine the structural differences between ligand-bound T and R conformers.

Importantly, Vertex-11e differs from classic type II inhibitors by showing higher selectivity for the R conformer, which is associated with the active state of ERK2. There are two implications of this finding. First, to the extent that the catalytic function of ERK2 involves $T \rightleftharpoons R$ interconversion, conformational selection by Vertex-11e would reinforce its ATP competitive effects on kinase inhibition by allowing it to work through a conformational trapping mechanism. Thus, the release of the barrier to dynamics by dual phosphorylation represents a new type of allosteric switch for “type II-like” conformation selective inhibitors to exploit. Second, the behavior of Vertex-11e predicts that it would preferentially bind the active form of ERK2 in cells. This might be advantageous under conditions where tissue levels of inhibitor are limiting, given that the cellular abundance of ERK1/2 typically reaches micromolar concentrations. Preferential binding to the active form of the kinase would minimize competition for inhibitor by the larger pool of inactive ERK. We propose that the $T \rightleftharpoons R$ conformational equilibrium in ERK2 provides a new property to exploit in structure–activity studies, which can impact inhibitor mechanism and cellular potency.

Finally, we asked about the degree to which $T \rightleftharpoons R$ conformational exchange accounts for the property of slow dissociation in Vertex-11e. Conformational selection has been associated with long residence times and inhibitor potency. For example, slow off rates have been reported for p38, MET, IGF1R, and VEGFR inhibitors and ascribed to their selective binding to the DFG-out conformation.^{40–44} Slow dissociation and a higher rate of cell retention of ponatinib have also been used to explain its higher potency toward BCR-ABL in primary acute myeloid leukemia cells.¹⁸ Thus, conformational selection in kinase inhibitors may prove to be a valuable means of conferring slow tight binding, leading to higher potency and cellular efficacy.⁴⁵ Initially, we suspected that the slow association or dissociation of Vertex-11e and SCH772984 might be explained by the interconversion between T and R conformers. The most commonly observed mechanism for slow tight binding is a two-step model in which enzyme and inhibitor associate rapidly followed by a slow conformational change to a tight-binding complex ($E + I \rightleftharpoons EI \rightleftharpoons EI^*$).^{46,47} However, two observations argue against this model here. First, the kinetics of both Vertex-11e and SCH772984 are most consistent with a one-step mechanism to form a tight-binding complex ($E + I \rightleftharpoons EI^*$). Although less common than the two-step model, the one-step model has been observed for other tight-binding protein kinase inhibitors.^{48,49} Second, previous NMR CPMG studies of the 2P-ERK2 apoenzyme measured rate constants for $T \rightleftharpoons R$ exchange of 240 and 60 s^{−1} in the forward and reverse directions, respectively. Therefore, it is unlikely that the slow association or dissociation of Vertex-11e is rate-limited by $T \rightleftharpoons R$.

interconversion. Instead, T and R interconvert rapidly in 2P-ERK2 prior to inhibitor binding, and the slow kinetics of binding and dissociation are most likely attributed to structural features in the R conformer, which remain to be determined. Further studies are needed to understand the structural basis of slow inhibition and examine the linkage between allostery and kinetics in ERK2.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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ABBREVIATIONS

ERK	extracellular signal-regulated kinase
MAP	mitogen-activated protein
0P-ERK2	unphosphorylated ERK2
2P-ERK2	dually phosphorylated ERK2
MKK	MAP kinase kinase
PDB	Protein Data Bank

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Figure 1.

Active site of ERK2 in complex with ATP and inhibitors. Shown are X-ray structures of OP-ERK2 bound to (A) ATP (PDB entry 4GT3), (B) olomoucine (PDB entry 4ERK), (C) SB220025 (PDB entry 3ERK), (D) FR180204 (PDB entry 1TVO), (E) Vertex-1 (PDB entry 1OJG), (F) Vertex-11e (PDB entry 4QTE), and (G) SCH772984 (PDB entry 4QTA). Key residues involved in ATP binding (Lys52, Met106, and Asp165) and the catalytic base (Asp147) are shown. In the ATP cocrystal structure, the Lys52 side chain is partially disordered. Graphics were prepared in VMD.³¹

Figure 2.

Representative data for time-dependent assays of rapid on/off inhibitors of ERK2, FR180204. (A) Reactions were initiated by addition of 2P-ERK2 (0.25 nM) in the presence of 20 μM sub-D, 100 μM ATP, and varying concentrations of FR180204. Data were fit with a linear function. (B) Reactions were initiated by 300-fold dilution of a mixture containing 25 nM ERK2 and 4 μM FR180204 into solutions containing 20 μM sub-D and 500 μM ATP. Data were fit with a linear function. (C) Lineweaver–Burk plot for the rapidly reversible inhibitor, FR180204. 2P-ERK2 (1.25 nM) was assayed in the presence of 20 μM sub-D at varying concentrations of ATP (15–500 μM) and FR180204 (32–2000 nM) in a fixed time point assay (5 min). The inset shows the dependence of the apparent K_m on the inhibitor concentration. Data were simultaneously fit using eight different instances of the Michaelis–Menten equation.

Figure 3.

Representative time-dependent data for a slow inhibitor of ERK2, Vertex-11e. (A) Reactions were initiated by adding 2P-ERK2 (0.25 nM) with 20 μM sub-D and 100 μM ATP at varying concentrations of Vertex-11e. Data were fit using eq 2. (B) Reactions were initiated by diluting a mixture containing 2.5 nM ERK2 and 16 nM Vertex-11e by 300-fold into solutions containing 20 μM sub-D and 500 μM ATP. Data were fit using eq 2.

Figure 4.

Representative data for the Morrison experiment with Vertex-11e or SCH772984. (A) 2P-ERK2 was preincubated with inhibitor, using 90 and 11 nM enzyme for Vertex-11e and SCH772984, respectively. Assays were initiated by adding a 10% volume of sub-D and ATP to final concentrations of 20 μ M sub-D and 100 μ M ATP. Data were fit using eq 3. (B) Representative data for a “double-Morrison experiment” developed to determine the affinity of Vertex-11e and SCH772984 for 0P-ERK2. Mixtures of 2P-ERK2 (4 nM) with varying concentrations of 0P-ERK2 were each preincubated with 8 nM inhibitor. Assays were initiated by adding a 10% volume of sub-D and ATP to final concentrations of 20 μ M sub-D and 100 μ M ATP. Data were fit using two instances of eq 3, which were simultaneously minimized to determine K_d values for 0P- and 2P-ERK2. In both panels A and B, the x -axis is plotted on a log scale.

Figure 5.

Chemical shift perturbations following binding of Vertex-11e to ERK2. (A and B) Ile, Leu, and Val methyls showing ^{13}C and ^1H combined chemical shift perturbations induced by binding of Vertex-11e to (A) 0P-ERK2 and (B) 2P-ERK2. The methyls for individual residues were mapped onto a cocrystal structure of 0P-ERK2 and Vertex-9a (blue), an analogue of Vertex-11e (PDB entry 2OJJ¹²). Spheres show Ile, Leu, and Val methyls assigned in the 0P-ERK2 and 2P-ERK2 apoenzymes.¹⁶ Side chains are shown as sticks when the methyl assignment is absent or ambiguous. Colors indicate chemical shift perturbations induced by Vertex-11e binding: <0.05 ppm (white), 0.05–0.1 ppm (green), 0.1–0.2 ppm (orange), and >0.2 ppm (red). Magenta indicates residues with peaks missing in the Vertex-11e–ERK2 complexes, most likely due to changes in methyl dynamics. Labeled residues showed significant differences in chemical shift perturbation upon Vertex-11e binding between ERK2 forms, increasing in 0P-ERK2 (A) or 2P-ERK2 (B) (see Figure S6 of the Supporting Information). In panel B, methyls labeled in italics indicate the positions of I72, V143, and L242, which show no chemical shift perturbation upon binding of Vertex-11e but report the populations of T and R conformers (see Figure 6).

Figure 6.

Vertex-11e stabilizes the R conformer in 2P-ERK2. (A) 2D ^{13}C - ^1H HMQC spectra collected at 25 °C, showing methyl peaks of key residues I72, V143, and L242, which report T and R conformers.¹⁶ Their locations in the structure are shown in Figure 5B. The spectra show that the Vertex-11e-0P-ERK2 complex (pink) adopts the T conformer, observed in the 0P-ERK2 apoenzyme (blue). In contrast, the Vertex-11e-2P-ERK2 complex stabilizes the R conformer (green), shifting the equilibrium between T and R conformers observed in the 2P-ERK2 apoenzyme (black). (B) The same methyl peaks as in panel A, but for spectra collected at 5 °C, showing the more pronounced shift in equilibrium toward the R conformer in the Vertex-11e-2P-ERK2 complex (green), compared to the 2P-ERK2 apoenzyme (black).

Table 1

Kinetic Parameters for Inhibition of 2P-ERK2

Inhibitor	Structure	IC ₅₀ (nM) Published previously	IC ₅₀ (nM) ^a This study	k _{on} (μM ⁻¹ s ⁻¹) ^b	k _{off} (h ⁻¹) ^b	k _{off} /k _{on} (nM)	K _i (nM) ^c
Olomoucine		27,000 (ref 27)	33,000 ± 3,500	n.d.	n.d.	n.d.	10,800 ± 2700
SB220025		18,000 (ref 27)	7,200 ± 1,400	n.d.	n.d.	n.d.	19,000 ± 1,000
FR180204		140 (ref 28)	330 ± 64	>0.8 ^d	n.d.	n.d.	63 ± 17
Vertex-1		2,300 (ref 29)	2,500 ± 550	n.d.	n.d.	n.d.	380 ± 86
Vertex-11e		<2 (ref 12)	n.d.	0.21 ± 0.04	0.21 ± 0.07	0.28 ± 0.073	0.34 ± 0.099 (2P); 2.5 ± 0.5 (0P)
SCH772984		0.24 (ref 10)	n.d.	2.8 ± 0.34	1.1 ± 0.31	0.11 ± 0.021	0.12 ± 0.10 (2P); 0.12 ± 0.04 (0P)

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^a Errors were derived from fits of duplicate experiments using eq 1.

^b Errors were derived from fits of three or more experiments.

^c All measurements describe 2P-ERK2, except where noted for Vertex-11e and SCH772984. Errors for olomoucine, SB220025, FR180204, and Vertex-1 were derived from fits of duplicate experiments. Errors for Vertex-11e and SCH772984 were derived from fits of three or more experiments.

^d FR180204 has a dissociation rate constant of $>0.05 \text{ s}^{-1}$ ($t_{1/2} < 14 \text{ s}$); therefore, its rate constant for association must be $>0.8 \mu\text{M}^{-1} \text{ s}^{-1}$.