



PI3K inhibitors: review and new strategies

Cite this: *Chem. Sci.*, 2020, **11**, 5855Mingzhen Zhang,^a Hyunbum Jang^a and Ruth Nussinov^{*,ab}

All publication charges for this article have been paid for by the Royal Society of Chemistry

The search is on for effective specific inhibitors for PI3K α mutants. PI3K α , a critical lipid kinase, has two subunits, catalytic and inhibitory. PIK3CA, the gene that encodes the p110 α catalytic subunit is a highly mutated protein in cancer. Dysregulation of PI3K α signalling is commonly associated with tumorigenesis and drug resistance. Despite its vast importance, only recently the FDA approved the first drug (alpelisib by Novartis) for breast cancer. A second (GDC0077), classified as PI3K α isoform-specific, is undergoing clinical trials. Not surprisingly, these ATP-competitive drugs commonly elicit severe concentration-dependent side effects. Here we briefly review PI3K α mutations, focus on PI3K drug repertoire and propose new, to-date unexplored PI3K α therapeutic strategies. These include (1) an allosteric and orthosteric inhibitor combination and (2) taking advantage of allosteric rescue mutations to guide drug discovery.

Received 20th March 2020
Accepted 18th May 2020

DOI: 10.1039/d0sc01676d

rsc.li/chemical-science

Introduction

Phosphatidylinositol-3 kinases (PI3Ks) are lipid kinases that phosphorylate signaling lipid PIP₂ to PIP₃.¹ PIP₃ molecules recruit proteins bearing PIP₃-binding pleckstrin homology (PH) domains such as Akt in the PI3K/Akt/mTOR signaling pathway to the plasma membrane. The PI3K pathway is among the most frequently activated in human cancers, impacting almost 50% of the malignancies.² Class IA isoforms PI3K α , β and δ are particularly strongly associated with cancer.³ Activating mutations in the PIK3CA gene,⁴ which encodes the p110 α catalytic subunit of PI3K α and suppressing mutations of tumor suppressor PTEN^{5,6} (a lipid phosphatase that dephosphorylates PIP₃, thus opposes PI3K α action) are the second and third most frequently mutated in cancer, making PI3K α a vastly important target for drug discovery.^{7,8} However, despite considerable efforts, to date the clinical outcome of PI3K inhibitor-based treatments¹ for solid tumors has been disappointing. Reasons include drug resistance such as that resulting from PTEN suppression and lack of specificity thus intolerable toxicity, making innovative drug development a major aim of the pharmaceutical industry. Structural understanding of PI3K α activation mechanism under physiological conditions and in the presence of activating oncogenic mutations is expected to provide a better grasp of its underlying principles toward this aim. The landmark work of Peter Vogt that has uncovered mechanistic principles of PI3K α gain-of-function mutations, including rare mutations, Roger Williams, who pointed out the

premise that oncogenic mutations mimic and enhance dynamic events in natural activation of p110 α , and Mario Amzel who solved the first full PI3K α structure elucidating oncogenic mutations have been major steps in this direction.⁹⁻¹²

PI3K α has two subunits, catalytic (p110) and inhibitory (p85). PIK3CA, which encodes the p110 α , is frequently mutated in cancer. Dysregulation of the PI3K α signaling pathway is commonly associated with tumorigenesis and drug resistance. Despite its vast importance, only in May 2019 the FDA approved the first drug (alpelisib by Novartis) for breast cancer.¹³ A second (GDC0077) is undergoing clinical trials. Although the drug has been classified as PI3K isoform-specific, severe concentration-dependent side effects are commonly observed. This is not unexpected. These drugs, like the previously developed ones, are ATP-competitive and the ATP binding site is almost identical among PI3K isoforms, and they target a major signaling pathway in the cell. Recently, it was discovered that double, triple, and multiple mutations can work conjointly,¹⁴ raising key questions, including why co-occurring mutations, why specific combinations, and how frequent can such events be in oncogenic proteins. Remarkably, alpelisib was observed to potently affect the signaling strength of particular combinations of mutations when targeting PI3K α oncogenes, raising the question of how the drug works.

Below, we briefly discuss the mechanism of PI3K α activation, its oncogenic mutations and their combinations. We review inhibitors developed to block it and their intrinsic off-target toxicities, and outline the challenges encountered in isoform-specific drug discovery. Finally, we propose two new possible strategies for PI3K α specific inhibitor development. The first combines PI3K α orthosteric and allosteric drugs; the second involves allosteric drugs at positions of rescue mutations. Drugs mimicking the allosteric effects of these mutants may rescue

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PI3K α function in cancer. We further delineate determinants to accomplish these approaches.

Allostery plays a role in PI3K α activation by RTK, but not by Ras

Recently we published in *Chem. Sci.* an atomistic-level mechanism of PI3K α activation.¹⁵ In that work we delineated how, *via* the phosphorylated motif at the C-terminal of a receptor tyrosine kinase (RTK) PI3K α relieves its autoinhibition by the nSH2 domain and the details of the allosteric population shifts of the PI3K α ensemble that take place to expose the active site of this lipid kinase at the membrane. The mechanism further clarifies how these conformational changes increase the population of ATP within a catalytically required distance, permitting phosphoryl ion transfer to the signaling lipid PIP₂ to produce PIP₃. In addition, it provided the structural role of Ras in PI3K α activation.¹⁶

Even though the binding of active Ras perturbs the PI3K α structure with allosteric propagation that promotes certain conformational changes,^{17–21} the key question is whether this action is the dominant contribution to activation. RTK recruits PI3K α to the membrane and promotes conformational change that relieves the autoinhibition; Ras acts to stabilize the PIP₂ lipid substrate in the relatively shallow active site that accommodates it. This enhances its population, with the ‘top’ hydrophobic portion of PIP₂ projecting into the plasma membrane. The allosteric conformational

change promoted by Ras is unlikely to play a significant role in activation.

Taken together, the detailed PI3K α activation mechanism helps in understanding how the strong cancer driver mutations activate this lipid kinase; why some other observed oncogenic mutations are weaker drivers, and why certain mutational combinations are more powerful and more potently targeted by these drugs. Hotspot mutations E542K and E545K mimic RTK’s action in releasing the autoinhibition. This action promotes the allosteric conformational changes that expose PI3K α active site at the membrane and reduce the barrier height of the transition state; the H1047R hotspot with a substitution of His by positively charged Arg mimics Ras’ action. The co-acting double mutations observed by Vasan *et al.*¹⁴ pair a strong hotspot with one or more weak activation mutations, such as E453Q/K, E726K and M1043V/I, which either introduce charge reversal at the membrane, or altered hydrophobicity. The changes of the residue charge and hydrophobicity strengthen the activation and signaling. Most of the weak mutations are far away from both the inhibitory nSH2 in the p85 α subunit and the catalytic site, confirming their allosteric roles in activation. Fig. 1 illustrates the PI3K α structure in the autoinhibited and active states, with the location of the mutations highlighted (Fig. 1A and B). While hotspot mutations occur in the helical and kinase domains, most weak mutations are at the surfaces of the ABD and C2 domains. ABD and C2 in p110 α are responsible for iSH2 interactions. The oncogenic mutations in ABD and C2 domains imply a more flexible iSH2 in activated PI3K α . The crystal

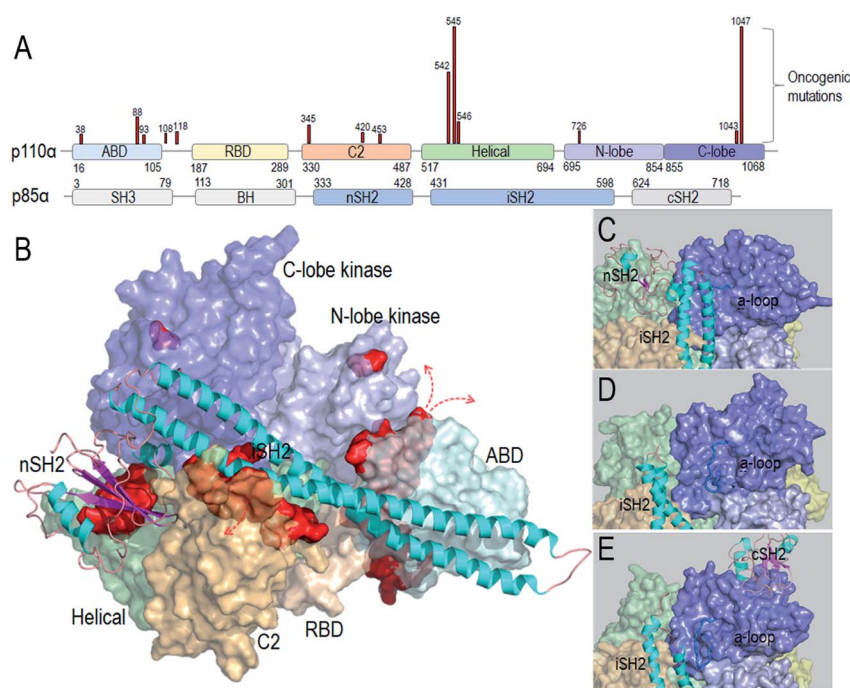


Fig. 1 PI3K α conformation and oncogenic mutations. (A) PI3K α sequence and the frequent oncogenic mutations in p110 α (data obtained from TCGA database). (B) PI3K α structures (PDB ID 4OVV) with the oncogenic mutations highlighted by the red surfaces. The close-up of (C) the PI3K α conformation with nSH2 in the crystal structure (PDB ID 4OVV), (D) the conformation of PI3K α with nSH2 released from the simulations,¹⁵ and (E) the conformation of the PI3K β with nSH2 released (PDB ID 2Y3A).



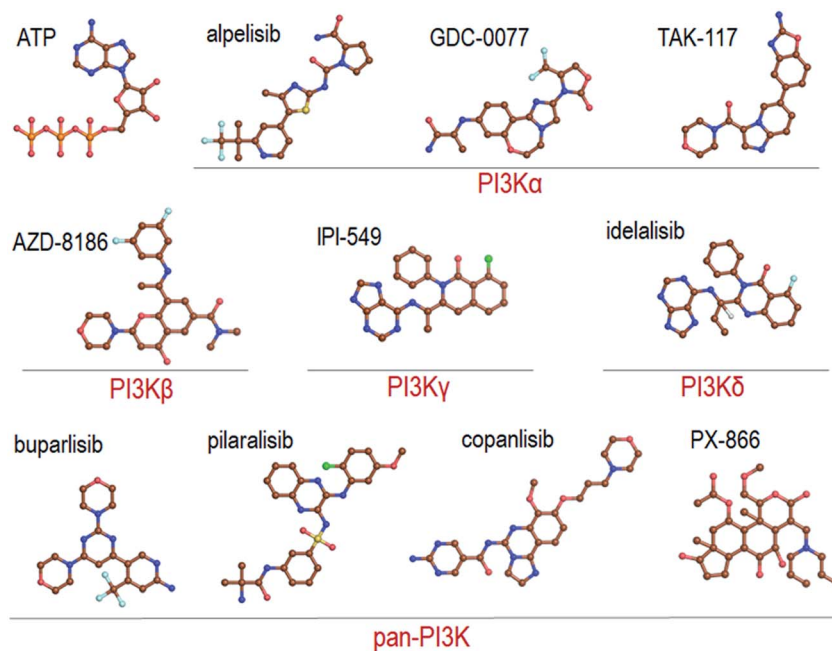


Fig. 2 Summary of the representative pan-PI3K and PI3K isoform-specific ATP-competitive inhibitors. The molecular structures are obtained from Drugbank.

structures of PI3K α containing the inhibitory nSH2 may represent an inactive conformation (Fig. 1C).^{22–24} The active conformation observed in our simulations following nSH2 release differs. Remarkably, the sampled PI3K α active conformation resembles the crystal structure of PI3K β with the inhibitory nSH2 removed (PDB ID 2Y3A), with similar iSH2 and activation loop (a-loop) conformations (Fig. 1D and E). PI3K β is inhibited by both nSH2 and cSH2. The crystal structure of PI3K β without the nSH2 domain may represent a partially active conformation of PI3K,²⁵ implying that PI3K isoforms may share conformational changes for activation.

PI3K α activation takes place at the plasma membrane. It is mediated by interactions with the phosphorylated motif at the C-terminal of an RTK, Ras and the membrane. The activating mutations mimic the effects of these binding events. However, current inhibitors are still stymied by insufficient selectivity, the outcome of the near identity among the ATP binding sites of the PI3K isoforms. At the same time, since ATP is the principal energy currency of the cell with structurally similar cavities hosting it,²⁶ attempts to overcome these hurdles by higher ATP-competitive drug levels encounter toxicity. This argues for allosteric drugs.^{27–34} Unfortunately, these efforts are yet to produce effective drugs. Notwithstanding, the encouraging successes with KRas4B, another hitherto undruggable oncogenic protein, where creative covalent allosteric drugs target KRas4B^{G12C}, and discovery of new PI3K α pockets,²⁷ breed cautious optimism.

Pan PI3K and PI3K α inhibitors

Recent reviews provide a comprehensive coverage of pan-PI3K inhibitors, PI3K α specific inhibitors and PI3K/mTOR

inhibitors, their status and the specific cancers that they target.^{35–38} We thus refer the readers to these. Here we provide a mere brief overview (Fig. 2).

Pan-PI3K inhibitors act on all class I isoforms.³⁸ The outcome is off-target side effects and toxicities.³⁷ Depending on the specific targeted cancer type and its status they include buparlisib^{39–41} and pictilisib (GDC-0941) in breast cancer,^{42,43} and pilaralisib,^{44–47} copanlisib,^{48,49} PX866,^{50–56} CH5132799,⁵⁷ and ZSTK474 and SF1126⁵⁸ in diverse cancers (Fig. 2). Their lack of isoform specificity makes them toxic albeit to different extents.^{37,59} Consequently, most were discontinued. Inhibitors gravitating toward isoform-specificity have lesser off-target toxicity. δ -Isoform inhibitor idelalisib is one example.^{37,60,61} Taselisib inhibits primarily p110 β with 30-fold lower potency⁵⁹ but also reduces mutant p110 α levels. Alpelisib (BYL719), described as PI3K α -specific, is another.⁶² Alpelisib was also prescribed in combination with the aromatase inhibitor letrozole.⁶³ Off-target toxicities were observed. A combination with the BRAF inhibitor encorafenib and anti-EGFR monoclonal antibody cetuximab pointed to clinical efficacy and lower toxicity profiles.⁶⁴ Additional combinations were tested as well, including alpelisib with inhibitors targeting PI3K pathway proteins,⁶⁵ although still with side effects. As expected, at higher concentrations it accomplishes strong PI3K (80%) inhibition. TAK-117 is another PI3K α inhibitor.⁶⁶ The new PI3K α inhibitor GDC-0077 currently undergoes clinical trials in combination with other drugs.⁶⁷ Further details are provided in the reviews cited above. Exhaustive structural analyses of the active sites of all PI3K α class I isoforms have also been reviewed and regions that can play a role in selectivity were described in detail.⁶⁸ Below, we briefly overview them focusing on PI3K α and the interactions with alpelisib.



PI3K α ATP binding site and inhibitor interactions

We illustrate the difficulty in optimizing an ATP-competitive inhibitor specifically to the PI3K α isoform. The ATP binding sites of class I PI3Ks are highly homologous, differing in only a few residues at the rim. The respective residues have been grouped into two regions, Regions 1 and 2.^{69,70} Non-conserved residues in the two regions can aid selectivity. The first non-conserved Region is near the hinge (four residues). The second less variable is at the P-loop. The chemical and conformational differences of these regions among the isoforms define their distinct interactions and have been meticulously probed with the hope that such analyses would help in improving inhibitor specificity.³⁵ The ATP binding site in PI3K isoforms is between the two lobes of the kinase domain, which are separated by a hinge. This is where the adenine ring of ATP and the inhibitors' rings anchor. The hydrophobic residues are conserved, but not the neighboring ones. The so-called specificity pocket shows certain isoform dependent affinity differences.⁷¹ A conformational change of the P-loop upon inhibitor binding exposes it.^{72–74} However, the residues involved are conserved even though those around the pocket are not and may be at play in inhibitor design. There are also differences in the conformational flexibilities. Nonetheless, the opening of the specificity pocket does not appear as determining inhibitor selectivity. ATP also does not penetrate the affinity pocket, which is encircled by the conserved residues.⁷⁵ Despite this, inhibitors entering the pocket have higher potency. P-loop

residues play a role in PI3K α selectivity for GDC-0941,⁷⁶ which was subsequently converted into a more selective covalent drug.⁷⁷ For PI3K α the first non-conserved region appears the key. An inhibitor that extends towards Region 1 (benzoxazepine) adjusts selectivity,^{76,78,79} leading to GDC-0326,⁶⁷ with the drug's carboxamide interacting with Gln859, and Ser854 H-bonding the drug as well.⁶⁷ Additional drugs and their interactions have also been described.⁶⁸

As a detailed example we describe alpelisib, the first drug to be approved by the FDA. The crystal structure of PI3K α with alpelisib has been solved (PDB ID 4JPS). Alpelisib binds to the ATP pocket in the kinase domain, with massive contacts with the P-loop and hinge region. Due to the high sequence similarity in PI3K isoforms, most alpelisib-contacting residues in PI3K α are conserved, with only five variable residues (⁸⁵²RNSH⁸⁵⁵ and Q859) in the hinge region (Fig. 3A and B). Here, we focus on two residues, R852 and Q859, since the side chains of the other three residues (NSH) are exposed to solvent and barely involve in the interactions with alpelisib. Q859 of PI3K α forms dual hydrogen bonds with alpelisib (Fig. 3C). Its importance in isoform selectivity has been indicated and confirmed.^{80,81} This residue becomes shorter in PI3K β (D856) and PI3K δ (N836), while the K890 in PI3K γ is still long enough for the hydrogen bond interaction with alpelisib (Fig. 3D–F). R852 in PI3K α forms a salt bridge with E798 in the kinase domain's N-lobe. A similar salt bridge is observed in PI3K γ (K831–E814). L829 of PI3K γ also establishes hydrophobic interactions with M762 and W760. However, PI3K β is quite different with no favorable interaction of S849. This indicates

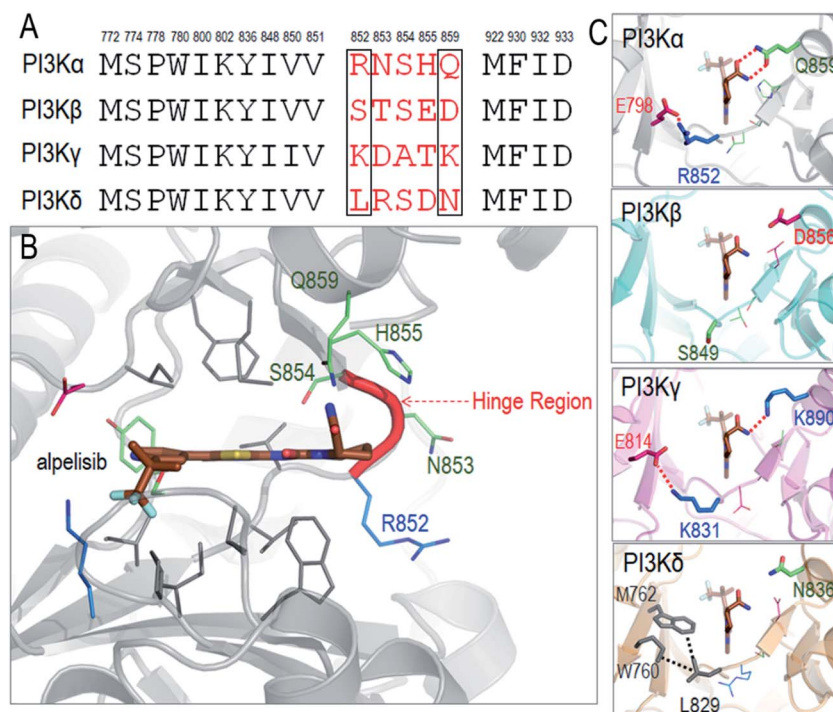


Fig. 3 Structural insights into selectivity of alpelisib in PI3K isoforms. (A) The sequence alignment indicates the conserved and variable residues in PI3K for the interactions with alpelisib. (B) The structure of alpelisib interacting with the ATP pocket in PI3K α (PDB ID 4JPS). (C) The structural comparison of the hinge regions in PI3K isoforms (PDB ID PI3K β -2Y3A; PI3K γ -1E8X; PI3K δ -4XE0). The alpelisib conformations in PI3K isoforms are modeled based on crystal structure of PI3K α with alpelisib (PDB ID 4JPS).



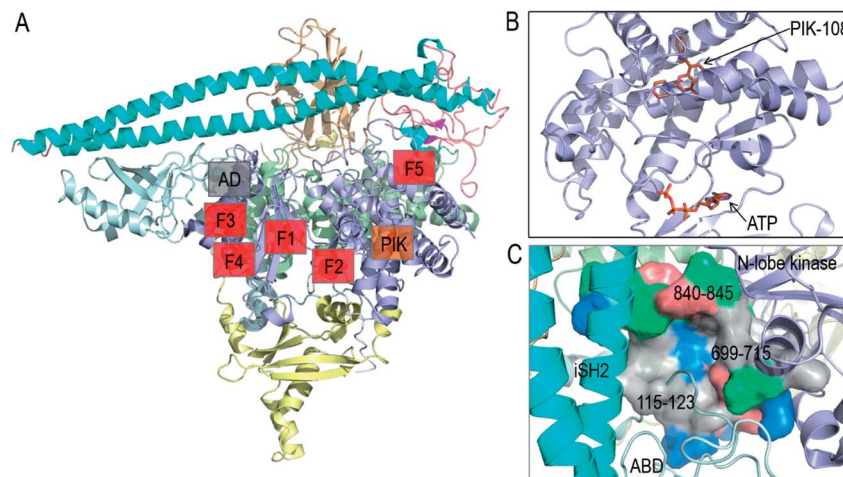


Fig. 4 Summary of PI3K α allosteric inhibitor pockets. (A) The full structure of PI3K α with the allosteric pocket highlighted. The allosteric pocket for PIK-108 (PIK in the figure) is identified from the crystal structure of PI3K α (PDB ID 4A55), the allosteric pockets F1–F5 are obtained from fragment-based screening²⁷ and the allosteric pocket between ABD and kinase domain (AD in the figure) is proposed in the work. (B) The snapshot of PIK-108 interacting with a non-ATP pocket in the C-lobe of kinase domain. The ATP is modeled based on the PI3K structure (PDB ID 1E8X). (C) The snapshot of the proposed allosteric pocket between ABD and N-lobe kinase domain.

that PI3K γ resembles more PI3K α in those residue contacts, followed by PI3K δ and PI3K β . It is interesting that the structural similarity of the hinge region relates to the selectivity of alpelisib to PI3K isoforms. The IC₅₀ values for PI3K α (~5 nM) is highest, followed by PI3K γ (~250 nM), PI3K δ (~290 nM) and PI3K β (~1200 nM).⁸¹ This emphasizes the significance of the hinge region in mediating the isoform selectivity.

In vitro screening of oncogenic mutants has been carried out and micromolar range binders have been identified.⁸² However, they are not mutant-selective. A non-ATP allosteric pocket has been identified in a crystal structure of PI3K α (PDB 4A55). It has been further studied by computational simulations.⁸³ Still, currently its efficacy has been unclear, leading the authors to conclude that this binding site is unlikely to yield productive allosteric inhibitors. Fragment-based screening was also carried out to identify the allosteric inhibitor pockets with promising results²⁷ with phosphopeptide binding at the E542K and E545K hotspots. These allosteric pockets are summarized in Fig. 4A and B. Here, we suggest another allosteric pocket. The oncogenic mutations in C2 and ABD suggest the flexible iSH2 in the activated PI3K α , which has been observed in the simulations of wild type PI3K α and the E545K mutant.^{15,84} The movement of ABD is coupled with the interacting iSH2 domain in PI3K α .⁸⁵ The more stable ABD is expected to interfere with PI3K α activation and increase the population of the inactive PI3K α conformation in the ensemble. By visualizing the PI3K α structure, a pocket was identified between the ABD and the kinase domain's N-lobe. The pocket is deep with considerable hydrophobicity (Fig. 4C). The involved residues include 115–123 in the ABD–RBD linker, and 699–715 and 840–845 in the kinase domain's N-lobe. An inhibitor targeting the pocket is expected to enhance the ABD–kinase domain interface, rendering a higher stability of ABD and iSH2 to restrict the PI3K α inactive conformation and target PI3K α with ABD mutations.

Orthosteric drug design focused on the ATP binding site, with attempts to obtain highly isoform specific inhibitors.

However, they encountered an almost impregnable hurdle: the near identity of the ATP binding sites across isoforms. At the same time, designing drugs to stably bind at the substrate binding site is difficult. With part of the PIP₂ lipid anchored in the membrane with high avidity, the substrate binding site is shallow. The membrane-buried protein surface further constrains small molecule access. Currently available inhibitors mimicking ATP require high concentrations, resulting in toxicities. The frequent emerging deletion or mutations of PTEN, which suppresses cancer by dephosphorylating the PI3K-phosphorylated lipid PIP₃ back to PIP₂, which is also with no currently available inhibitors, aggravates the pharmacological scenario. PTEN is highly challenging to target: to function the inhibitor needs to activate, not suppress it. Drug-induced activation of an enzyme is difficult to accomplish.

To-date unexplored PI3K α therapies

We propose exploring two new strategies for PI3K α inhibition: (i) combination of allosteric and orthosteric drugs and (ii) taking advantage of rescue mutations to guide drug discovery. They can also be merged. Below, we describe the background, rationale, and map a strategy for their implementation.

(i) Allosteric plus orthosteric drugs

A few years ago, a remarkable observation was made: a Cys to Tyr mutation (C1156Y) in metastatic anaplastic lymphoma kinase (ALK) desensitized it to crizotinib, an ATP-competitive drug that was given earlier. Lorlatinib was then prescribed; however, drug resistance emerged. Sequencing indicated that an emerging allosteric mutation involving L1198F substitution conferred resistance to lorlatinib, also an ATP-competitive small-molecule inhibitor of ALK and the related tyrosine kinase ROS1. The L1198F mutation led to steric interference



with lorlatinib binding. ALK L1198F has been reported as an ALK gain-of-function mutation. However, remarkably the allosteric L1198F mutation resensitized ALK to crizotinib. The patient harboring this mutation received crizotinib again, and the cancer-related symptoms abated.⁸⁶ In this example L1198F is a rescue mutation. The mutation redistributed the ALK conformational ensemble, now re-populating a crizotinib binding-favored state. In line with this observation, loss of G2032R resistance mutation upon chemotherapy treatment enabled successful crizotinib rechallenge in a patient with ROS1-rearranged cancer.⁸⁷ Thus, once the mutation occurred, there is no need for an allosteric drug that would mimic its action to combine with the ATP-competitive crizotinib. Had there not been such a mutation at that position, an allosteric drug mimicking it could conceivably accomplish this role, promoting a drug-accommodating conformational change at the ATP binding site. Drug resistance often emerges from conformational changes.⁸⁸ Allosteric drugs can resensitize an active site in the same way that an allosteric mutation does, restoring the actions of orthosteric drugs. As we describe below, experiments in Nathaniel Gray's Lab on BCR-ABL kinase toward treatment of chronic myeloid leukemia (CML) along with subsequent work provide an outstanding example.

Imatinib, an orthosteric ATP-competitive drug, can inhibit oncogenic mutants of BCR-ABL. However, mutations often affect BCR-ABL's ATP-binding site or the activation loop can result in drug resistance. Among these, T315 substitutions that elicit steric clashes with the drug are particularly potent. By screening drug libraries the Gray team first identified an allosteric drug GNF-2 with similar antiproliferative outcome as the orthosteric drug.⁹⁰ Subsequent optimization obtained the allosteric GNF-5 compound.⁹⁰ Remarkably, like the L1198F rescue mutation in ALK, the GNF drug series shifted the protein ensemble toward an active site that re-sensitized BCR-ABL to imatinib and nilotinib. NMR, crystallography, mutagenesis and hydrogen exchange mass spectrometry all indicated that the drug binds to the C-terminal myristate pocket of BCR-ABL. The structure unraveled formation of considerable hydrophobic as well as vdW interactions and water-mediated H-bonds, indicating how the combination of the GNF-series allosteric drug collaborates with the ATP-competitive imatinib, overcoming the T315 drug-resistant hotspots.^{89,90}

ABL001 (asciminib) is also a potent and selective allosteric ABL1 inhibitor that binds to the myristoyl pocket of ABL1 in a manner resembling the GNF-series. It interacts with active site residues *via* the same type of interactions as the GNF inhibitors, and shifts the ensemble toward the inactive kinase conformation. Each drug alone, ABL001 and the catalytic ATP-competitive inhibitor nilotinib are blocked by distinct drug resistant mutations. However, as above, their combination eradicates the tumors.⁹¹ Thus, even though mutations at the myristate site can confer resistance to ABL001, they are sensitive to nilotinib or other ATP-competitive inhibitors making their combination (or ABL001 with other ATP-competitive drugs, like imatinib or dasatinib) a viable BCR-ABL inhibition strategy.⁹² Combining the allosteric inhibitor asciminib with ATP-competitive ponatinib was recently found to even more powerfully suppress the

emergence of mutants (Y253H and E255V) and restore efficacy against highly resistant BCR-ABL1 mutants.⁹³ Additional examples of such combinations are available.^{94,95}

The concept of allosteric drug working conjointly with an orthosteric drug is not new. We only need to consider that a 'ligand' can be an orthosteric drug, with the allosteric drug acting to modulate it.⁹⁶ This powerful combinatorial concept has shown promise for the protein kinases above; but has not been tested for PI3K α . Allosteric drugs may overcome PI3K α drug resistance by altering the ATP binding site and resensitizing it, *i.e.* making it accessible to sterically blocked ATP-competitive orthosteric drugs, or by producing an altered, druggable PIP₂ binding site. Cross-correlations of the motions and analysis of the conformational ensembles in long timescale molecular dynamics simulations may be one way to help identify likely targetable allosteric sites.

(ii) Rescue mutations provide a powerful concept

That mutations can rescue, or resensitize, a mutant protein to an allosteric drug has been shown for a number of diseases. As such they have been dubbed "natural gene therapy" and "nature's proof of principle that you can fix those mutations" in the case of Wiskott-Aldrich syndrome (WAS).⁹⁷ Additional examples include the p.Q188R mutation in classic galactosemia⁹⁸ that gives rise to a protein with lower conformational stability and lower catalytic activity⁹⁹ and p53.¹⁰⁰ Drugs mimicking the allosteric effects of these mutants may rescue function. To the best of our knowledge, in practice this however has not been followed up, likely due to the design complexity which is involved. Nonetheless, identifying them in protein structures, and their potential substitutions will provide allosteric sites and potentially certain orthosteric-drug favored conformational changes. Here we propose attempting this strategy in targeting PI3K α .

How to *a priori* identify the positions of such mutations? Earlier we have taken this up for von Hippel-Lindau tumor suppressor protein (pVHL).¹⁰¹ To identify possible candidate residues, we used molecular dynamics simulations, seeking the least stable regions in the unbound state of the protein. This flagged the interface between the two domains. We then designed several stabilizing mutations at the interdomain interface and in the domains. Experiments showed that disease mutant Y98N at the HIF binding site destabilizes pVHL and lowers its affinity to hypoxia-inducible factor (HIF). pVHL is part of the E3 ubiquitin ligase complex that regulates the degradation of HIF. The simulations indicated that pVHL stability and binding affinity are allosterically regulated. We thus sought to allosterically stabilize pVHL and its interactions with mutations away from binding sites. The mutations were selected based on sequence alignment with a pVHL from other species (worm pVHL sequence). The worm-based substitutions stabilized pVHL, its Y98N mutant, and the complexes of pVHL with HIF, lowering its binding free energy, and the Y98N mutant with elongin C complex, thus rescuing pVHL-HIF affinity and function. The reason for choosing the worm sequence was our observation that the worm p53 is significantly more stable than



the human form.¹⁰² In PI3K α , rescuing mutations could improve the affinity of the ATP-competitive drug to its binding site without increasing its concentration, by allosterically altering its conformation. The mutations in the kinase domain's P-loop may be promising. Such a strategy may also be considered for PTEN tumor suppressor.

Alternative strategies and considerations to map and harness allosteric candidates to improve PI3K α isoform-specific inhibitor design were also proposed.¹⁰³ Key steps in successful design of allosteric drugs include identification of allosteric sites^{103–105} including hidden cryptic allosteric pockets^{106,107} shown to underlie allosteric modulator selectivity at muscarinic GPCRs,¹⁰⁸ allosteric mutations^{19,109–118} and allosteric drug design.^{119,120} Allosteric drugs can constitute 'anchors' and 'drivers'.¹⁰³ The anchor docks into an allosteric pocket whose conformation is unchanged during the transition between the inactive and active states. The driver 'pulls' and/or 'pushes' protein atoms (side-chain or backbone) and triggers shifts of the receptor population to a drug binding-favored state. Based on crystal structures, characteristics of anchors and drivers have been outlined. The recent computational tools quantifying the allosteric signaling, mutation effects and their combinational actions in the signaling promote the discovery of allosteric effectors.^{121,122}

Conclusions

To understand the behavior of biomacromolecules we need to look at the distributions of their conformational ensembles and how these are affected by their environments. Under physiological conditions PI3K α populates the inactive state. Oncogenic mutations will shift the ensemble to populate the active state. The extent of the shift determines the strength of the mutation. The chemistry of the surface matters: the membrane surface is negatively charged, with lipid hydrophobicity playing a major role. Mutations that contribute positive charge will enhance the interactions. Mutations (orthosteric or allosteric) that weaken the interactions of p85 α with p110 α or between the p85 α domains or strengthen the interaction with the membrane will shift the population toward the active state. Double (strong and weak) mutations can act to more fully shift the population. Drug resistance mutations can alter the ATP binding site, shifting the ensemble toward orthosteric drug-inaccessible state. Allosteric modulator drug can shift the population toward drug accessibility. Rescue mutations can identify regulatory hot spots that nature uses for functional control. Small molecule allosteric inhibitors can take aim at such hot spots in tough-to-drug targets such as kinases and phosphatases, like the SHP2.

Here we described the PI3K α saga from the free energy landscape standpoint¹²³ and its dynamic shifts following changes in the environment.^{110,124–126} Allostery is a powerful ensemble property used by nature. Harnessing it to favor an ATP-competitive drug exploits "nature's proof of principle" may be one way to target PI3K α , the second most highly mutated protein in cancer. Here we proposed two new possible venues not exploited yet in PI3K α drug discovery: allosteric plus

orthosteric drug combinations and following mother nature rescue mutation strategy.

Conflicts of interest

No potential conflict of interest was disclosed.

Acknowledgements

This project has been funded in whole or in part with federal funds from the National Cancer Institute, National Institutes of Health, under contract HHSN26120080001E. The content of this publication does not necessarily reflect the views or policies of the Department of Health and Human Services, nor does mention of trade names, commercial products, or organizations imply endorsement by the U.S. Government. This research was supported [in part] by the Intramural Research Program of the NIH, National Cancer Institute, Center for Cancer Research.

References

- 1 A. C. Carrera and R. Anderson, *J. Cell Sci.*, 2019, **132**, jcs228395.
- 2 T. A. Yap, L. Bjerke, P. A. Clarke and P. Workman, *Curr. Opin. Pharmacol.*, 2015, **23**, 98–107.
- 3 S. J. Klemptner, A. P. Myers and L. C. Cantley, *Cancer Discovery*, 2013, **3**, 1345–1354.
- 4 *S. I. C. database*, <http://www.sanger.ac.uk/genetics/CGP/cosmic/>.
- 5 K. K. Wong, J. A. Engelman and L. C. Cantley, *Curr. Opin. Genet. Dev.*, 2010, **20**, 87–90.
- 6 K. D. Courtney, R. B. Corcoran and J. A. Engelman, *J. Clin. Oncol.*, 2010, **28**, 1075–1083.
- 7 P. A. Clarke and P. Workman, *J. Clin. Oncol.*, 2012, **30**, 331–333.
- 8 M. S. Lawrence, P. Stojanov, C. H. Mermel, J. T. Robinson, L. A. Garraway, T. R. Golub, M. Meyerson, S. B. Gabriel, E. S. Lander and G. Getz, *Nature*, 2014, **505**, 495–501.
- 9 L. Zhao and P. K. Vogt, *Proc. Natl. Acad. Sci. U. S. A.*, 2008, **105**, 2652–2657.
- 10 M. Gymnopoulos, M. A. Elsliger and P. K. Vogt, *Proc. Natl. Acad. Sci. U. S. A.*, 2007, **104**, 5569–5574.
- 11 J. E. Burke, O. Perisic, G. R. Masson, O. Vadas and R. L. Williams, *Proc. Natl. Acad. Sci. U. S. A.*, 2012, **109**, 15259–15264.
- 12 C. H. Huang, D. Mandelker, O. Schmidt-Kittler, Y. Samuels, V. E. Velculescu, K. W. Kinzler, B. Vogelstein, S. B. Gabelli and L. M. Amzel, *Science*, 2007, **318**, 1744–1748.
- 13 A. Markham, *Drugs*, 2019, **79**, 1249–1253.
- 14 N. Vasani, P. Razavi, J. L. Johnson, H. Shao, H. Shah, A. Antoine, E. Ladewig, A. Gorelick, T. Y. Lin, E. Toska, G. Xu, A. Kazmi, M. T. Chang, B. S. Taylor, M. N. Dickler, K. Jhaveri, S. Chandralapaty, R. Rabadan, E. Reznik, M. L. Smith, R. Sebra, F. Schimmoller, T. R. Wilson, L. S. Friedman, L. C. Cantley, M. Scaltriti and J. Baselga, *Science*, 2019, **366**, 714–723.



- 15 M. Z. Zhang, H. Jang and R. Nussinov, *Chem. Sci.*, 2019, **10**, 3671–3680.
- 16 M. Z. Zhang, H. Jang and R. Nussinov, *Phys. Chem. Chem. Phys.*, 2019, **21**, 12021–12028.
- 17 R. Nussinov, C. J. Tsai and H. Jang, *Front. Oncol.*, 2019, **9**, 1231.
- 18 C. J. Tsai and R. Nussinov, *PLoS Comput. Biol.*, 2014, **10**, e1003394.
- 19 J. Liu and R. Nussinov, *PLoS Comput. Biol.*, 2016, **12**, e1004966.
- 20 R. Nussinov, *Chem. Rev.*, 2016, **116**, 6263–6266.
- 21 D. D. Boehr, R. Nussinov and P. E. Wright, *Nat. Chem. Biol.*, 2009, **5**, 789–796.
- 22 M. S. Miller, O. Schmidt-Kittler, D. M. Bolduc, E. T. Brower, D. Chaves-Moreira, M. Allaire, K. W. Kinzler, I. G. Jennings, P. E. Thompson, P. A. Cole, L. M. Amzel, B. Vogelstein and S. B. Gabelli, *Oncotarget*, 2014, **5**, 5198–5208.
- 23 W. C. Hon, A. Berndt and R. L. Williams, *Oncogene*, 2012, **31**, 3655–3666.
- 24 J. Yu, C. Wjasow and J. M. Backer, *J. Biol. Chem.*, 1998, **273**, 30199–30203.
- 25 X. X. Zhang, O. Vadas, O. Perisic, K. E. Anderson, J. Clark, P. T. Hawkins, L. R. Stephens and R. L. Williams, *Mol. Cell*, 2011, **41**, 567–578.
- 26 S. Y. Lu, W. K. Huang, Q. Wang, Q. C. Shen, S. Li, R. Nussinov and J. Zhang, *Biophys. J.*, 2015, **108**, 528a.
- 27 M. S. Miller, S. Maheshwari, F. M. McRobb, K. W. Kinzler, L. M. Amzel, B. Vogelstein and S. B. Gabelli, *Bioorg. Med. Chem.*, 2017, **25**, 1481–1486.
- 28 X. Lu, J. B. Smaill and K. Ding, *Angew. Chem., Int. Ed. Engl.*, 2020, **59**, 2–15.
- 29 A. E. Leroux, L. Z. F. Gross, M. Sacerdoti and R. M. Biondi, *Adv. Exp. Med. Biol.*, 2019, **1163**, 279–311.
- 30 X. Cheng and H. Jiang, *Adv. Exp. Med. Biol.*, 2019, **1163**, 1–23.
- 31 R. Nussinov and C. J. Tsai, *Annu. Rev. Pharmacol. Toxicol.*, 2015, **55**, 249–267.
- 32 S. Lu, Y. Qiu, D. Ni, X. He, J. Pu and J. Zhang, *Drug Discovery Today*, 2020, **25**, 177–184.
- 33 A. Zorba, V. Nguyen, A. Koide, M. Hoemberger, Y. Zheng, S. Kutter, C. Kim, S. Koide and D. Kern, *Proc. Natl. Acad. Sci. U. S. A.*, 2019, **116**, 13937–13942.
- 34 Q. Tang, M. T. Villar, A. Artigues, J. P. Thyfault, U. Apte, H. Zhu, K. R. Peterson and A. W. Fenton, *Sci. Rep.*, 2019, **9**, 9031.
- 35 A. E. Garces and M. J. Stocks, *J. Med. Chem.*, 2019, **62**, 4815–4850.
- 36 J. Yang, J. Nie, X. L. Ma, Y. Q. Wei, Y. Peng and X. W. Wei, *Mol. Cancer*, 2019, **18**, 26.
- 37 F. Janku, T. A. Yap and F. Meric-Bernstam, *Nat. Rev. Clin. Oncol.*, 2018, **15**, 273–291.
- 38 R. Arafah and Y. Samuels, *Semin. Cancer Biol.*, 2019, **59**, 36–49.
- 39 J. Baselga, S. A. Im, H. Iwata, J. Cortes, M. De Laurentiis, Z. Jiang, C. L. Arteaga, W. Jonat, M. Clemons, Y. Ito, A. Awada, S. Chia, A. Jagiello-Gruszfeld, B. Pistilli, L. M. Tseng, S. Hurvitz, N. Masuda, M. Takahashi, P. Vuylsteke, S. Hachemi, B. Dharan, E. Di Tomaso, P. Urban, C. Massacesi and M. Campone, *Lancet Oncol.*, 2017, **18**, 904–916.
- 40 A. Di Leo, S. Johnston, K. S. Lee, E. Ciruelos, P. E. Lonning, W. Janni, R. O'Regan, M. A. Mouret-Reynier, D. Kalev, D. Egle, T. Csozsi, R. Bordonaro, T. Decker, V. C. G. Tjan-Heijnen, S. Blau, A. Schirone, D. Weber, M. El-Hashimy, B. Dharan, D. Sellami and T. Bachelot, *Lancet Oncol.*, 2018, **19**, 87–100.
- 41 M. Martin, A. Chan, L. Dirix, J. O'Shaughnessy, R. Hegg, A. Manikhas, M. Shtivelband, P. Krivorotko, N. Batista Lopez, M. Campone, M. Ruiz Borrego, Q. J. Khan, J. T. Beck, M. Ramos Vazquez, P. Urban, S. Goteti, E. Di Tomaso, C. Massacesi and S. Delaloge, *Ann. Oncol.*, 2017, **28**, 313–320.
- 42 I. E. Krop, I. A. Mayer, V. Ganju, M. Dickler, S. Johnston, S. Morales, D. A. Yardley, B. Melichar, A. Forero-Torres, S. C. Lee, R. de Boer, K. Petrakova, S. Vallentin, E. A. Perez, M. Piccart, M. Ellis, E. Winer, S. Gendreau, M. Derynck, M. Lackner, G. Levy, J. Qiu, J. He and P. Schmid, *Lancet Oncol.*, 2016, **17**, 811–821.
- 43 P. Vuylsteke, M. Huizing, K. Petrakova, R. Roylance, R. Laing, S. Chan, F. Abell, S. Gendreau, I. Rooney, D. Apt, J. Zhou, S. Singel and L. Fehrenbacher, *Ann. Oncol.*, 2016, **27**, 2059–2066.
- 44 S. Tolaney, H. Burris, E. Gartner, I. A. Mayer, C. Saura, M. Maurer, E. Ciruelos, A. A. Garcia, F. Campana, B. Wu, Y. Xu, J. Jiang, E. Winer and I. Krop, *Breast Canc. Res. Treat.*, 2015, **149**, 151–161.
- 45 V. G. Abramson, J. G. Supko, T. Ballinger, J. M. Cleary, J. F. Hilton, S. M. Tolaney, N. G. Chau, D. C. Cho, J. Pearlberg, J. Lager, G. I. Shapiro and C. L. Arteaga, *Clin. Cancer Res.*, 2017, **23**, 3520–3528.
- 46 J. Wheler, D. Mutch, J. Lager, C. Castell, L. Liu, J. Jiang and A. M. Traynor, *Oncologist*, 2017, **22**, 377–e37.
- 47 J. C. Soria, P. LoRusso, R. Bahleda, J. Lager, L. Liu, J. Jiang, J. F. Martini, S. Mace and H. Burris, *Oncologist*, 2015, **20**, 245–246.
- 48 A. Patnaik, L. J. Appleman, A. W. Tolcher, K. P. Papadopoulos, M. Beeram, D. W. Rasco, G. J. Weiss, J. C. Sachdev, M. Chadha, M. Fulk, S. Ejadi, J. M. Mountz, M. T. Lotze, F. G. S. Toledo, E. Chu, M. Jeffers, C. Pena, C. Xia, S. Reif, I. Genvresse and R. K. Ramanathan, *Ann. Oncol.*, 2016, **27**, 1928–1940.
- 49 M. Dreyling, A. Santoro, L. Mollica, S. Leppa, G. A. Follows, G. Lenz, W. S. Kim, A. Nagler, P. Panayiotidis, J. Demeter, M. Ozcan, M. Kosinova, K. Bouabdallah, F. Morschhauser, D. A. Stevens, D. Trevarthen, M. Giurescu, L. Cupit, L. Liu, K. Kochert, H. Seidel, C. Pena, S. X. Yin, F. Hiemeyer, J. Garcia-Vargas, B. H. Childs and P. L. Zinzani, *J. Clin. Oncol.*, 2017, **35**, 3898–3905.
- 50 D. S. Hong, D. W. Bowles, G. S. Falchook, W. A. Messersmith, G. C. George, C. L. O'Bryant, A. C. H. Vo, K. Klucher, R. S. Herbst, S. G. Eckhardt, S. Peterson, D. F. Hausman, R. Kurzrock and A. Jimeno, *Clin. Cancer Res.*, 2012, **18**, 4173–4182.



- 51 D. W. Bowles, W. W. Ma, N. Senzer, J. R. Brahmer, A. A. Adjei, M. Davies, A. J. Lazar, A. Vo, S. Peterson, L. Walker, D. Hausman, C. M. Rudin and A. Jimeno, *Br. J. Cancer*, 2013, **109**, 1085–1092.
- 52 C. Yam, X. W. Xu, M. A. Davies, P. A. Gimotty, J. J. D. Morrisette, M. T. Tetzlaff, K. M. Wani, S. J. Liu, W. L. Deng, M. Buckley, J. H. Zhao, R. K. Amaravadi, N. B. Haas, R. R. Kudchadkar, A. C. Pavlick, J. A. Sosman, H. Tawbi, L. Walker, L. M. Schuchter, G. C. Karakousis and T. C. Gangadhar, *Clin. Cancer Res.*, 2018, **24**, 22–32.
- 53 A. Jimeno, J. E. Bauman, C. Weissman, D. Adkins, I. Schnadig, P. Beauregard, D. W. Bowles, A. Spira, B. Levy, N. Seetharamu, D. Hausman, L. Walker, C. M. Rudin and K. Shirai, *Oral Oncol.*, 2015, **51**, 383–388.
- 54 A. Jimeno, K. Shirai, M. Choi, J. Laskin, M. Kochenderfer, A. Spira, V. Cline-Burkhardt, E. Winquist, D. Hausman, L. Walker and R. B. Cohen, *Ann. Oncol.*, 2015, **26**, 556–561.
- 55 D. W. Bowles, N. Senzer, D. Hausman, S. Peterson, A. Vo, L. Walker, R. B. Cohen and A. Jimeno, *Invest. New Drugs*, 2014, **32**, 1197–1203.
- 56 D. W. Bowles, M. Kochenderfer, A. Cohn, L. Sideris, N. Nguyen, V. Cline-Burkhardt, I. Schnadig, M. Choi, L. Nabell, A. Chaudhry, R. Ruxer, A. Ucar, D. Hausman, L. Walker, A. Spira and A. Jimeno, *Clin. Colorectal Cancer*, 2016, **15**, 337–344.
- 57 S. Blagden, A. Omlin, D. Josephs, C. Stavrou, A. Zivi, D. J. Pinato, A. Anthoney, S. Decordova, K. Swales, R. Riisnaes, L. Pope, K. Noguchi, R. Shiokawa, M. Inatani, J. Prince, K. Jones, C. Twelves, J. Spicer and U. Banerji, *Clin. Cancer Res.*, 2014, **20**, 5908–5917.
- 58 D. Mahadevan, E. G. Chiorean, W. B. Harris, D. D. Von Hoff, A. Stejskal-Barnett, W. Qi, S. P. Anthony, A. E. Younger, D. M. Rensvold, F. Cordova, C. F. Shelton, M. D. Becker, J. R. Garlich, D. L. Durden and R. K. Ramanathan, *Eur. J. Cancer*, 2012, **48**, 3319–3327.
- 59 F. Janku, *Cancer Treat. Rev.*, 2017, **59**, 93–101.
- 60 C. Y. Cheah and N. H. Fowler, *Blood*, 2016, **128**, 331–336.
- 61 B. W. Miller, D. Przepiorka, R. A. de Claro, K. Lee, L. Nie, N. Simpson, R. Gudi, H. Saber, S. Shord, J. Bullock, D. Marathe, N. Mehrotra, L. S. Hsieh, D. Ghosh, J. Brown, R. C. Kane, R. Justice, E. Kaminskis, A. T. Farrell and R. Pazdur, *Clin. Cancer Res.*, 2015, **21**, 1525–1529.
- 62 C. Fritsch, A. Huang, C. Chatenay-Rivauday, C. Schnell, A. Reddy, M. Liu, A. Kauffmann, D. Guthy, D. Erdmann, A. De Pover, P. Furet, H. Gao, S. Ferretti, Y. Wang, J. Trappe, S. M. Brachmann, S. M. Maira, C. Wilson, M. Boehm, C. Garcia-Echeverria, P. Chene, M. Wiesmann, R. Cozens, J. Lehar, R. Schlegel, G. Caravatti, F. Hofmann and W. R. Sellers, *Mol. Cancer Ther.*, 2014, **13**, 1117–1129.
- 63 I. A. Mayer, V. G. Abramson, L. Formisano, J. M. Balko, M. V. Estrada, M. E. Sanders, D. Juric, D. Solit, M. F. Berger, H. H. Won, Y. Li, L. C. Cantley, E. Winer and C. L. Arteaga, *Clin. Cancer Res.*, 2017, **23**, 26–34.
- 64 R. M. J. M. van Geel, J. Taberero, E. Elez, J. C. Bendell, A. Spreafico, M. Schuler, T. Yoshino, J. P. Delord, Y. Yamada, M. P. Lolkema, J. E. Faris, F. A. L. M. Eskens, S. Sharma, R. Yaeger, H. J. Lenz, Z. A. Wainberg, E. Avsar, A. Chatterjee, S. Jaeger, E. Tan, K. Maharry, T. Demuth and J. H. M. Schellens, *Cancer Discovery*, 2017, **7**, 610–619.
- 65 S. Jain, A. N. Shah, C. A. Santa-Maria, K. Siziopikou, A. Rademaker, I. Helenowski, M. Cristofanilli and W. J. Gradishar, *Breast Cancer Res. Treat.*, 2018, **171**, 371–381.
- 66 D. Juric, J. S. de Bono, P. M. LoRusso, J. Nemunaitis, E. I. Heath, E. L. Kwak, T. Macarulla Mercade, E. Geuna, M. Jose de Miguel-Luken, C. Patel, K. Kuida, S. Sankoh, E. H. Westin, F. Zohren, Y. Shou and J. Taberero, *Clin. Cancer Res.*, 2017, **23**, 5015–5023.
- 67 R. Arafah and Y. Samuels, *Semin. Cancer Biol.*, 2019, **59**, 36–49.
- 68 M. S. Miller, P. E. Thompson and S. B. Gabelli, *Biomolecules*, 2019, **9**, 82.
- 69 M. Frazzetto, C. Suphioglu, J. Zhu, O. Schmidt-Kittler, I. G. Jennings, S. L. Cranmer, S. P. Jackson, K. W. Kinzler, B. Vogelstein and P. E. Thompson, *Biochem. J.*, 2008, **414**, 383–390.
- 70 Z. Zheng, S. I. Amran, P. E. Thompson and I. G. Jennings, *Mol. Pharmacol.*, 2011, **80**, 657–664.
- 71 Z. A. Knight, B. Gonzalez, M. E. Feldman, E. R. Zunder, D. D. Goldenberg, O. Williams, R. Loewith, D. Stokoe, A. Balla, B. Toth, T. Balla, W. A. Weiss, R. L. Williams and K. M. Shokat, *Cell*, 2006, **125**, 733–747.
- 72 C. A. Evans, T. Liu, A. Lescarbeau, S. J. Nair, L. Grenier, J. A. Pradeilles, Q. Glenadel, T. Tibbitts, A. M. Rowley, J. P. DiNitto, E. E. Brophy, E. L. O'Hearn, J. A. Ali, D. G. Winkler, S. I. Goldstein, P. O'Hearn, C. M. Martin, J. G. Hoyt, J. R. Soglia, C. Cheung, M. M. Pink, J. L. Proctor, V. J. Palombella, M. R. Tremblay and A. C. Castro, *ACS Med. Chem. Lett.*, 2016, **7**, 862–867.
- 73 V. Certal, J. C. Carry, F. Halley, A. Virone-Oddos, F. Thompson, B. Filoche-Romme, Y. El-Ahmad, A. Karlsson, V. Charrier, C. Delorme, A. Rak, P. Y. Abecassis, C. Amara, L. Vincent, H. Bonnevaux, J. P. Nicolas, M. Mathieu, T. Bertrand, J. P. Marquette, N. Michot, T. Benard, M. A. Perrin, O. Lemaitre, S. Guerif, S. Perron, S. Monget, F. Gruss-Leleu, G. Doerflinger, H. Guizani, M. Brollo, L. Delbarre, L. Bertin, P. Richepin, V. Loyau, C. Garcia-Echeverria, C. Lengauer and L. Schio, *J. Med. Chem.*, 2014, **57**, 903–920.
- 74 Z. H. Zheng, M. S. Miller, I. G. Jennings and P. E. Thompson, *ACS Chem. Biol.*, 2013, **8**, 679–683.
- 75 A. Berndt, S. Miller, O. Williams, D. D. Le, B. T. Houseman, J. I. Pacold, F. Gorrec, W. C. Hon, Y. Liu, C. Rommel, P. Gaillard, T. Ruckle, M. K. Schwarz, K. M. Shokat, J. P. Shaw and R. L. Williams, *Nat. Chem. Biol.*, 2010, **6**, 117–124.
- 76 T. P. Heffron, B. Q. Wei, A. Olivero, S. T. Staben, V. Tsui, S. Do, J. Dotson, A. J. Folkes, P. Goldsmith, R. Goldsmith, J. Gunzner, J. Lesnick, C. Lewis, S. Mathieu, J. Nonomiya, S. Shuttleworth, D. P. Sutherland, N. C. Wan, S. M. Wang, C. Wiesmann and B. Y. Zhu, *J. Med. Chem.*, 2011, **54**, 7815–7833.
- 77 M. Nacht, L. X. Qiao, M. P. Sheets, T. St Martin, M. Labenski, H. Mazdiyasi, R. Karp, Z. D. Zhu,



- P. Chaturvedi, D. Bhavsar, D. Q. Niu, W. Westlin, R. C. Petter, A. P. Medikonda and J. Singh, *J. Med. Chem.*, 2013, **56**, 712–721.
- 78 S. T. Staben, C. Ndubaku, N. Blaquiere, M. Belvin, R. J. Bull, D. Dudley, K. Edgar, D. Gray, R. Heald, T. P. Heffron, G. E. Jones, M. Jones, A. Kolesnikov, L. Lee, J. Lesnick, C. Lewis, J. Murray, N. J. McLean, J. Nonomiya, A. G. Olivero, R. Ord, J. Pang, S. Price, W. W. Prior, L. Rouge, L. Salphati, D. Sampath, J. Wallin, L. Wang, B. Wei, C. Weismann and P. Wu, *Bioorg. Med. Chem. Lett.*, 2013, **23**, 2606–2613.
- 79 C. O. Ndubaku, T. P. Heffron, S. T. Staben, M. Baumgardner, N. Blaquiere, E. Bradley, R. Bull, S. Do, J. Dotson, D. Dudley, K. A. Edgar, L. S. Friedman, R. Goldsmith, R. A. Heald, A. Kolesnikov, L. Lee, C. Lewis, M. Nannini, J. Nonomiya, J. Pang, S. Price, W. W. Prior, L. Salphati, S. Sideris, J. J. Wallin, L. Wang, B. Wei, D. Sampath and A. G. Olivero, *J. Med. Chem.*, 2013, **56**, 4597–4610.
- 80 Z. H. Zheng, S. I. Amran, J. X. Zhu, O. Schmidt-Kittler, K. W. Kinzler, B. Vogelstein, P. R. Shepherd, P. E. Thompson and I. G. Jennings, *Biochem. J.*, 2012, **444**, 529–535.
- 81 P. Furet, V. Guagnano, R. A. Fairhurst, P. Imbach-Weese, I. Bruce, M. Knapp, C. Fritsch, F. Blasco, J. Blanz, R. Aichholz, J. Hamon, D. Fabbro and G. Caravatti, *Bioorg. Med. Chem. Lett.*, 2013, **23**, 3741–3748.
- 82 J. Wang, G. Q. Gong, Y. Zhou, W. J. Lee, C. M. Buchanan, W. A. Denny, G. W. Rewcastle, J. D. Kendall, J. M. J. Dickson, J. U. Flanagan, P. R. Shepherd, D. H. Yang and M. W. Wang, *Acta Pharmacol. Sin.*, 2018, **39**, 1816–1822.
- 83 P. Gkeka, A. Papafotika, S. Christoforidis and Z. Cournia, *J. Phys. Chem. B*, 2015, **119**, 1002–1016.
- 84 I. Galdadas, F. L. Gervasio and Z. Cournia, *Chem. Sci.*, 2020, **11**, 3511–3515.
- 85 H. Leontiadou, I. Galdadas, C. Athanasiou and Z. Cournia, *Sci. Rep.*, 2018, **8**, 15544.
- 86 A. T. Shaw, L. Friboulet, I. Leshchiner, J. F. Gainor, S. Bergqvist, A. Brooun, B. J. Burke, Y. L. Deng, W. Liu, L. Dardaie, R. L. Frias, K. R. Schultz, J. Logan, L. P. James, T. Smeal, S. Timofeevski, R. Katayama, A. J. Iafrate, L. Le, M. McTigue, G. Getz, T. W. Johnson and J. A. Engelman, *N. Engl. J. Med.*, 2016, **374**, 54–61.
- 87 S. Michels, M. Scheffler, S. Wagener, D. Plenker, A. Scheel, L. Nogova, A. Schultheis, R. N. Fischer, D. S. Y. Abdulla, R. Riedel, A. Bunck, C. Kobe, W. Baus, S. Merkelbach-Bruse, M. L. Sos, R. Buttner and J. Wolf, *JCO Precis Oncol.*, 2018, **2**, 1–6.
- 88 J. Rotow and T. G. Bivona, *Nat. Rev. Cancer*, 2017, **17**, 637–658.
- 89 F. J. Adrian, Q. Ding, T. B. Sim, A. Velentza, C. Sloan, Y. Liu, G. B. Zhang, W. Hur, S. Ding, P. Manley, J. Mestan, D. Fabbro and N. S. Gray, *Nat. Chem. Biol.*, 2006, **2**, 95–102.
- 90 J. M. Zhang, F. J. Adrian, W. Jahnke, S. W. Cowan-Jacob, A. G. Li, R. E. Iacob, T. Sim, J. Powers, C. Dierks, F. X. Sun, G. R. Guo, Q. Ding, B. Okram, Y. Choi, A. Wojciechowski, X. M. Deng, G. X. Liu, G. Fendrich, A. Strauss, N. Vajpai, S. Grzesiek, T. Tuntland, Y. Liu, B. Bursulaya, M. Azam, P. W. Manley, J. R. Engen, G. Q. Daley, M. Warmuth and N. S. Gray, *Nature*, 2010, **463**, 501–506.
- 91 A. A. Wylie, J. Schoepfer, W. Jahnke, S. W. Cowan-Jacob, A. Loo, P. Furet, A. L. Marzinzik, X. Pelle, J. Donovan, W. J. Zhu, S. Buonamici, A. Q. Hassan, F. Lombardo, V. Iyer, M. Palmer, G. Berellini, S. Dodd, S. Thohan, H. Bitter, S. Branford, D. M. Ross, T. P. Hughes, L. Petruzzelli, K. G. Vanasse, M. Warmuth, F. Hofmann, N. J. Keen and W. R. Sellers, *Nature*, 2017, **543**, 733–737.
- 92 J. Schoepfer, W. Jahnke, G. Berellini, S. Buonamici, S. Cotesta, S. W. Cowan-Jacob, S. Dodd, P. Drueckes, D. Fabbro, T. Gabriel, J. M. Groell, R. M. Grotzfeld, A. Q. Hassan, C. Henry, V. Iyer, D. Jones, F. Lombardo, A. Loo, P. W. Manley, X. Pelle, G. Rummel, B. Salem, M. Warmuth, A. A. Wylie, T. Zoller, A. L. Marzinzik and P. Furet, *J. Med. Chem.*, 2018, **61**, 8120–8135.
- 93 C. A. Eide, M. S. Zabriskie, S. L. S. Stevens, O. Antelope, N. A. Vellore, H. Than, A. R. Schultz, P. Clair, A. D. Bowler, A. D. Pomicter, D. Q. Yan, A. V. Senina, W. Qiang, T. W. Kelley, P. Szankasi, M. C. Heinrich, J. W. Tyner, D. Rea, J. M. Cayuela, D. W. Kim, C. E. Tognon, T. O'Hare, B. J. Druker and M. W. Deininger, *Cancer Cell*, 2019, **36**, 431–433.
- 94 X. Y. Lu, L. Yu, Z. Zhang, X. M. Ren, J. B. Smaill and K. Ding, *Med. Res. Rev.*, 2018, **38**, 1550–1581.
- 95 C. To, J. Jang, T. Chen, E. Park, M. Mushajiang, D. J. H. De Clercq, M. Xu, S. Wang, M. D. Cameron, D. E. Heppner, B. H. Shin, T. W. Gero, A. Yang, S. E. Dahlberg, K. K. Wong, M. J. Eck, N. S. Gray and P. A. Janne, *Cancer Discovery*, 2019, **9**, 926–943.
- 96 A. F. Abdel-Magid, *ACS Med. Chem. Lett.*, 2015, **6**, 104–107.
- 97 M. May, *Nat. Med.*, 2011, **17**, 405–407.
- 98 M. Filteau, V. Hamel, M. C. Pouliot, I. Gagnon-Arsenault, A. K. Dube and C. R. Landry, *Mol. Syst. Biol.*, 2015, **11**, 832.
- 99 M. Haskovic, B. Derks, L. van der Ploeg, J. Trommelen, J. Nyakayiru, L. J. C. van Loon, S. Mackinnon, W. W. Yue, R. W. A. Peake, L. Zha, D. Demirbas, W. S. Qi, X. P. Huang, G. T. Berry, J. Achten, J. Bierau, M. E. Rubio-Gozalbo and A. I. Coelho, *Orphanet J. Rare Dis.*, 2018, **13**, 212.
- 100 A. C. Joerger, H. C. Ang, D. B. Veprintsev, C. M. Blair and A. R. Fersht, *J. Biol. Chem.*, 2005, **280**, 16030–16037.
- 101 J. Liu and R. Nussinov, *Proc. Natl. Acad. Sci. U. S. A.*, 2008, **105**, 901–906.
- 102 Y. P. Pan, B. Y. Ma, A. J. Levine and R. Nussinov, *Biochemistry*, 2006, **45**, 3925–3933.
- 103 R. Nussinov and C. J. Tsai, *Trends Pharmacol. Sci.*, 2014, **35**, 256–264.
- 104 K. Song, Q. Li, W. Gao, S. Y. Lu, Q. C. Shen, X. Y. Liu, Y. Y. Wu, B. Q. Wang, H. W. Lin, G. Q. Chen and J. Zhang, *Nucleic Acids Res.*, 2019, **47**, W315–W321.
- 105 Z. W. Tan, W. V. Tee, E. Guarnera, L. Booth and I. N. Berezovsky, *Nucleic Acids Res.*, 2019, **47**, D265–D270.



- 106 D. B. Kokh, S. Richter, S. Henrich, P. Czodrowski, F. Rippmann and R. C. Wade, *J. Chem. Inf. Model.*, 2013, **53**, 1235–1252.
- 107 P. Cimermancic, P. Weinkam, T. J. Rettenmaier, L. Bichmann, D. A. Keedy, R. A. Woldeyes, D. Schneidman-Duhovny, O. N. Demerdash, J. C. Mitchell, J. A. Wells, J. S. Fraser and A. Sali, *J. Mol. Biol.*, 2016, **428**, 709–719.
- 108 S. A. Hollingsworth, B. Kelly, C. Valant, J. A. Michaelis, O. Mastromihalis, G. Thompson, A. J. Venkatakrisnan, S. Hertig, P. J. Scammells, P. M. Sexton, C. C. Felder, A. Christopoulos and R. O. Dror, *Nat. Commun.*, 2019, **10**, 3289.
- 109 H. S. Hayatshahi, E. Ahuactzin, P. Tao, S. Y. Wang and J. Liu, *J. Chem. Inf. Model.*, 2019, **59**, 4691–4705.
- 110 J. J. Guo and H. X. Zhou, *Chem. Rev.*, 2016, **116**, 6503–6515.
- 111 E. Guarnera and I. N. Berezovsky, *PLoS Comput. Biol.*, 2016, **12**, e1004678.
- 112 S. W. Lockless and R. Ranganathan, *Science*, 1999, **286**, 295–299.
- 113 C. Kaya, A. Armutlulu, S. Ekesan and T. Haliloglu, *Nucleic Acids Res.*, 2013, **41**, W249–W255.
- 114 R. Kalescky, J. Liu and P. Tao, *J. Phys. Chem. A*, 2015, **119**, 1689–1700.
- 115 A. T. VanWart, J. Eargle, Z. Luthey-Schulten and R. E. Amaro, *J. Chem. Theory Comput.*, 2012, **8**, 2949–2961.
- 116 S. Bowerman and J. Wereszczynski, *Methods Enzymol.*, 2016, **578**, 429–447.
- 117 G. La Sala, S. Decherchi, M. De Vivo and W. Rocchia, *ACS Cent. Sci.*, 2017, **3**, 949–960.
- 118 I. N. Berezovsky, *Biochim. Biophys. Acta, Proteins Proteomics*, 2013, **1834**, 830–835.
- 119 R. Nussinov and C. J. Tsai, *Annu. Rev. Pharmacol.*, 2015, **55**, 249–267.
- 120 W. K. Huang, R. Nussinov and J. Zhang, *Methods Mol. Biol.*, 2017, **1529**, 439–446.
- 121 Z. W. Tan, E. Guarnera, W. V. Tee and I. N. Berezovsky, *Nucleic Acids Res.*, 2020, DOI: 10.1093/nar/gkaa338.
- 122 W. V. Tee, E. Guarnera and I. N. Berezovsky, *J. Mol. Biol.*, 2019, **431**, 3933–3942.
- 123 H. Frauenfelder, S. G. Sligar and P. G. Wolynes, *Science*, 1991, **254**, 1598–1603.
- 124 B. Ma, S. Kumar, C. J. Tsai and R. Nussinov, *Protein Eng.*, 1999, **12**, 713–720.
- 125 K. Gunasekaran, B. Y. Ma and R. Nussinov, *Proteins*, 2004, **57**, 433–443.
- 126 J. Liu and R. Nussinov, *Proc. Natl. Acad. Sci. U. S. A.*, 2017, **114**, 7480–7482.

