# Discovery of pyrazolo-thieno[3,2-d]pyrimidinylamino-phenyl acetamides as type-II pan-tropomyosin receptor kinase (TRK) inhibitors: Design, synthesis, and biological evaluation 

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# European Journal of Medicinal Chemistry <br> Discovery of pyrozolo-thienopyrimidinylamino-phenyl acetamides as type-II pantropomyosin receptor kinase (TRK) inhibitors: design, synthesis, and biological evaluation <br> --Manuscript Draft-- 

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| Abstract: | Tropomyosin receptor kinase (TRK) represents an attractive oncology target for cancer therapy related to its critical role in cancer formation and progression. NTRK fusions are found to occur in $3.3 \%$ of lung cancers, $2.2 \%$ of colorectal cancers, $16.7 \%$ of thyroid cancers, $2.5 \%$ of glioblastomas, and $7.1 \%$ of pediatric gliomas. In this paper, we described the discovery of the type-II pan-TRK inhibitor 4c through the structurebased drug design strategy from the original hits 1 b and 2 b . Compound 4 c exhibited excellent in vitro TRKA, TRKB, and TRKC kinase inhibitory activity and antiproliferative activity against human colorectal carcinoma derived cell line KM12. In the $\mathrm{NCI}-60$ human cancer cell lines screen, compound 4 g demonstrated nearly $80 \%$ of growth inhibition for KM12, while only minimal inhibitory activity was observed for the remaining 59 cancer cell lines. Western blot analysis demonstrated that 4c and its urea cousin 4 k suppressed the TPM3-TRKA autophosphorylation at the concentrations of 100 nM and 10 nM , respectively. The work presented that 2-(4-(thieno[3,2- d ]pyrimidin-4-ylamino)phenyl)acetamides could serve as a novel scaffold for the discovery and development of type-II pan-TRK inhibitors for the treatment of TRK driven cancers. |
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31\% TRKA inhibition @ $20 \mu \mathrm{M}$ KM12 IC $_{50}=\mathbf{2 5} \boldsymbol{\mu} \mathrm{M}$


Lead

## Highlights:

- A series of novel thieno[3,2-d]pyrimidine type-II TRKA inhibitors were designed and synthesized.
- Representative compound $\mathbf{4 c}$ was discovered through structure-based drug design (SBDD) approach
- Compound 4c showed potent in vitro anti-proliferative activity.
- Compound 4c exhibited an effective blockage of TRKA autophosphorylation.

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Click here to access/download Supplementary Material - For Publication Online supporting information_20210108.pdf acetamides as type-II pan-tropomyosin receptor kinase (TRK) inhibitors: design, synthesis, and biological evaluation Wei Yan, ${ }^{1, \dagger}$ Lingtian Zhang, ${ }^{1, \dagger}$ Fengping Lv, ${ }^{1}$ Marialuisa Moccia, ${ }^{2}$ Francesca Carlomagno, ${ }^{2,3}$ Christophe Landry, ${ }^{4}$ Massimo Santoro, ${ }^{2}$ Fabien Gosselet, ${ }^{4}$ Brendan Frett, ${ }^{1}$ and Hong-yu Lil, ${ }^{1,}$
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Abstract

Tropomyosin receptor kinase (TRK) represents an attractive oncology target for cancer therapy related to its critical role in cancer formation and progression. NTRK fusions are found to occur in $3.3 \%$ of lung cancers, $2.2 \%$ of colorectal cancers, $16.7 \%$ of thyroid cancers, $2.5 \%$ of glioblastomas, and $7.1 \%$ of pediatric gliomas. In this paper, we described the discovery of the type-II pan-TRK inhibitor $\mathbf{4 c}$ through the structure-based drug design strategy from the original hits $\mathbf{1 b}$ and $\mathbf{2 b}$. Compound $\mathbf{4 c}$ exhibited excellent in vitro TRKA, TRKB, and TRKC kinase inhibitory activity and anti-proliferative activity against human colorectal carcinoma derived cell line KM12. In the NCI-60 human cancer cell lines screen, compound $\mathbf{4 g}$ demonstrated nearly $80 \%$ of growth inhibition for KM12, while only minimal inhibitory activity was observed for the remaining 59 cancer cell lines. Western blot analysis demonstrated that $\mathbf{4 c}$ and its urea cousin $\mathbf{4 k}$ suppressed the TPM3-TRKA autophosphorylation at the concentrations of 100 nM and 10 nM , respectively. The work presented that 2-(4-(thieno[3,2-d]pyrimidin-4-

Keywords: kinase inhibitor, TRK, type-II, colorectal cancer

## Introduction

The tropomyosin receptor kinase (TRK) family enzymes belong to the transmembrane receptor tyrosine kinases superfamily. The TRKs are primarily expressed in human neuronal tissues that regulate synaptic strength and plasticity in the mammalian nervous system.[1-2] The TRK family consists of three subtypes, and each is activated by a specific ligand: TRKA and TRKB are activated by nerve growth factor (NGF) and brainderived neurotrophic factor (BDNF), respectively, while TRKC is primarily turned on by neurotrophin 3 (NT3).[13] Upon ligand binding, TRKs become dimerized and autophosphorylated thus trigger downstream signal transduction pathways including RAS/mitogen-activated protein kinase (MAPK), phosphatidylinositol 3 kinase (PI3K)/protein kinase B (AKT), and phospholipase C-y (PLC-y)/PKC pathways, which consequently modulate cell growth, proliferation, survival, and differentiation.[4] Initially, the TRKA oncogene was discovered in colon cancer.[5-6] So far, NTRK fusions are found to occur in $3.3 \%$ of lung cancers, $2.2 \%$ of colorectal cancers, $16.7 \%$ of thyroid cancers, $2.5 \%$ of glioblastomas, and $7.1 \%$ of pediatric gliomas.[3] Within the TRK family, TRKA is the most commonly identified oncogene, which is found at a rate of approximately $7.4 \%$ across multiple tumor types ( $0.4 \%$ for TRKB and $3.4 \%$ for TRKA, respectively).[3, 7] Moreover, it has been found that TRKA also mediates the stimulation of early tumor progression.[8] Therefore, the blockade of TRKA signaling is considered as an attractive clinical approach for the treatment of cancers.


Larotrectinib


Entrectinib

Figure 1. Chemical structures of larotrectinib (Vitrakvi®®) and entrectinib (Rozlytrek®).

To date, only two TRKA inhibitors were approved by U.S. food and drug administration. Larotrectinib viecerso approved in November 2018 for solid tumors with NTRK gene fusions;[9] and entrectinib was approved in August 2019 for NTRK gene fusion-positive solid tumors or ROS1-positive non-small cell lung cancer.[10] According to the classification of Shokat et al., [11] these two drugs both belong to type-I kinase inhibitor. Type-I inhibitors bind to the kinase only at the highly conserved adenosine triphosphate (ATP) binding site in the "DFG-motif in" (active) conformation and are thus less selective or active. Distinct from type-I inhibitors, type-II inhibitors extend themselves into an allosteric hydrophobic pocket adjacent to the ATP binding site and bind to the kinase in the "DFG-motif out" (inactive) conformation, which makes type-II inhibitors more active; moreover, given that the allosteric pocket residues are much less conserved between kinases, type-II inhibitors are therefore more selective as well.

One concern in the development of TRK kinase inhibitors is the potential risk of central nervous system (CNS) adverse events, regarding to their critical roles in the CNS normal function.[12-15] Unwanted adverse events related to the drug exposure in the CNS had been reported.[16] Therefore, the development of peripherally restricted TRK inhibitors is a rational and feasible approach to minimize the potential CNS toxicity.[17] Because of their binding modes, type-II kinase inhibitors generally bear a larger molecular weight and are more polar than type-I inhibitors, and are thus considered to be less permeable through the blood-brain barrier (BBB).[18-19] For example, the classical type-II BCR-ABL inhibitor imatinib (Gleevec®) exhibited excellent efficacy and safety profile and low CNS exposure.[20] In this context, the development of type-II TRKA inhibitors is highly desired. Currently, several type-II TRK inhibitors have been reported, but the majority of them are in the early discovery stage.[3, 21-23] One of the most advanced type-II TRKA inhibitor CE-245677 was discontinued in Phase I clinical trials;[17] its successor PF-06273340 completed the Phase I clinical trial in 2014, but no further progress was reported ever since.[24] Therefore, the exploitation of novel chemical scaffolds for type-II TRKA inhibitors is highly demanded.

As a part of our continuous effort on the development of kinase inhibitors,[25-30] especially TRKA inhibitors,[31] we recently screened a small in-house kinase compounds library, and compounds $\mathbf{1 b}$ and $\mathbf{2 b}$ were found to exhibit moderate TRKA inhibitory activity. Although thienopyrimidines derivatives had been reported having a variety of biological activities including anti-parasitic activity[32]; Bertrand et al, disclosed the co-crystal
struo , e of EX429 with TRKA and TRKB,[33] but the medicinal chemistry effort towards the discovery arsolinpound remained unknown. Therefore, we hypothesized that our thienopyrimidine derivativescould serve earsors. a starting point for the discovery and development of novel TRK pan-inhibitors. From the initial hits $\mathbf{1 b} \mathbf{a n d} \mathbf{2 b}$, utilizing the structure-based drug design (SBDD) approach, a series of potent TRKA inhibitors was designed and synthesized. The representative lead $\mathbf{4 c}$ exhibited strong in vitro TRKA inhibition and anti-proliferative activity. In this paper, we report the design, synthesis, and biological evaluation of these compounds.

## Results and discussion

## Structure and activity relationship (SAR)

We quickly synthesized a few analogues of compounds $\mathbf{1 b}$ and $\mathbf{2 b}$ to explore the SAR of these two closely related scaffolds. Unfortunately, all compounds 1a-1c and 2a-2c were only moderately or weakly active in enzymatic and cellular assays (Tables $1 \& 2$ ). To understand the binding mode and further improve the potency, compounds $\mathbf{1 b}$ and $\mathbf{2 b}$ were docked to the TRKA protein (PDB ID: 6PL1). As shown in Figure 2A, compound 1b bound to the TRKA protein in a classical type-II manner: a $\pi-\pi$ interaction was found between the phenyl ring and Phe589, two hydrogen bonds with Glu560 and DFG-motif residue Asp668 were formed from the amide moiety, and the allosteric DFG pocket was occupied with the 3-(tert-butyl)-1-methyl-1H-pyrazole moiety; while in compound $\mathbf{2 b}$, a hydrogen bond from the N 1 of 4 -amino-thieno[3,2-d]pyrimidine was formed with the hinge region residue Met592 (Figure 2B). Compounds 1a-1c features with the direct carbon-carbon connection between the phenyl ring and the thieno-pyrimidine core. Although the structural feature had been proved to be favored in a few kinase inhibitors (e.g. JAK inhibitor ruxolitinib and BTK inhibitor ibrutinib), [34] based on the molecular docking result, we discovered that this structural type is too rigid to have a strong binding with the TRKA kinase. In compounds $\mathbf{2 a - 2} \mathbf{c}$, the phenyl ring was bridged to the thieno-pyrimidine core through a nitrogen atom, which allows a flexible binding. As shown in the superimposed structures of $\mathbf{1 b}$ and $\mathbf{2 b}$ (Figure 2C), the spatial distance between the phenyl rings of $\mathbf{1 b}$ and $\mathbf{2 b}$ was only $1.8 \AA$. Therefore, we envisioned that a hybrid of these two scaffolds would be able to retain the original crucial interactions with the TRKA protein while maintaining a flexible binding for the optimal affinity.
Entrectinib
${ }^{a} \mathrm{C}_{50}$ values are shown as the mean $\pm$ SD of two independent assays.
${ }^{\text {In }}$ Inhibition percentage values are shown as the average of two independent assays.

Table 2. Structure-activity relationship of series II compounds. ${ }^{a}$


II

| Compound | $\mathrm{R}^{2}$ | TRKA Inhibition | KM12 cell |
| :---: | :---: | :---: | :---: |
|  |  | $\%(@ 20 \mu \mathrm{M})^{b}$ | $\mathrm{IC}_{50}(\mu \mathrm{M})$ |
| $\mathbf{2 a}$ | O | 18.0 | $102.6 \pm 10.3$ |

2b
${ }^{a} C_{50}$ values are shown as the mean $\pm$ SD of two independent assays.
${ }^{\text {b }}$ Inhibition percentage values are shown as the average of two independent assays.


Figure 2. Proposed binding modes of $\mathbf{1 b}$ and $\mathbf{2 b}$ with TRKA. The protein structure of TRKA is shown as orange ribbon. The hinge region of TRKA is colored in violet, and the DFG-motif is colored in blue. Key residues are colored as follows: carbon grey, oxygen red, nitrogen blue, and hydrogen white. The hydrogen bonds are colored in black, and $\pi-\pi$ interaction is colored in teal. (A) The atoms of $\mathbf{1 b}$ are colored as follows: carbon green, oxygen red, nitrogen blue, sulfur yellow, and hydrogen white. (B) The atoms of $\mathbf{2 b}$ are colored as follows: carbon brown, nitrogen blue, sulfur yellow, fluorine cyan, and hydrogen white. (C) Superimposed structures of $\mathbf{1 b}$ and $\mathbf{2 b}$. Zoomed view on the phenyl rings.

Table 3. Structure-activity relationship of series III compounds. ${ }^{a}$


III

| Compound | $\mathrm{R}^{3}$ | TRKA | KM12 cell |
| :--- | :---: | :---: | :---: |
|  |  | $\mathrm{IC}_{50}(\mu \mathrm{M})$ | $\mathrm{IC} 50(\mu \mathrm{M})$ |

3e
${ }^{\text {a }} \mathrm{C}_{50}$ values are shown as the mean $\pm$ SD of two independent assays.

A series of compounds were then synthesized and evaluated. The SAR study showed that the nature of $R^{3}$ group in the allosteric pocket had significant impacts on the activity (Table 3). Similar to that observed in the compound 1 series, aliphatic group substituted derivatives ( $\mathbf{3}$ a and $\mathbf{3 b}$ ) were 2-40 fold less active than aromatic substituted derivatives (3c-f) in enzymatic assays, and the cellular potency was completely abolished. Among the aromatic substituted derivatives, the five-membered heterocyclic rings (3e-f) were more favored than substituted phenyls (3c-d). Moreover, the ethynyl derivative 3d was about 3-fold less active than its trifluoromethyl analogue 3c, probably due to a steric clash between the ethynyl fragment and the allosteric pocket residues. Compound $\mathbf{3 e}$ was docked with TRKA (Figure 3). As expected, the $\pi-\pi$ interaction between the phenyl acetamide fragment and the enzyme was retained, and a hydrogen bond was found between the hinge region residue Met592 and the N 1 atom of compound $3 \mathbf{e}$.


Figure 3. Proposed binding mode of compounds 3 e with TRKA. The protein structure of TRKA is shown as orange ribbon. The hinge region of TRKA is colored in violet, and the DFG-motif is colored in blue. Key residues are colored as follows: carbon grey, oxygen red, nitrogen blue, and hydrogen white. The hydrogen bonds are colored in black, and pi-pi interaction is colored in teal. The atoms of 3 e are colored as follows: carbon green, oxygen red, nitrogen blue, sulfur yellow, and hydrogen white.

Although compounds in the series III displayed excellent enzymatic potency in the low micromolar range, but the cellular potency was relatively weak. We envisioned that the low potency in cells could be due to poor aqueous solubility. As it was observed in the binding study of compound $3 \mathbf{e}$ with TRKA kinase (Figure 3), the thiophenyl moiety of thieno-pyrimidine orients toward to the solvent front. We hypothesized that the introduction of a polar substituent group to the thiophenyl moiety could improve aqueous solubility while maintaining good ligand-protein interactions. Therefore, a methyl-pyrazole group was introduced to the 6-position of the thienopyrimidine moiety. To our delight, the resulting compounds $\mathbf{4 a}$ and $\mathbf{4 b}$ displayed approximately 10 -fold increase in both enzymatic and cellular potencies than the parent compounds 3c and 3d (Table 3); strikingly, 4c demonstrated a 20 -fold improvement in enzymatic potency and 100 -fold improvement in cellular potency than the parent compound $\mathbf{3 e}$.

Encouraged by these promising results, we proceeded to synthesize more derivatives to further explore SAR in the allosteric pocket. As shown in Table 3, the replacement of methyl-pyrazole in $\mathbf{4 c}$ with thiazole (compound 4d) resulted in 20- and 150-fold decreases in enzymatic and cellular potencies, respectively. Compound $\mathbf{4 e}$ with morpholinoethyl attached to the pyrazole, exhibited a comparable enzymatic potency to $\mathbf{4 c}$; but was far less potent in the cellular assay. It can be explained with that the high polarity of compound $\mathbf{4 e}$ limits its cell membrane permeability. Several compounds bearing a phenyl ring ( $\mathbf{4 f}-\mathbf{h}$ ) showed comparable cellular
mer-g). When the terminal phenyl group was replaced with a benzylic group (4i), a 25 -fold drop in potency vasorm observed (vs 4a), indicating a potential steric clash with a surrounding residue; while a pyridine replacement (4j) resulted in a 70-fold decrease in enzymatic potency, probably due to an unfavored electrostatic interaction between pyridine and surrounding residues. Diaryl urea is an important pharmacophore in constructing type-II anticancer kinase inhibitors, exemplified by the success of sorafenib (Nexavar®). Therefore, compound 4k bearing a urea linker was prepared and evaluated, which demonstrated comparable activities in both enzymatic and cellular levels as its amide partner 4c.

Table 4. Structure-activity relationship of series IV compounds (part 1). ${ }^{\text {a }}$
4e
4f
$\mathrm{CH}_{2}$
 $0.092 \pm 0.054$
$1.02 \pm 0.16$
4g
$\mathrm{CH}_{2}$
 $0.094 \pm 0.016$ $0.09 \pm 0.01$
4h
$\mathrm{CH}_{2}$

$1.58 \pm 0.17$
$1.38 \pm 0.20$
$4 i$
$\mathrm{CH}_{2}$

$3.61 \pm 0.34$
>200
4j
$\mathrm{CH}_{2}$

$9.70 \pm 1.59$
$\mathrm{ND}^{b}$
4k
NH

$0.058 \pm 0.008$
$0.13 \pm 0.04$
Entrectinib --
--
$0.0027 \pm 0.0007$
$0.012 \pm 0.00002$
${ }^{\text {a }} \mathrm{C}_{50}$ values are shown as the mean $\pm$ SD of two independent assays.
${ }^{\mathrm{b}}$ Not determined.


Figure 4. Proposed binding mode of compounds $\mathbf{4 c}$ and $\mathbf{~} \mathbf{e}$ with TRKA. The protein structure of TRKA is shown as orange ribbon. The hinge region of TRKA is colored in violet, and the DFG-motif is colored in blue. Key residues are colored as follows: carbon grey, oxygen red, nitrogen blue, and hydrogen white. The hydrogen bonds are colored in black. (A) The atoms of $\mathbf{4 c}$ are colored as follows: carbon green, oxygen red, nitrogen blue, sulfur yellow, and hydrogen white. (B) The protein structure of TRKA is shown as an azure surface. (C)
.nposed structures of $\mathbf{4 c}$ and $\mathbf{4 e}$. The atoms of $\mathbf{4 e}$ are colored as follows: carbon brown, oxygen re jen blue, sulfur yellow, and hydrogen white. Zoomed view on the morpholine moiety of $\mathbf{4 e}$.

As illustrated in the proposed binding mode of $\mathbf{4 c}$ with the TRKA kinase (Figure 4A), all interactions observed between $\mathbf{3 e}$ and the TRKA enzyme were retained with compound $\mathbf{4 c}$, with the terminal pyrazole group laying at the solvent front (Figure 4B). To rationalize the excellent enzymatic potency, compound $\mathbf{4 e}$ was docked with the TRKA protein as well. Compound $\mathbf{4 e}$ nicely superimposed with $\mathbf{4 c}$; meanwhile, the morpholine moiety of $\mathbf{4 e}$ extended out of the allosteric hydrophobic pocket and was able to form a salt bridge with the carboxylic acid group of Glu560 or Asp668 (Figure 4C). This proposed binding mode explained its excellent enzymatic potency very well. The low potency could be explained with the low cell permeability of compound $\mathbf{4 e}$.

To explore the $S A R$ in the $R^{5}$ region, the methyl-pyrazole was replaced (Table 5). 4(Methylsulfonyl)phenyl group was established as a bioisosteric group of methyl-pyrazole which can avoid the formation of de-methylation metabolite.[28] However, the resulting compounds 41 and 4 m totally abolished the potency in the enzymatic assay. Although it is unexpected, the loss of the potency can be partially explained with potential steric clash between the bulky methylsulfonyl (4I) or trimethoxyl group (4m) with surrounding residues; because as observed in Figure 4B, the methyl-pyrazole of $\mathbf{4 c}$ was tightly surrounded by the solvent front residues. In line with the above speculation, a less hindered pyrrolidin-ethoxy-phenyl group was well tolerated and compound $\mathbf{4 n}$ displayed a comparable enzymatic and cellular potency to compound $\mathbf{4 h}$. The tolerability of a solubilizing group from the pyrazole moiety is an important SAR discovery which would enable the further optimization of solubility and drug properties.

Table 5. Structure-activity relationship of series IV compounds (part 2). ${ }^{\text {a }}$


IV

| Compound | $\mathrm{R}^{5}$ | TRKA | KM12 cell |
| :--- | :---: | :---: | :---: |
|  |  | $\mathrm{IC}_{50}(\mu \mathrm{M})$ | $\mathrm{IC}_{50}(\mu \mathrm{M})$ |

4h
${ }^{a} C_{50}$ values are shown as the mean $\pm$ SD of two independent assays.
${ }^{b}$ Not determined.

## Western Blot



Figure 5. Inhibition of TPM3-TRKA phosphorylation by $\mathbf{4 c}$, and $\mathbf{4 k}$ in human colorectal carcinoma derived cell line KM12. KM12 cell line were serum starved and treated for 2 hours with the indicated concentrations of the compounds. Total cell lysates ( $50 \mu \mathrm{~g}$ ) were subjected to immunoblotting with anti-phospho-TRKA (apY490 and apY4674/675) and anti-phospho-Tyrosine ( $\alpha \mathrm{pY}$ ) antibodies. The blots were normalized using anti-TRK (apanTRK) antibody.

In order to explore the ability of these new compounds to inhibit TRK-derived oncoproteins in cells, we analyzed their effects on the phosphorylation of TRKA using the human colorectal cancer derived cell line KM12 which endogenously harbours TPM3-TRKA rearrangement. As shown in Figure 5, compounds 4c and 4k inhibited the TPM3-TRKA autophosphorylation in a dose-dependent manner. Both compounds completely inhibited TPM3-TRKA autophosphorylation at the concentration of $100 \mathbf{n M}$; while $\mathbf{4 k}$ was able to significantly autophosphorylation with a relatively low concentration in cells, which is correlated with the anti-proliferative activity.

Table 6. Kinase selectivity profile of compound $\mathbf{4 c} .^{a}$

| Kinase | $\mathrm{IC}_{50}(\mu \mathrm{M})$ |
| :---: | :---: |
| TRKA | $0.075 \pm 0.001$ |
| TRKB | $0.074 \pm 0.007$ |
| TRKC | $0.055 \pm 0.014$ |
| Aurora A | $0.893 \pm 0.006$ |
| Aurora B | $1.99 \pm 0.24$ |
| CSF-1R | $13.6 \pm 0.3$ |
| MAP4K4 | $41.5 \pm 7.8$ |
| EGFR | $176.09 \pm 18.21$ |
| NEK2 | $>200$ |

${ }^{\text {a }} \mathrm{C}_{50}$ values are shown as the mean $\pm$ SD of two independent assays.


Figure 6. NCI-60 human cancer cell lines screen of compound $\mathbf{4 g}$.

## Selectivity profile

To assess the selectivity of our compounds, inhibitory activities of $\mathbf{4 c}$ and $\mathbf{~} \mathbf{g}$ were tested against a panel of kinases and $\mathrm{NCl}-60$ human cancer cell lines, respectively. As shown in Table 6, 4c exhibited pan-TRK inhibitory activity and was almost equal potent against TRKA-C, but was much less active against other kinases. Furthermore, as illustrated in Figure $6,4 \mathrm{~g}$ also exhibited excellent selectivity against TRKA-driven KM12 cells. At the drug concentration of $10 \mu \mathrm{M}, \mathbf{4 g}$ demonstrated nearly $80 \%$ of growth inhibition for KM12, while only minimal inhibitory activity was observed for the remaining 59 cancer cell lines. These data indicated our compounds are selective against TRK oncogenes at both enzymatic and cellular levels.

## Blog -brain barrier permeability



Figure 7. Transport assessment of entrectinib and $\mathbf{4 c}$ across in vitro BBB model.
To evaluate the BBB permeability, entrectinib and $\mathbf{4 c}$ were subjected to the in vitro BBB transport assay. As shown in Figure 7, the diffusion of entrectinib (permeability coefficient $(\mathrm{Pe})=1.02 \times 10^{-3} \mathrm{~cm} / \mathrm{min}$ ) across the BBB was observed in our human in vitro BBB model, which was in line with published data that entrectinib crossed the BBB in the mouse xenograft model.[35] Compound $\mathbf{4 c}\left(\mathrm{Pe}=3.03 \times 10^{-3} \mathrm{~cm} / \mathrm{min}\right)$ was more permeable than entrectinib, which was predicted from its lower molecular weight (500.6 Da vs 560.6 Da of entrectinib) and higher predicted LogP (5.15 vs 4.45 of entrectinib). We envisioned that by strategically increasing molecular weight and lowing LogP could lead to the discovery of derivatives having desirable cancer cell killing activity and low BBB permeability.

## Chemistry

As shown in Scheme 1, the key intermediates acids 7, 11, 14, and 15 were synthesized through either a telescope coupling-hydrolysis reaction ( $\mathbf{7}$ and 14) or a successive substitution-hydrolysis reaction (11 and 15). Final compounds 1, 3, and 4a-j were synthesized from the corresponding acids and amines using HATU as the acylation reagent in DMF at room temperature; amide 16 was synthesized from acid 15 following the same procedure, and then compound 16 was coupled with the corresponding boronic acids to give final compounds 4I-n. Compounds 12 and 17 were irradiated under microwave to produce 18, which underwent a successive coupling and de-protection procedure to generate amine 19. Urea compound $\mathbf{4 k}$ was prepared from aniline 19 and 3-(tert-butyl)-1-methyl-1 H-pyrazol-5-amine in the presence of triphosgene. To synthesize intermediate 24, $\mathbf{2 0}$ was reacted with an excessive amount of TsCl to give compound $\mathbf{2 1}$; compound 21 was heated in morpholine
rersonse the amino group was hindered by the bulky Ts groups. The Ts-protection group was removed by stirticiso to give compound $\mathbf{2 4}$ in $\mathrm{H}_{2} \mathrm{SO}_{4}$ at ambient temperature. Amine $\mathbf{2 4}$ was too hydrophilic to be extracted from water.

Therefore, the reaction mixture was concentrated after base quenching, and the resulting crude was used directly for next step without purification.


Scheme 1. Synthesis of final compounds. Reagents and conditions: (a) $\mathrm{Pd}\left(\mathrm{PPh}_{3}\right)_{4}, \mathrm{NaHCO}_{3}, \mathrm{THF}, \mathrm{H}_{2} \mathrm{O}$, microwave, $140^{\circ} \mathrm{C}, 0.5 \mathrm{~h}$; (b) LiOH, THF, $\mathrm{H}_{2} \mathrm{O}$, microwave, $80^{\circ} \mathrm{C}, 0.5 \mathrm{~h}$; (c) amines, HATU, Et N , DMF, R.T., overnight; (d) ${ }^{\circ} \mathrm{PrOH}$, microwave, $100^{\circ} \mathrm{C}, 0.5 \mathrm{~h}$.



Scheme 3. Synthesis of intermediate 24. Reagents and conditions: (a) $\mathrm{TsCl}^{2} \mathrm{Et}_{3} \mathrm{~N}, \mathrm{DCM}$, R.T. to reflux, overnight; (b) morpholine, $120^{\circ} \mathrm{C}$, overnight; (c) $\mathrm{H}_{2} \mathrm{SO}_{4}$, R.T., overnight.

## Conclusion

In summary, we have discovered a novel series of 2-(4-(thieno[3,2-d]pyrimidin-4ylamino)phenyl)acetamide derivatives as potent TRKA inhibitors. By utilizing the SBDD strategy, compound 4c was identified through the optimization of the starting hits $\mathbf{1}$ and $\mathbf{2}$. Compound $\mathbf{4 c}$ exhibited excellent in vitro TRKA, TRKB, and TRKC kinase inhibitory and anti-proliferative activities in the TRKA driven cell line. In the NCI60 human cancer cell lines screen, compound $\mathbf{4 g}$ demonstrated nearly $80 \%$ of growth inhibition for KM12, while only minimal inhibitory activity was observed for the remaining 59 cancer cell lines at $10 \mu \mathrm{M}$. Western blot analysis indicated that compounds $\mathbf{4 c}$ and $\mathbf{4 k}$ suppressed the TPM3-TRKA autophosphorylation at $10-100 \mathrm{nM}$ concentration, implying that the inhibitory activity in cells was derived from the inhibition of TRKA. Molecular modeling study demonstrated that this series of compounds bond to the TRKA kinase in a type-II manner. The work presented provided convincing evidence that 2-(4-(thieno[3,2-d]pyrimidin-4-ylamino)phenyl)acetamides could serve as a novel scaffold for the discovery and development of potent and selective type-II TRK inhibitors.

## Experimental Section

## Chemistry.

All starting materials and reagents were either obtained from commercial suppliers or prepared according to literature reported procedures. All purchased chemicals and solvents were used without further purification
mass spectrometry (LC-MS) (Thermo Surveyor HPLC System with Thermo Finnigan LCQ Deca). ${ }^{1} \mathrm{H}$ NMR spectral data were recorded on Varian 400 NMR spectrometer, and ${ }^{13} \mathrm{C}$ NMR was recorded on Varian Mercury 101 NMR spectrometer at ambient temperature. Chemicals shifts ( $\delta$ ) were reported in parts per million, coupling constants ( $\mathcal{J}$ ) values were in hertz, and the splitting patterns were abbreviated as follows: s for singlet; d for doublet; t for triplet; q for quartet; and m for multiplet. All final compounds for biological evaluation were purified to $\geq 95 \%$ purity as determined by Agilent 1100 HPLC instrument (column: Phenomenex kinetex $150 \times 4.6 \mathrm{~mm}, 5$ $\mu \mathrm{m}$; solvent $\mathrm{A}: \mathrm{H}_{2} \mathrm{O} / 0.1 \%$ TFA; solvent $\mathrm{B}: ~ \mathrm{MeOH} / 0.1 \%$ TFA; run time 12 min.; time/\%B: 0.0/30, 6.0/100, 7.0/100, 7.01/30,12.0/30; flow rate: $0.8 \mathrm{~mL} / \mathrm{min}$.; wavelength: 254 nm ).

General procedure A: Synthesis of compounds 1, 3, 4a-j, and 16. Acids such as 7, 11, 14, or 15 ( 0.14 mmol ), amine ( 0.16 mmol ), HATU ( 0.16 mmol ), and $\mathrm{Et}_{3} \mathrm{~N}(0.28 \mathrm{mmol})$ were dissolved in anhydrous DMF ( 1 mL ). The reaction mixture was stirred at ambient temperature overnight. The completion of the reaction was monitored by LC-MS analysis. The reaction mixture was poured into water and extracted with ethyl acetate. The organic phase was washed with water and brine twice, and concentrated in vacuum. The resulting residue was purified by silica gel chromatography to get the title compound.

N-((1R,2R)-2-hydroxycyclohexyl)-2-(4-(thieno[3,2-d]pyrimidin-4-yl)phenyl)acetamide (1a). The title compound was synthesized from 7 and ( $1 R, 2 R$ )-2-aminocyclohexanol hydrochloride following general procedure $A$. White solid (73\%). ${ }^{1} \mathrm{H}$ NMR (400 MHz, DMSO-d ${ }_{6}$ ) $\delta 9.25(\mathrm{~s}, 1 \mathrm{H}), 8.56(\mathrm{~d}, J=5.5 \mathrm{~Hz}, 1 \mathrm{H}), 8.12(\mathrm{~d}, J=7.9 \mathrm{~Hz}, 2 \mathrm{H})$, $7.81(\mathrm{~d}, J=8.0 \mathrm{~Hz}, 1 \mathrm{H}), 7.72(\mathrm{~d}, J=5.6 \mathrm{~Hz}, 1 \mathrm{H}), 7.54(\mathrm{~d}, J=8.0 \mathrm{~Hz}, 2 \mathrm{H}), 4.62(\mathrm{~d}, J=3.8 \mathrm{~Hz}, 1 \mathrm{H}), 3.74(\mathrm{~s}$, $1 \mathrm{H}), 3.71-3.62(\mathrm{~m}, 1 \mathrm{H}), 3.61(\mathrm{~s}, 2 \mathrm{H}), 1.71-1.49(\mathrm{~m}, 4 \mathrm{H}), 1.47-1.36(\mathrm{~m}, 2 \mathrm{H}), 1.33-1.17(\mathrm{~m}, 2 \mathrm{H}) .{ }^{13} \mathrm{C}$ NMR (101 MHz, DMSO- $d_{6}$ ) $\delta 168.86,161.87,158.81,154.58,140.37,138.43,134.71,129.79,128.07,127.15$, 124.44, 66.82, $50.87,42.09,31.69,26.63,23.76,19.57$. LC-MS (ESI) $m / z .[M+H]^{+} 368.1$. HPLC purity: $100 \%$, $t_{R}($ retention time $)=6.87 \mathrm{~min}$.

N-(3-(tert-butyl)-1-methyl-1H-pyrazol-5-yl)-2-(4-(thieno[3,2-d]pyrimidin-4-yl)phenyl)acetamide (1b). The title compound was synthesized from 7 and 3-(tert-butyl)-1-methyl-1H-pyrazol-5-amine following general procedure A. White solid (82\%). ${ }^{1} \mathrm{H}$ NMR ( $400 \mathrm{MHz}, ~ D M S O-d_{6}$ ) $\delta 10.15$ (s, 1H), 9.27 (s, 1H), 8.57 (d, J=5.5 Hz, 1H), 8.17 (d, J = 7.0 Hz, 2H), $7.73(\mathrm{~d}, J=5.5 \mathrm{~Hz}, 1 \mathrm{H}), 7.61(\mathrm{~d}, J=7.2 \mathrm{~Hz}, 2 \mathrm{H}), 6.08(\mathrm{~s}, 1 \mathrm{H}), 3.84(\mathrm{~s}, 2 \mathrm{H}), 3.61(\mathrm{~s}, 3 \mathrm{H})$, (ESI) $m / z:[\mathrm{M}+\mathrm{H}]^{+} 406.3$. HPLC purity: $100 \%, t_{\mathrm{R}}=7.53 \mathrm{~min}$.

N-(5-(tert-butyl)isoxazol-3-yl)-2-(4-(thieno[3,2-d]pyrimidin-4-yl)phenyl)acetamide (1c). The title compound was synthesized from 7 and 5-(tert-butyl)isoxazol-3-amine following general procedure A. White solid ( $85 \%$ ). ${ }^{1} \mathrm{H}$ NMR (400 MHz, DMSO- $d_{6}$ ) $\delta 11.28(\mathrm{~s}, 1 \mathrm{H}), 9.26(\mathrm{~d}, J=1.1 \mathrm{~Hz}, 1 \mathrm{H}), 8.57(\mathrm{~d}, J=5.6 \mathrm{~Hz}, 1 \mathrm{H}), 8.15(\mathrm{~d}, J=7.5$ $\mathrm{Hz}, 2 \mathrm{H}), 7.73(\mathrm{~d}, J=5.5 \mathrm{~Hz}, 1 \mathrm{H}), 7.59(\mathrm{~d}, J=7.8 \mathrm{~Hz}, 2 \mathrm{H}), 6.59(\mathrm{~s}, 1 \mathrm{H}), 3.83(\mathrm{~s}, 2 \mathrm{H}), 1.27(\mathrm{~s}, 9 \mathrm{H}) .{ }^{13} \mathrm{C}$ NMR ( $101 \mathrm{MHz}, \mathrm{DMSO}_{6}$ ) $\delta 180.47,168.94,161.90,158.71,157.87,154.59,138.67,138.50,135.21,130.00$ (2C), 128.27 (2C), 127.23, 124.45, 93.12, 42.26, 32.47, 28.29 (3C). LC-MS (ESI) $m / z:[M+H]^{+} 393.3$. HPLC purity: $97.9 \%, t_{\mathrm{R}}=8.02 \mathrm{~min}$.

N-((1S,2S)-2-hydroxycyclohexyl)-2-(4-(thieno[3,2-d]pyrimidin-4-ylamino)phenyl)acetamide (3a). The title compound was synthesized from 11 and (1R,2R)-2-aminocyclohexanol hydrochloride following general procedure A. Beige solid (67\%). ${ }^{1} \mathrm{H}$ NMR ( $400 \mathrm{MHz}, \mathrm{DMSO}-d_{6}$ ) $\delta 9.64$ (s, 1H), 8.55 (d, J=1.8 Hz, 1H), 8.21 $8.18(\mathrm{~m}, 1 \mathrm{H}), 7.68-7.64(\mathrm{~m}, 2 \mathrm{H}), 7.62(\mathrm{~d}, J=8.1 \mathrm{~Hz}, 1 \mathrm{H}), 7.45(\mathrm{~d}, J=5.4 \mathrm{~Hz}, 1 \mathrm{H}), 7.25(\mathrm{~d}, J=8.3 \mathrm{~Hz}, 2 \mathrm{H})$, $4.61-4.56(\mathrm{~m}, 1 \mathrm{H}), 3.71(\mathrm{~s}, 1 \mathrm{H}), 3.64(\mathrm{~s}, 1 \mathrm{H}), 3.45(\mathrm{~s}, 2 \mathrm{H}), 1.70-1.47(\mathrm{~m}, 4 \mathrm{~h}), 1.46-1.34(\mathrm{~m}, 2 \mathrm{H}), 1.31-$ 1.16 ( $\mathrm{m}, 2 \mathrm{H}$ ). ${ }^{13} \mathrm{C}$ NMR ( $101 \mathrm{MHz}, \mathrm{DMSO}_{6}$ $\mathrm{d}_{6} \delta 169.48,160.33,155.27,154.02,137.18,134.14,132.13$, 129.03 (2C), 124.34, 122.28 (2C), 115.28, 66.86, 50.77, 41.79, 39.52, 31.72, 26.64, 23.77, 19.56. LC-MS (ESI) $m / z:[M+H]^{+} 383.3$. HPLC purity: $100 \%, t_{R}=5.26 \mathrm{~min}$.

N-((1R,2S)-2-hydroxycyclohexyl)-2-(4-(thieno[3,2-d]pyrimidin-4-ylamino)phenyl)acetamide (3b). The title compound was synthesized from 11 and (1R,2S)-2-aminocyclohexanol following general procedure A. Beige solid (70\%). ${ }^{1} \mathrm{H}$ NMR (400 MHz, DMSO-d ${ }_{6}$ ) $\delta 9.64(\mathrm{~s}, 1 \mathrm{H}), 8.54(\mathrm{~s}, 1 \mathrm{H}), 8.23-8.18(\mathrm{~m}, 1 \mathrm{H}), 7.65(\mathrm{~d}, \mathrm{~J}=8.4$ $\mathrm{Hz}, 2 \mathrm{H}), 7.62(\mathrm{~d}, J=8.1 \mathrm{~Hz}, 1 \mathrm{H}), 7.45(\mathrm{~d}, J=5.4 \mathrm{~Hz}, 1 \mathrm{H}), 7.25(\mathrm{~d}, J=8.3 \mathrm{~Hz}, 2 \mathrm{H}), 4.59(\mathrm{~d}, J=3.9 \mathrm{~Hz}, 1 \mathrm{H})$, $3.71(\mathrm{~s}, 1 \mathrm{H}), 3.63(\mathrm{t}, \mathrm{J}=8.6 \mathrm{~Hz}, 1 \mathrm{H}), 3.44(\mathrm{~s}, 2 \mathrm{H}), 3.19-3.14(\mathrm{~m}, 1 \mathrm{H}), 1.70-1.18(\mathrm{~m}, 7 \mathrm{H}) .{ }^{13} \mathrm{C}$ NMR (101 MHz, DMSO- $d_{6}$ ) $\delta 169.47,160.33,155.27,154.02,137.18,134.15,132.14,129.02$ (2C), 124.35, 122.27 (2C), 115.27, 66.85, 50.76, 41.78, 31.72, 26.64, 23.76, 19.56. LC-MS (ESI) $m / z .[M+H]^{+} 383.4$. HPLC purity: $100 \%$, $t_{\mathrm{R}}=5.34 \mathrm{~min}$. NMR (400 MHz, DMSO-d $\mathrm{d}_{6}$ ס 10.50 (s, 1H), 9.69 (s, 1H), 8.56 (s, 1H), $8.21(\mathrm{~d}, J=5.4 \mathrm{~Hz}, 1 \mathrm{H}), 8.12(\mathrm{~s}, 1 \mathrm{H})$, $7.81(\mathrm{~d}, J=8.2 \mathrm{~Hz}, 1 \mathrm{H}), 7.72(\mathrm{~d}, J=8.2 \mathrm{~Hz}, 2 \mathrm{H}), 7.55(\mathrm{t}, J=8.0 \mathrm{~Hz}, 1 \mathrm{H}), 7.46(\mathrm{~d}, J=5.4 \mathrm{~Hz}, 1 \mathrm{H}), 7.39(\mathrm{~d}, J=$
 153.99, 139.95, 137.66, 134.19, 130.76, 129.96, $129.44(q, J=32 H z), 129.21(2 C), 124.09(q, J=272 \mathrm{~Hz})$, 124.34, 122.60, 122.31 (2C), 119.52 ( $q, J=4 \mathrm{~Hz}$ ), 115.37, 115.11 ( $\mathrm{q}, J=4 \mathrm{~Hz}$ ), 42.78. LC-MS (ESI) $\mathrm{m} / \mathrm{z}$ : $[M+H]^{+}$429.0. HPLC purity: $99.5 \%, t_{R}=6.72 \mathrm{~min}$.
$N$-(3-ethynylphenyl)-2-(4-(thieno[3,2-d]pyrimidin-4-ylamino)phenyl)acetamide (3d). The title compound was synthesized from 11 and 3-ethynylaniline following general procedure A. Beige solid (83\%). ${ }^{1} \mathrm{H}$ NMR (400 $\left.\mathrm{MHz}, \mathrm{DMSO}-\mathrm{d}_{6}\right) \delta 10.27(\mathrm{~s}, 1 \mathrm{H}), 9.69(\mathrm{~s}, 1 \mathrm{H}), 8.56(\mathrm{~s}, 1 \mathrm{H}), 8.20(\mathrm{~d}, J=5.4 \mathrm{~Hz}, 1 \mathrm{H}), 7.82(\mathrm{~s}, 1 \mathrm{H}), 7.72(\mathrm{~d}, J=$ $8.3 \mathrm{~Hz}, 2 \mathrm{H}), 7.59(\mathrm{~d}, J=8.3 \mathrm{~Hz}, 1 \mathrm{H}), 7.46(\mathrm{~d}, J=5.4 \mathrm{~Hz}, 1 \mathrm{H}), 7.37-7.28(\mathrm{~m}, 3 \mathrm{H}), 7.15(\mathrm{~d}, J=7.6 \mathrm{~Hz}, 1 \mathrm{H})$, $4.16(\mathrm{~s}, 1 \mathrm{H}), 3.65(\mathrm{~s}, 2 \mathrm{H}) .{ }^{13} \mathrm{C}$ NMR (101 MHz, DMSO-d ) $\delta$ 169.51, 160.34, 155.24, 154.00, 139.41, 137.59, 134.17, 130.97, 129.18, 129.16 (2C), 126.43, 124.35, 122.32 (2C), 122.00, 121.94, 119.69, 115.35, 83.35 , 80.51, 42.79. LC-MS (ESI) $m / z:[M+H]^{+} 385.1$. HPLC purity: $99.2 \%, t_{R}=6.18 \mathrm{~min}$.

N-(3-(tert-butyl)-1-methyl-1H-pyrazol-5-yl)-2-(4-(thieno[3,2-d]pyrimidin-4-ylamino)phenyl)acetamide (3e). The title compound was synthesized from 11 and 3-(tert-butyl)-1-methyl-1H-pyrazol-5-amine following general procedure A. Beige solid (72\%). ${ }^{1} \mathrm{H}$ NMR ( $400 \mathrm{MHz}, \mathrm{DMSO}_{6}$ ) $\delta 10.03(\mathrm{~s}, 1 \mathrm{H}), 9.68(\mathrm{~s}, 1 \mathrm{H}), 8.56(\mathrm{~s}, 1 \mathrm{H}), 8.21$ (d, $J=5.4 \mathrm{~Hz}, 1 \mathrm{H}), 7.72(\mathrm{~d}, J=7.9 \mathrm{~Hz}, 2 \mathrm{H}), 7.46(\mathrm{~d}, J=5.4 \mathrm{~Hz}, 1 \mathrm{H}), 7.33(\mathrm{~d}, J=8.0 \mathrm{~Hz}, 2 \mathrm{H}), 6.06(\mathrm{~s}, 1 \mathrm{H})$, 3.66 ( $\mathrm{s}, 2 \mathrm{H}$ ), $3.58(\mathrm{~s}, 3 \mathrm{H}), 1.19(\mathrm{~s}, 9 \mathrm{H}) .{ }^{13} \mathrm{C}$ NMR (101 MHz, DMSO-d $\mathrm{d}_{6}$ ) $\delta 169.10,160.37,158.60,155.24$, 154.01, 137.62, 136.27, 134.19, 130.85, 129.15 (2C), 124.36, 122.30 (2C), 115.36, 95.02, 41.64, 35.27, 31.78, 30.32 (3C). LC-MS (ESI) $m / z:[M+H]^{+} 421.2$. HPLC purity: $100 \%, t_{\mathrm{R}}=6.15 \mathrm{~min}$.

N-(5-(tert-butyl)isoxazol-3-yl)-2-(4-(thieno[3,2-d]pyrimidin-4-ylamino)phenyl)acetamide (3f). The title compound was synthesized from 11 and 5-(tert-butyl)isoxazol-3-amine following general procedure A. Beige solid (69\%). ${ }^{1} \mathrm{H}$ NMR (400 MHz, DMSO-d $\mathrm{d}_{6}$ ס 11.17 (s, 1H), 9.68 (s, 1H), 8.55 (s, 1H), 8.21 (dd, J=5.4, 1.3 Hz, 1H), 7.70 (d, $J=7.4 \mathrm{~Hz}, 2 \mathrm{H}), 7.45(\mathrm{dd}, J=5.4,1.3 \mathrm{~Hz}, 1 \mathrm{H}), 7.30(\mathrm{~d}, J=7.7 \mathrm{~Hz}, 2 \mathrm{H}), 6.58(\mathrm{~s}, 1 \mathrm{H}), 3.65(\mathrm{~s}, 2 \mathrm{H}), 1.27$ (s, 9H). ${ }^{13} \mathrm{C}$ NMR (101 MHz, $\mathrm{DMSO}_{6}$ ( $\mathrm{C}_{6}$ ס 180.40, 169.54, 160.36, 157.94, 155.22, 153.99, 137.68, 134.19,

## 2-(4-((6-(1-methyl-1H-pyrazol-4-yl)thieno[3,2-d]pyrimidin-4-yl)amino)phenyl)-N-(3-

(trifluoromethyl)phenyl)acetamide (4a). The title compound was synthesized from 14 and 3(trifluoromethyl)aniline following general procedure A. Beige solid ( $62 \%$ ). ${ }^{1} \mathrm{H}$ NMR ( $400 \mathrm{MHz}, \mathrm{DMSO}-d_{6}$ ) $\delta$ $10.50(\mathrm{~s}, 1 \mathrm{H}), 9.60(\mathrm{~s}, 1 \mathrm{H}), 8.51(\mathrm{~s}, 1 \mathrm{H}), 8.28(\mathrm{~s}, 1 \mathrm{H}), 8.12(\mathrm{~s}, 1 \mathrm{H}), 7.96(\mathrm{~s}, 1 \mathrm{H}), 7.81(\mathrm{~d}, J=8.3 \mathrm{~Hz}, 1 \mathrm{H}), 7.71$ (d, $J=8.3 \mathrm{~Hz}, 2 \mathrm{H}), 7.57-7.52(\mathrm{~m}, 2 \mathrm{H}), 7.39(\mathrm{~d}, J=7.8 \mathrm{~Hz}, 1 \mathrm{H}), 7.33(\mathrm{~d}, J=8.3 \mathrm{~Hz}, 2 \mathrm{H}), 3.91(\mathrm{~s}, 3 \mathrm{H}), 3.67(\mathrm{~s}$, 2H). ${ }^{13} \mathrm{C}$ NMR (101 MHz, DMSO- $d_{6}$ ) $\delta$ 169.88, 161.21, 154.56, 154.29, 141.85, 139.96, 137.79, 136.90, $130.56,129.97,129.44(q, J=32 H z), 129.22(2 C), 129.18,124.09(q, J=272 H z), 122.61,122.08(2 C)$, $119.52(q, J=4 H z), 118.22,115.31,115.11(q, J=4 H z), 113.20,54.89,42.79$. LC-MS (ESI) $m / z:[M+H]^{+}$ 509.2. HPLC purity: $99.4 \%, t_{\mathrm{R}}=6.96 \mathrm{~min}$. HRMS (ESI) m/z: $[\mathrm{M}+\mathrm{H}]^{+}$calcd for $\mathrm{C}_{25} \mathrm{H}_{20} \mathrm{~F}_{3} \mathrm{~N}_{6} \mathrm{OS} 509.1371$; found 509.1354.

N-(3-ethynylphenyl)-2-(4-((6-(1-methyl-1H-pyrazol-4-yl)thieno[3,2-d]pyrimidin-4-yl)amino)phenyl)acetamide (4b). The title compound was synthesized from 14 and 3-ethynylaniline following general procedure A. Beige solid (73\%). ${ }^{1} \mathrm{H}$ NMR (400 MHz, DMSO-d ${ }^{2}$ ) $\delta 10.26(\mathrm{~s}, 1 \mathrm{H}), 9.59(\mathrm{~s}, 1 \mathrm{H}), 8.51(\mathrm{~s}, 1 \mathrm{H}), 8.28(\mathrm{~s}, 1 \mathrm{H}), 7.96(\mathrm{~s}$, $1 \mathrm{H}), 7.81(\mathrm{~s}, 1 \mathrm{H}), 7.70(\mathrm{~d}, J=7.6 \mathrm{~Hz}, 2 \mathrm{H}), 7.59(\mathrm{~d}, J=7.6 \mathrm{~Hz}, 1 \mathrm{H}), 7.54(\mathrm{~d}, J=1.1 \mathrm{~Hz}, 1 \mathrm{H}), 7.31(\mathrm{dd}, J=7.4$, $5.1 \mathrm{~Hz}, 3 \mathrm{H}), 7.17-7.12(\mathrm{~m}, 1 \mathrm{H}), 4.16(\mathrm{~s}, 1 \mathrm{H}), 3.91(\mathrm{~s}, 3 \mathrm{H}), 3.64(\mathrm{~s}, 2 \mathrm{H}) .{ }^{13} \mathrm{C}$ NMR ( 101 MHz, DMSO-d $) ~ \delta$ $169.53,161.21,154.57,154.30,141.83,139.42,137.72,136.90,130.78,129.19,129.16$ (2C), 126.42, 122.09 (2C), 121.99, 121.93, 119.68, 118.21, 115.31, 113.18, 83.35, 80.51, 42.79, 38.83. LC-MS (ESI) m/z: $[M+H]^{+}$ 465.2. HPLC purity: $98.8 \%, t_{R}=6.54$ min. $\mathrm{HRMS}(E S I) \mathrm{m} / \mathrm{z}:[\mathrm{M}+\mathrm{H}]^{+}$calcd for $\mathrm{C}_{26} \mathrm{H}_{21} \mathrm{~N}_{6} \mathrm{OS} 465.1498$; found 465.1491.

N-(3-(tert-butyl)-1-methyl-1H-pyrazol-5-yl)-2-(4-((6-(1-methyl-1H-pyrazol-4-yl)thieno[3,2-d]pyrimidin-4yl)amino)phenyl)acetamide (4c). The title compound was synthesized from 14 and 3-(tert-butyl)-1-methyl-1H-pyrazol-5-amine following general procedure A. White solid (58\%). ${ }^{1} \mathrm{H}$ NMR ( $400 \mathrm{MHz}, \mathrm{DMSO}-d_{6}$ ) $\delta 10.03$ (s, $1 \mathrm{H}), 9.60(\mathrm{~s}, 1 \mathrm{H}), 8.52(\mathrm{~s}, 1 \mathrm{H}), 8.29(\mathrm{~s}, 1 \mathrm{H}), 7.96(\mathrm{~s}, 1 \mathrm{H}), 7.70(\mathrm{~d}, J=7.7 \mathrm{~Hz}, 2 \mathrm{H}), 7.54(\mathrm{~s}, 1 \mathrm{H}), 7.32(\mathrm{~d}, J=7.9$ Hz, 2H), 6.06 (s, 1H), 3.91 (s, 3H), 3.66 (s, 2H), 3.58 (s, 3H), 1.19 (s, 9H). ${ }^{13} \mathrm{C}$ NMR (101 MHz, DMSO-d ${ }_{6}$ ) $169.10,161.13,158.59,154.57,154.26,141.88,137.70,136.90,136.26,130.69,129.18,129.15$ (2C), 122.10
(2C) s $18.16,115.28,113.16,95.01,41.62,38.82,35.27,31.77,30.31$ (3C). LC-MS (ESI) $\mathrm{m} / \mathrm{z}:[\mathrm{M}+\mathrm{H}]^{+}$ C purity: $100 \%, t_{R}=7.02 \mathrm{~min}$. HRMS (ESI) m/z: $[\mathrm{M}+\mathrm{H}]^{+}$calcd for $\mathrm{C}_{26} \mathrm{H}_{29} \mathrm{~N}_{8} \mathrm{OS} 501.2185$; found 501

N-(4-(tert-butyl)thiazol-2-yl)-2-(4-((6-(1-methyl-1H-pyrazol-4-yl)thieno[3,2-d]pyrimidin-4-
$y$ l)amino)phenyl)acetamide (4d). The title compound was synthesized from 14 and 4-(tert-butyl)thiazol-2-amine following general procedure A. Yellow solid ( $62 \%$ ). ${ }^{1} \mathrm{H}$ NMR ( $400 \mathrm{MHz}, \mathrm{DMSO}-d_{6}$ ) $\delta 12.30(\mathrm{~s}, 1 \mathrm{H}), 9.59(\mathrm{~s}, 1 \mathrm{H})$, $8.51(\mathrm{~s}, 1 \mathrm{H}), 8.28(\mathrm{~s}, 1 \mathrm{H}), 7.96(\mathrm{~s}, 1 \mathrm{H}), 7.69(\mathrm{~d}, J=8.3 \mathrm{~Hz}, 2 \mathrm{H}), 7.54(\mathrm{~s}, 1 \mathrm{H}), 7.31(\mathrm{~d}, J=8.3 \mathrm{~Hz}, 2 \mathrm{H}), 3.91(\mathrm{~s}$, 3 H ), $3.73(\mathrm{~s}, 2 \mathrm{H}), 1.26(\mathrm{~s}, 9 \mathrm{H}) .{ }^{13} \mathrm{C}$ NMR ( 101 MHz , DMSO- $\mathrm{d}_{6}$ ) $\delta 169.24,161.21,160.12,157.15,154.54$, $154.28,141.85,137.88,136.89,130.06,129.25$ (2C), 129.17, 122.11 (2C), 118.20, 115.29, 113.18, 104.60, 41.16, 38.82, 34.10, 29.75 (3C). LC-MS (ESI) $m / z:[M+H]^{+} 504.1$. HPLC purity: $95.7 \%, t_{R}=7.28 \mathrm{~min}$. HRMS (ESI) $m / z:[M+H]^{+}$calcd for $\mathrm{C}_{25} \mathrm{H}_{26} \mathrm{~N}_{7} \mathrm{OS}_{2}$ 504.1640; found 504.1635.

N-(3-(tert-butyl)-1-(2-morpholinoethyl)-1H-pyrazol-5-yl)-2-(4-((6-(1-methyl-1H-pyrazol-4-yl)thieno[3,2-dlpyrimidin-4-yl)amino)phenyl)acetamide (4e). The title compound was synthesized from 14 and 25 following general procedure A. Beige solid (55\%). ${ }^{1} \mathrm{H}$ NMR (400 MHz, DMSO-d ${ }_{6}$ ) $\delta 10.26$ (s, 1H), 9.79 (s, 1H), 8.71 (s, $1 \mathrm{H}), 8.48(\mathrm{~s}, 1 \mathrm{H}), 8.16(\mathrm{~s}, 1 \mathrm{H}), 7.93(\mathrm{~d}, J=8.3 \mathrm{~Hz}, 2 \mathrm{H}), 7.74(\mathrm{~s}, 1 \mathrm{H}), 7.52(\mathrm{~d}, J=8.3 \mathrm{~Hz}, 2 \mathrm{H}), 6.26(\mathrm{~s}, 1 \mathrm{H}), 4.18$ (t, $J=7.0 \mathrm{~Hz}, 2 \mathrm{H}), 4.10(\mathrm{~s}, 3 \mathrm{H}), 3.85(\mathrm{~s}, 2 \mathrm{H}), 3.74-3.65(\mathrm{~m}, 4 \mathrm{~h}), 3.52(\mathrm{~s}, 2 \mathrm{H}), 2.75(\mathrm{t}, J=7.0 \mathrm{~Hz}, 2 \mathrm{H}), 2.70(\mathrm{~s}$, 2 H ), 1.39 ( $\mathrm{s}, 9 \mathrm{H}$ ). ${ }^{13} \mathrm{C}$ NMR ( $101 \mathrm{MHz}, \mathrm{DMSO}^{-} \mathrm{d}_{6}$ ) $\delta$ 169.16, 161.24, 158.93, 154.51, 154.28, 141.84, 137.86, 136.90, 135.94, 130.53, 129.18, 129.08 (2C), 121.87 (2C), 118.22, 115.29, 113.21, 95.38, 66.09 (2C), 57.29 , 53.19 (2C), 45.23, 41.83, 38.82, 31.81, 30.30 (3C). LC-MS (ESI) $m / z:[M+H]^{+} 600.4$. HPLC purity: $100 \%, t_{R}=$ 6.00 min . HRMS (ESI) $\mathrm{m} / \mathrm{z}:[\mathrm{M}+\mathrm{H}]^{+}$calcd for $\mathrm{C}_{31} \mathrm{H}_{38} \mathrm{~N}_{9} \mathrm{O}_{2} \mathrm{~S} 600.2869$; found 600.2854 .

N-(2,3-dihydrobenzo[b][1,4]dioxin-6-yl)-2-(4-((6-(1-methyl-1H-pyrazol-4-yl)thieno[3,2-d]pyrimidin-4$y$ l)amino)phenyl)acetamide (4f). The title compound was synthesized from 14 and 2,3-dihydrobenzo[b][1,4]dioxin-6-amine following general procedure A. White solid (69\%). ${ }^{1} \mathrm{H}$ NMR ( 400 MHz , DMSO-d $\mathrm{d}_{6}$ ס $9.98(\mathrm{~s}, 1 \mathrm{H}), 9.58(\mathrm{~s}, 1 \mathrm{H}), 8.51(\mathrm{~s}, 1 \mathrm{H}), 8.28(\mathrm{~s}, 1 \mathrm{H}), 7.96(\mathrm{~s}, 1 \mathrm{H}), 7.68(\mathrm{~d}, \mathrm{~J}=8.2 \mathrm{~Hz}, 2 \mathrm{H}), 7.53(\mathrm{~s}$, $1 \mathrm{H}), 7.30(\mathrm{~d}, J=8.2 \mathrm{~Hz}, 2 \mathrm{H}), 7.25(\mathrm{~d}, J=1.8 \mathrm{~Hz}, 1 \mathrm{H}), 6.98(\mathrm{dd}, J=8.7,1.8 \mathrm{~Hz}, 1 \mathrm{H}), 6.77(\mathrm{~d}, J=8.7 \mathrm{~Hz}, 1 \mathrm{H})$, 4.19 (dd, $J=8.9,4.5 \mathrm{~Hz}, 4 \mathrm{~h}$ ), $3.91(\mathrm{~s}, 3 \mathrm{H}), 3.58(\mathrm{~s}, 2 \mathrm{H}) .{ }^{13} \mathrm{C}$ NMR ( $101 \mathrm{MHz}, \mathrm{DMSO}-\mathrm{d}_{6}$ ) $\delta 168.75,161.19$, $154.59,154.30,142.87,141.83,139.22,137.60,136.90,132.94,131.16,129.16,129.11$ (2C), 122.12 (2C), 118.20, 116.71, 115.30, 113.15, 112.33, 108.23, 64.16, 63.89, 42.73, 38.82. LC-MS (ESI) $m / z:[M+H]^{+} 499.2$.

N-(4-(tert-butyl)phenyl)-2-(4-((6-(1-methyl-1H-pyrazol-4-yl)thieno[3,2-d]pyrimidin-4-yl)amino)phenyl)acetamide $(4 g)$. The title compound was synthesized from 14 and 4-(tert-butyl)aniline following general procedure A. White solid (73\%). ${ }^{1} \mathrm{H}$ NMR ( $\left.400 \mathrm{MHz}, ~ D M S O-d_{6}\right) \delta 10.08(\mathrm{~s}, 1 \mathrm{H}), 9.59(\mathrm{~s}, 1 \mathrm{H}), 8.51(\mathrm{~s}, 1 \mathrm{H}), 8.28(\mathrm{~s}, 1 \mathrm{H}), 7.95$ (s, 1H), $7.69(\mathrm{~d}, J=7.9 \mathrm{~Hz}, 2 \mathrm{H}), 7.56-7.48(\mathrm{~m}, 3 \mathrm{H}), 7.32(\mathrm{~d}, J=5.8 \mathrm{~Hz}, 2 \mathrm{H}), 7.30(\mathrm{~d}, J=5.9 \mathrm{~Hz}, 2 \mathrm{H}), 3.91(\mathrm{~s}$, 3 H ), $3.61(\mathrm{~s}, 2 \mathrm{H}), 1.25(\mathrm{~s}, 9 \mathrm{H}) .{ }^{13} \mathrm{C}$ NMR ( $101 \mathrm{MHz}, \mathrm{DMSO}^{2}-\mathrm{d}_{6}$ ) $\delta 169.00,161.13,154.59,154.27,145.47$, $141.85,137.59,136.89,136.66,131.21,129.17,129.10$ (2C), 125.26 (2C), 122.13 (2C), 118.91 (2C), 118.17, 115.30, 113.15, 42.76, 38.82, 33.96, 31.17 (3C). LC-MS (ESI) $m / z:[M+H]^{+} 497.2$. HPLC purity: $99.1 \%, t_{R}=$ 7.35 min. HRMS (ESI) $m / z:[M+H]^{+}$calcd for $\mathrm{C}_{28} \mathrm{H}_{29} \mathrm{~N}_{6} \mathrm{OS} 497.2124$; found 497.2100.

N-(3-chloro-4-fluorophenyl)-2-(4-((6-(1-methyl-1H-pyrazol-4-yl)thieno[3,2-d]pyrimidin-4-
$y l() a m i n o)$ phenyl)acetamide (4h). The title compound was synthesized from 14 and 3-chloro-4-fluoroaniline following general procedure A. Beige solid (70\%). ${ }^{1} \mathrm{H}$ NMR (400 MHz, DMSO-d ${ }_{6}$ ) $\overline{10.37}(\mathrm{~s}, 1 \mathrm{H}), 9.59(\mathrm{~s}, 1 \mathrm{H})$, $8.51(\mathrm{~s}, 1 \mathrm{H}), 8.29(\mathrm{~s}, 1 \mathrm{H}), 7.98-7.92(\mathrm{~m}, 2 \mathrm{H}), 7.70(\mathrm{~d}, J=8.4 \mathrm{~Hz}, 2 \mathrm{H}), 7.54(\mathrm{~s}, 1 \mathrm{H}), 7.52-7.46(\mathrm{~m}, 1 \mathrm{H}), 7.37$ ( $\mathrm{t}, J=9.1 \mathrm{~Hz}, 1 \mathrm{H}$ ), $7.31(\mathrm{~d}, J=8.4 \mathrm{~Hz}, 2 \mathrm{H}), 3.91(\mathrm{~s}, 3 \mathrm{H}), 3.63(\mathrm{~s}, 2 \mathrm{H}) .{ }^{13} \mathrm{C}$ NMR (101 MHz, DMSO-d $\left.\mathrm{d}_{6}\right)$ ס 169.52, 161.23, 154.55, 154.30, $152.99(J=243 \mathrm{~Hz}), 141.83,137.76,136.90,136.45(J=3 \mathrm{~Hz}), 130.58,129.17(2 \mathrm{C})$, $122.07(2 \mathrm{C}), 120.45,119.36(J=7 \mathrm{~Hz}), 119.07(J=18 \mathrm{~Hz}), 118.22,116.92(J=21 \mathrm{~Hz}), 115.29,113.17,42.68$, 38.83. LC-MS (ESI) $m / z:[M+H]^{+}$493.1. HPLC purity: $98.8 \%, t_{\mathrm{R}}=6.91 \mathrm{~min}$. $\mathrm{HRMS}\left(E S I^{+}\right) \mathrm{m} / z:[\mathrm{M}+\mathrm{H}]^{+}$calcd for $\mathrm{C}_{24} \mathrm{H}_{19} \mathrm{CIFN}_{6} \mathrm{OS} 493.1014$; found 493.0999 .
$N$-(3-fluorobenzyl)-2-(4-((6-(1-methyl-1H-pyrazol-4-yl)thieno[3,2-d]pyrimidin-4-yl)amino)phenyl)acetamide (4i). The title compound was synthesized from 14 and (3-fluorophenyl)methanamine following general procedure A. Yellow solid (58\%). ${ }^{1} \mathrm{H}$ NMR ( $400 \mathrm{MHz}, \mathrm{DMSO}_{6}$ ) $\delta 9.57(\mathrm{~s}, 1 \mathrm{H}), 8.57(\mathrm{t}, J=5.8 \mathrm{~Hz}, 1 \mathrm{H}), 8.51(\mathrm{~s}, 1 \mathrm{H}), 8.28(\mathrm{~s}$, $1 \mathrm{H}), 7.96(\mathrm{~s}, 1 \mathrm{H}), 7.67(\mathrm{~d}, J=8.1 \mathrm{~Hz}, 2 \mathrm{H}), 7.54(\mathrm{~s}, 1 \mathrm{H}), 7.35(\mathrm{~d}, J=6.4 \mathrm{~Hz}, 1 \mathrm{H}), 7.27(\mathrm{~d}, J=8.2 \mathrm{~Hz}, 2 \mathrm{H}), 7.11$ $-7.01(\mathrm{~m}, 3 \mathrm{H}), 4.30(\mathrm{~d}, \mathrm{~J}=5.9 \mathrm{~Hz}, 2 \mathrm{H}), 3.91(\mathrm{~s}, 3 \mathrm{H}), 3.49(\mathrm{~s}, 2 \mathrm{H}) .{ }^{13} \mathrm{C}$ NMR (101 MHz, DMSO-d $\mathrm{d}_{6}$ ) 170.41, 161.22, $162.18(J=243 \mathrm{~Hz}), 154.58,154.31,142.57(J=8 \mathrm{~Hz}), 141.81,137.53,136.89,131.38,130.18(J=8$ $\mathrm{Hz})$, 129.17, $129.05(2 \mathrm{C}), 123.10(J=2 \mathrm{~Hz}), 122.04(2 \mathrm{C}), 118.21,115.29,113.76(J=21 \mathrm{~Hz}), 113.46(J=21$ $\mathrm{Hz}), 113.13,41.81,41.70$, 38.82. LC-MS (ESI) $\mathrm{m} / \mathrm{z}:[\mathrm{M}+\mathrm{H}]^{+} 473.1$. HPLC purity: $98.5 \%, t_{\mathrm{R}}=6.19 \mathrm{~min}$. HRMS (ESI) $m / z:[\mathrm{M}+\mathrm{H}]^{+}$calcd for $\mathrm{C}_{25} \mathrm{H}_{22} \mathrm{FN} \mathrm{F}_{6} \mathrm{OS} 473.1560$; found 473.1550. (66\%). ${ }^{1} \mathrm{H}$ NMR (400 MHz, DMSO-d ${ }_{6}$ ) $\delta 10.39(\mathrm{~s}, 1 \mathrm{H}), 9.60(\mathrm{~s}, 1 \mathrm{H}), 8.76(\mathrm{~s}, 1 \mathrm{H}), 8.51(\mathrm{~s}, 1 \mathrm{H}), 8.29(\mathrm{~s}, 1 \mathrm{H})$, $8.26(\mathrm{~d}, J=4.5 \mathrm{~Hz}, 1 \mathrm{H}), 8.05(\mathrm{~d}, J=8.4 \mathrm{~Hz}, 1 \mathrm{H}), 7.96(\mathrm{~s}, 1 \mathrm{H}), 7.70(\mathrm{~d}, J=8.3 \mathrm{~Hz}, 2 \mathrm{H}), 7.54(\mathrm{~s}, 1 \mathrm{H}), 7.34(\mathrm{t}, J=$ $8.7 \mathrm{~Hz}, 3 \mathrm{H}), 3.91(\mathrm{~s}, 3 \mathrm{H}), 3.67(\mathrm{~s}, 2 \mathrm{H}) .{ }^{13} \mathrm{C}$ NMR ( $101 \mathrm{MHz}, \mathrm{DMSO}-d_{6}$ ) $\delta 169.84,161.22,154.57,154.30$, $144.14,141.83,140.75,137.74,136.90,130.64,129.21$ (2C), 129.16, 126.03, 123.61, 122.11(2C), 118.21, 115.30, 113.17, 42.58, 38.82. LC-MS (ESI) $m / z:[M+H]^{+} 442.3$. HPLC purity: $99.7 \%, t_{R}=4.94 \mathrm{~min}$. HRMS (ESI) $m / z:[\mathrm{M}+\mathrm{H}]^{+}$calcd for $\mathrm{C}_{23} \mathrm{H}_{20} \mathrm{~N}_{7} \mathrm{OS} 442.1450$; found 442.1440.

2-(4-((6-bromothieno[3,2-d]pyrimidin-4-yl)amino)phenyl)-N-(4-chloro-3-fluorophenyl)acetamide (16). The title compound was synthesized from 15 and 3-chloro-4-fluoroaniline following general procedure A. Yellow solid ( $88 \%$ ). ${ }^{1} \mathrm{H}$ NMR ( $400 \mathrm{MHz}, \mathrm{DMSO}_{6}$ ) $\delta 10.40(\mathrm{~s}, 1 \mathrm{H}), 9.76(\mathrm{~s}, 1 \mathrm{H}), 8.51(\mathrm{~s}, 1 \mathrm{H}), 7.94(\mathrm{dd}, J=6.9,2.5 \mathrm{~Hz}, 1 \mathrm{H})$, $7.67(\mathrm{~s}, 1 \mathrm{H}), 7.65(\mathrm{~d}, J=8.4 \mathrm{~Hz}, 2 \mathrm{H}), 7.51-7.46(\mathrm{~m}, 1 \mathrm{H}), 7.37(\mathrm{t}, J=9.1 \mathrm{~Hz}, 1 \mathrm{H}), 7.33(\mathrm{~d}, J=8.4 \mathrm{~Hz}, 2 \mathrm{H})$, 3.64 (s, 2H). LC-MS (ESI) m/z: [M + H] ${ }^{+}$491.2/493.2.

General procedure B: Synthesis of compounds 2, 10, 13, and 18. Chloride 5, or 12 ( 0.2 mmol ), amine 8, 9, or 17 ( 0.2 mmol ) were suspended in isopropanol ( 4 mL ). The reaction mixture was irradiated under microwave at $100^{\circ} \mathrm{C}$ for 0.5 h . The completion of the reaction was monitored by LC-MS analysis. The reaction mixture was filtered to get the titled compound.
$N$-(4-methoxyphenyl)thieno[3,2-d]pyrimidin-4-amine (2a). The title compound was synthesized from 5 and 4methoxyaniline following general procedure B. Yellow solid (92\%). ${ }^{1} \mathrm{H}$ NMR ( $400 \mathrm{MHz}, \mathrm{DMSO}-d_{6}$ ) $\delta 11.46$ (s, $1 \mathrm{H}), 8.87(\mathrm{~s}, 1 \mathrm{H}), 8.47(\mathrm{~d}, J=5.3 \mathrm{~Hz}, 1 \mathrm{H}), 7.58(\mathrm{~d}, J=5.4 \mathrm{~Hz}, 1 \mathrm{H}), 7.54(\mathrm{~d}, J=8.6 \mathrm{~Hz}, 2 \mathrm{H}), 7.04(\mathrm{~d}, J=8.8$ $\mathrm{Hz}, 2 \mathrm{H}), 3.80(\mathrm{~s}, 3 \mathrm{H}) .{ }^{13} \mathrm{C}$ NMR ( 101 MHz , DMSO- $\mathrm{d}_{6}$ ) $\delta 168.62,157.13,148.83,147.83,140.02,128.85$, 126.93, 118.21 (2C), 114.18 (2C), 55.41. LC-MS (ESI) $m / z:[M+H]^{+} 258.1$. HPLC purity $99.8 \%, t_{R}=6.93 \mathrm{~min}$.

N -(4-fluorophenyl)thieno[3,2-d]pyrimidin-4-amine (2b). The title compound was synthesized from 5 and 4fluoroaniline following general procedure B. Beige solid ( $88 \%$ ). ${ }^{1} \mathrm{H}$ NMR ( 400 MHz , DMSO- $d_{6}$ ) $\delta 11.38(\mathrm{~s}, 1 \mathrm{H})$, $8.89(\mathrm{~s}, 1 \mathrm{H}), 8.50(\mathrm{~d}, J=5.4 \mathrm{~Hz}, 1 \mathrm{H}), 7.74-7.68(\mathrm{~m}, 2 \mathrm{H}), 7.59(\mathrm{~d}, J=5.5 \mathrm{~Hz}, 1 \mathrm{H}), 7.36-7.29(\mathrm{~m}, 2 \mathrm{H}) .{ }^{13} \mathrm{C}$ NMR ( $\left.101 \mathrm{MHz}, \mathrm{DMSO}-d_{6}\right) \delta 160.17(J=248 \mathrm{~Hz}), 158.94,156.75,149.49,145.77,139.59,133.03,126.80(J$ $=12 \mathrm{~Hz}, 2 \mathrm{C}), 119.00,115.67(\mathrm{~J}=23 \mathrm{~Hz}, 2 \mathrm{C})$. LC-MS (ESI) $\mathrm{m} / \mathrm{z}:[\mathrm{M}+\mathrm{H}]^{+} 246.0$. HPLC purity $97.1 \%, t_{\mathrm{R}}=7.88$ min. NMR (400 MHz, DMSO-d d $_{6}$ ס 11.57 (s, 1H), 8.93 (s, 1H), 8.55 (d, J=5.4 Hz, 1H), 7.81 (d, J= $\left.8.3 \mathrm{~Hz}, 2 \mathrm{H}\right), 7.63$ $(\mathrm{d}, J=5.4 \mathrm{~Hz}, 1 \mathrm{H}), 7.46(\mathrm{~d}, J=8.3 \mathrm{~Hz}, 2 \mathrm{H}), 3.56-3.35(\mathrm{~m}, 4 \mathrm{H}), 1.68-1.42(\mathrm{~m}, 6 \mathrm{H}) .{ }^{13} \mathrm{C}$ NMR ( 101 MHz , DMSO- $d_{6}$ ) $\delta 168.33,156.41,149.38,139.76,137.80,133.76,127.39$ (2C), 123.65, 118.99 (2C), 116.30, 109.55, 48.14 (2C), $25.52(2 \mathrm{C}), 24.05$. LC-MS (ESI) $m / z:[M+H]^{+} 339.1$. HPLC purity $99.5 \%, t_{R}=7.43 \mathrm{~min}$. ethyl 2-(4-(thieno[3,2-d]pyrimidin-4-ylamino)phenyl)acetate (10). The title compound was synthesized from 5 and 9 following general procedure B. Beige solid ( $91 \%$ ). ${ }^{1} \mathrm{H}$ NMR ( 400 MHz , DMSO- $d_{6}$ ) $\delta 11.33(\mathrm{~s}, 1 \mathrm{H}), 8.87$ (s, 1H), $8.50(\mathrm{~d}, J=5.3 \mathrm{~Hz}, 1 \mathrm{H}), 7.61(\mathrm{~d}, J=8.1 \mathrm{~Hz}, 2 \mathrm{H}), 7.58(\mathrm{~d}, J=5.4 \mathrm{~Hz}, 1 \mathrm{H}), 7.36(\mathrm{~d}, J=8.1 \mathrm{~Hz}, 2 \mathrm{H})$, $4.09(\mathrm{q}, J=7.1 \mathrm{~Hz}, 2 \mathrm{H}), 3.71(\mathrm{~s}, 2 \mathrm{H}), 1.20(\mathrm{t}, J=7.1 \mathrm{~Hz}, 3 \mathrm{H}) . \operatorname{LC}-\mathrm{MS}(\mathrm{ESI}) \mathrm{m} / \mathrm{z}:[\mathrm{M}+\mathrm{H}]^{+} 314.3$.
ethyl 2-(4-((6-bromothieno[3,2-d]pyrimidin-4-yl)amino)phenyl)acetate (13). The title compound was synthesized from 12 and 9 following general procedure B. Beige solid (87\%). ${ }^{1} \mathrm{H}$ NMR ( 400 MHz , DMSO- $d_{6}$ ) $\delta 10.84$ (s, $1 \mathrm{H}), 8.73(\mathrm{~s}, 1 \mathrm{H}), 7.73(\mathrm{~s}, 1 \mathrm{H}), 7.61(\mathrm{~d}, J=8.2 \mathrm{~Hz}, 2 \mathrm{H}), 7.33(\mathrm{~d}, J=8.3 \mathrm{~Hz}, 2 \mathrm{H}), 4.09(\mathrm{q}, J=7.3 \mathrm{~Hz}, 2 \mathrm{H}), 3.70$ ( $\mathrm{s}, 2 \mathrm{H}$ ), $1.19(\mathrm{t}, \mathrm{J}=7.3 \mathrm{~Hz}, 3 \mathrm{H}) . \mathrm{LC}-\mathrm{MS}(\mathrm{ESI}) \mathrm{m} / \mathrm{z}:[\mathrm{M}+\mathrm{H}]^{+}$392.1/394.3.
tert-butyl (4-((6-bromothieno[3,2-d]pyrimidin-4-yl)amino)phenyl)carbamate (18). The title compound was synthesized from 12 and 18 following general procedure B. Beige solid ( $93 \%$ ). ${ }^{1} \mathrm{H}$ NMR ( 400 MHz , DMSO- $d_{6}$ ) $\delta$ $10.97(\mathrm{~s}, 1 \mathrm{H}), 9.53(\mathrm{~s}, 1 \mathrm{H}), 8.74(\mathrm{~s}, 1 \mathrm{H}), 7.72(\mathrm{~d}, J=1.5 \mathrm{~Hz}, 1 \mathrm{H}), 7.54(\mathrm{~d}, J=9.0 \mathrm{~Hz}, 2 \mathrm{H}), 7.49(\mathrm{~d}, J=8.9 \mathrm{~Hz}$, 2H), 1.49 (s, 9H). LC-MS (ESI) $m / z:[M+H]^{+}$421.3/423.3.

## Synthesis of compound 4k.

Step 1.18 ( 0.3 mmol ), 1-methyl-4-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)-1 H -pyrazole ( 0.5 mmol ), and $\mathrm{NaHCO}_{3}(1.2 \mathrm{mmol})$ were dissolved in THF (2 mL) and $\mathrm{H}_{2} \mathrm{O}(2 \mathrm{~mL})$. The solution was degassed for 5 min , then $\mathrm{Pd}\left(\mathrm{PPh}_{3}\right)_{4}(0.03 \mathrm{mmol})$ was added. The reaction mixture was irradiated under microwave at $140^{\circ} \mathrm{C}$ for 0.5 h . The consumption of starting materials was detected by LC-MS analysis. The reaction mixture was concentrated in vacuum. The resulting residue was purified by silica gel chromatography to get yellow solid (120 mg, yield 86\%). ${ }^{1} \mathrm{H}$ NMR ( $400 \mathrm{MHz}, \mathrm{CDCl}_{3}$ ) $\delta 8.56(\mathrm{~s}, 1 \mathrm{H}), 7.70(\mathrm{~d}, J=8.4 \mathrm{~Hz}, 2 \mathrm{H}), 7.52-7.45(\mathrm{~m}, 3 \mathrm{H})$, $7.42(\mathrm{~d}, \mathrm{~J}=8.4 \mathrm{~Hz}, 2 \mathrm{H}), 6.64(\mathrm{~s}, 1 \mathrm{H}), 3.94(\mathrm{~s}, 3 \mathrm{H}), 1.55(\mathrm{~s}, 9 \mathrm{H}) . \operatorname{LC}-\mathrm{MS}(\mathrm{ESI}) \mathrm{m} / \mathrm{z}:[\mathrm{M}+\mathrm{H}]^{+} 423.2$. The solid was dissolved in DCM ( 2 mL ) and TFA ( 1 mL ) was added. The mixture was stirred at ambient temperature

Over , int. The completion of the reaction was monitored by LC-MS analysis. The reaction mixture was resonvicentrated in vacuum. The resulting residue was dissolved in DCM $(100 \mathrm{~mL})$ and washed with saturated $\mathrm{NaHCO}_{3}$ solution, water and brine twice. The organic phase was concentrated in vacuum to get 19 as a yellow solid (85 mg, 93\%). ${ }^{1} \mathrm{H}$ NMR (400 MHz, $\mathrm{CDCl}_{3}$ ) $\delta 8.55(\mathrm{~s}, 1 \mathrm{H}), 7.70(\mathrm{~s}, 1 \mathrm{H}), 7.63(\mathrm{~s}, 1 \mathrm{H}), 7.30(\mathrm{~s}, 1 \mathrm{H}), 7.22(\mathrm{~d}$, $J=8.6 \mathrm{~Hz}, 2 \mathrm{H}), 6.74(\mathrm{~d}, J=8.7 \mathrm{~Hz}, 2 \mathrm{H}), 3.93(\mathrm{~s}, 3 \mathrm{H}) . \mathrm{LC}-\mathrm{MS}(\mathrm{ESI}) \mathrm{m} / \mathrm{z}:[\mathrm{M}+\mathrm{H}]^{+} 323.1$.

Step 2. 3-(tert-butyl)-1-methyl-1 H-pyrazol-5-amine ( 0.06 mmol ) and triphogene ( 0.03 mmol ) was dissolved in DCM ( 1 mL ). A solution of $E t_{3} \mathrm{~N}$ in $\mathrm{DCM}\left(0.15 \mathrm{mmol}_{\mathrm{Et}}^{3} \mathrm{~N}\right.$ in 1 mL DCM$)$ was added slowly at $0^{\circ} \mathrm{C}$. The reaction mixture was warmed to ambient temperature and stirred for 2 h . The reaction mixture was added slowly into a solution of $19(0.05 \mathrm{mmol})$ in $\mathrm{DCM}(1 \mathrm{~mL})$ at ambient temperature. The resulting reaction mixture was stirred at ambient temperature for 5 h . The completion of the reaction was monitored by LC-MS analysis. The reaction mixture was poured into water and extracted with DCM. The organic phase was washed with water and brine twice, and concentrated in vacuum. The resulting residue was purified by silica gel chromatography to get $\mathbf{4 k}$ as a yellow solid ( $20 \mathrm{mg}, 66 \%$ ). ${ }^{1} \mathrm{H} \operatorname{NMR}\left(400 \mathrm{MHz}, \mathrm{CDCl}_{3}\right) \delta 8.45(\mathrm{~s}, 1 \mathrm{H}), 7.78(\mathrm{~s}, 1 \mathrm{H}), 7.70(\mathrm{~s}, 1 \mathrm{H}), 7.49(\mathrm{~d}, \mathrm{~J}$ $=8.8 \mathrm{~Hz}, 2 \mathrm{H}), 7.36(\mathrm{~d}, \mathrm{~J}=8.8 \mathrm{~Hz}, 2 \mathrm{H}), 7.32(\mathrm{~s}, 1 \mathrm{H}), 6.22(\mathrm{~s}, 1 \mathrm{H}), 3.89(\mathrm{~s}, 3 \mathrm{H}), 3.71(\mathrm{~s}, 3 \mathrm{H}), 1.25(\mathrm{~s}, 9 \mathrm{H}) . \mathrm{LC}-$ MS (ESI) $m / z:[\mathrm{M}+\mathrm{H}]^{+}$502.0. HPLC purity: $97.2 \%, t_{\mathrm{R}}=7.90 \mathrm{~min}$. HRMS (ESI) $m / z:[\mathrm{M}+\mathrm{H}]^{+}$calcd for $\mathrm{C}_{25} \mathrm{H}_{28} \mathrm{~N}_{9} \mathrm{OS} 502.2138$; found 502.2128.

General procedure C: Synthesis of compounds 4I-n. 16 ( 0.1 mmol ), boronic acid ( 0.15 mmol ), and $\mathrm{NaHCO}_{3}$ ( 0.3 mmol ) were dissolved in THF $(2 \mathrm{~mL})$ and $\mathrm{H}_{2} \mathrm{O}(2 \mathrm{~mL})$. The solution was degassed for 5 min , then $\mathrm{Pd}^{\left(\mathrm{PPh}_{3}\right)_{4}}$ ( 0.005 mmol ) was added. The reaction mixture was irradiated under microwave at $140^{\circ} \mathrm{C}$ for 0.5 h . The consumption of starting materials was detected by LC-MS analysis. The reaction mixture was poured into water and extracted with ethyl acetate. The organic phase was washed with water and brine twice, and concentrated in vacuum. The resulting residue was purified by silica gel chromatography to get the title compound.

## N-(3-chloro-4-fluorophenyl)-2-(4-((6-(4-(methylsulfonyl)phenyl)thieno[3,2-d]pyrimidin-4-

$y l) a m i n o) p h e n y l) a c e t a m i d e ~(4 I)$. The title compound was synthesized from 16 and (4-
(methylsulfonyl)phenyl)boronic acid following general procedure C. Yellow solid ( $66 \%$ ). ${ }^{1} \mathrm{H}$ NMR ( 400 MHz , DMSO-d $d_{6}$ ס $10.39(\mathrm{~s}, 1 \mathrm{H}), 9.83(\mathrm{~s}, 1 \mathrm{H}), 8.59(\mathrm{~s}, 1 \mathrm{H}), 8.17-8.10(\mathrm{~m}, 3 \mathrm{H}), 8.06(\mathrm{~d}, \mathrm{~J}=8.5 \mathrm{~Hz}, 2 \mathrm{H}), 7.94(\mathrm{dd}, \mathrm{J}$ $=6.8,2.5 \mathrm{~Hz}, 1 \mathrm{H}), 7.73(\mathrm{~d}, J=8.5 \mathrm{~Hz}, 2 \mathrm{H}), 7.52-7.47(\mathrm{~m}, 1 \mathrm{H}), 7.40-7.31(\mathrm{~m}, 3 \mathrm{H}), 3.65(\mathrm{~s}, 2 \mathrm{H}), 3.29(\mathrm{~s}, 3 \mathrm{H})$. 120.45, $119.36(J=6 \mathrm{~Hz}), 119.09(J=18 \mathrm{~Hz}), 116.92(J=21 \mathrm{~Hz}), 115.80,43.42,42.68$. LC-MS (ESI) $\mathrm{m} / \mathrm{z}$ : $[\mathrm{M}+\mathrm{H}]^{+} 567.3$. HPLC purity $95.0 \%, t_{\mathrm{R}}=7.06 \mathrm{~min}$.

N-(3-chloro-4-fluorophenyl)-2-(4-((6-(3,4,5-trimethoxyphenyl)thieno[3,2-d]pyrimidin-4$y l) a m i n o)$ phenyl)acetamide (4m). The title compound was synthesized from 16 and (3,4,5trimethoxyphenyl)boronic acid following general procedure C. Beige solid ( $62 \%$ ). ${ }^{1} \mathrm{H}$ NMR ( 400 MHz , DMSO$\left.d_{6}\right) \delta 10.37(\mathrm{~s}, 1 \mathrm{H}), 9.68(\mathrm{~s}, 1 \mathrm{H}), 8.57(\mathrm{~s}, 1 \mathrm{H}), 7.96-7.92(\mathrm{~m}, 2 \mathrm{H}), 7.75(\mathrm{~d}, J=8.6 \mathrm{~Hz}, 2 \mathrm{H}), 7.51-7.46(\mathrm{~m}$, $1 \mathrm{H}), 7.38(\mathrm{~d}, \mathrm{~J}=9.1 \mathrm{~Hz}, 1 \mathrm{H}), 7.35-7.30(\mathrm{~m}, 2 \mathrm{H}), 7.14(\mathrm{~s}, 2 \mathrm{H}), 3.90(\mathrm{~s}, 6 \mathrm{H}), 3.73(\mathrm{~s}, 3 \mathrm{H}), 3.64(\mathrm{~s}, 2 \mathrm{H}) .{ }^{13} \mathrm{C}$ NMR ( 101 MHz, DMSO- $d_{6}$ ) $\delta 169.52,161.04,154.60,154.46,153.40(2 \mathrm{C}), 152.99(J=243 \mathrm{~Hz}), 149.59$, $138.76,137.77,136.45(J=3 \mathrm{~Hz}), 130.58,129.21(2 \mathrm{C}), 128.24,121.88(2 \mathrm{C}), 120.48,120.45,119.36(J=7$ $\mathrm{Hz}), 119.07(J=18 \mathrm{~Hz}), 116.92(J=21 \mathrm{~Hz}), 114.67,103.81(2 C), 60.17,56.09(2 \mathrm{C})$, 42.67. LC-MS (ESI) $\mathrm{m} / \mathrm{z}$ : $[\mathrm{M}+\mathrm{H}]^{+} 579.1$. HPLC purity $98.0 \%, t_{\mathrm{R}}=7.49 \mathrm{~min}$.

N-(3-chloro-4-fluorophenyl)-2-(4-((6-(4-(2-(pyrrolidin-1-yl)ethoxy)phenyl)thieno[3,2-d]pyrimidin-4$y l)$ amino)phenyl)acetamide (4n). The title compound was synthesized from 16 and (4-(2-(pyrrolidin-1yl)ethoxy)phenyl)boronic acid following general procedure C. Yellow solid (52\%). ${ }^{1} \mathrm{H}$ NMR ( 400 MHz , DMSO$\left.d_{6}\right) \delta 10.35(\mathrm{~s}, 1 \mathrm{H}), 9.61(\mathrm{~s}, 1 \mathrm{H}), 8.49(\mathrm{~s}, 1 \mathrm{H}), 7.91(\mathrm{dd}, J=6.9,2.5 \mathrm{~Hz}, 1 \mathrm{H}), 7.75(\mathrm{~d}, J=8.7 \mathrm{~Hz}, 2 \mathrm{H}), 7.72(\mathrm{~s}$, $1 \mathrm{H}), 7.67(\mathrm{~d}, J=8.4 \mathrm{~Hz}, 2 \mathrm{H}), 7.48-7.42(\mathrm{~m}, 1 \mathrm{H}), 7.33(\mathrm{t}, J=9.1 \mathrm{~Hz}, 1 \mathrm{H}), 7.28(\mathrm{~d}, J=8.4 \mathrm{~Hz}, 2 \mathrm{H}), 7.04(\mathrm{~d}, J=$ $8.8 \mathrm{~Hz}, 2 \mathrm{H}), 4.10(\mathrm{t}, J=5.8 \mathrm{~Hz}, 2 \mathrm{H}), 3.60(\mathrm{~s}, 2 \mathrm{H}), 2.78(\mathrm{t}, J=5.8 \mathrm{~Hz}, 2 \mathrm{H}), 2.48-2.44(\mathrm{~m}, 3 \mathrm{H}), 1.69-1.61(\mathrm{~m}$, 4H). ${ }^{13} \mathrm{C}$ NMR ( 101 MHz , DMSO- $d_{6}$ ) $\delta 169.54,161.34,159.71,154.66,154.39,153.01(J=243 \mathrm{~Hz}), 149.59$, 137.71, $136.47(J=3 \mathrm{~Hz}), 130.73,129.21(2 \mathrm{C}), 127.74(2 \mathrm{C}), 125.13,122.22(2 \mathrm{C}), 120.47,119.36(J=7 \mathrm{~Hz})$, $119.08(J=18 \mathrm{~Hz}), 118.92,116.93(J=22 \mathrm{~Hz}), 115.29(2 \mathrm{C}), 114.16,66.96,54.18,53.98(2 \mathrm{C}), 42.70,23.15$ (2C). LC-MS (ESI) $m / z:[M+H]^{+} 602.2$. HPLC purity $98.1 \%, t_{R}=6.34 \mathrm{~min}$.
 (ersolillol), and $\mathrm{NaHCO}_{3}(14.0 \mathrm{mmol})$ were dissolved in THF $(5 \mathrm{~mL})$ and $\mathrm{H}_{2} \mathrm{O}(5 \mathrm{~mL})$. The solution was degassea for 5 min , then $\mathrm{Pd}\left(\mathrm{PPh}_{3}\right)_{4}(0.2 \mathrm{mmol})$ was added. The reaction mixture was irradiated under microwave at 140 ${ }^{\circ} \mathrm{C}$ for 0.5 h . The consumption of starting materials was detected by LC-MS analysis. $\mathrm{LiOH}(14.0 \mathrm{mmol})$ was added and the reaction mixture was irradiated under microwave at $100^{\circ} \mathrm{C}$ for 0.5 h . The consumption of starting materials was detected by LC-MS analysis. The resulting mixture was adjusted to $\mathrm{pH} 3-4$ with 1 N HCl aq solution. The resulting suspension was stirred for 5 h then filtered. The residue was dried under vacuum at $45^{\circ} \mathrm{C}$ overnight to get the titled compound.

2-(4-(thieno[3,2-d]pyrimidin-4-yl)phenyl)acetic acid (7). The title compound was synthesized from 5 and 6 following general procedure D. Grey solid (yield, 82\%) ${ }^{1} \mathrm{H}$ NMR ( 400 MHz , DMSO-d $\mathrm{d}_{6}$ ס 9.26 (s, 1H), 8.57 (d, J $=5.5 \mathrm{~Hz}, 1 \mathrm{H}), 8.14(\mathrm{~d}, J=8.2 \mathrm{~Hz}, 2 \mathrm{H}), 7.73(\mathrm{~d}, J=5.5 \mathrm{~Hz}, 1 \mathrm{H}), 7.55(\mathrm{~d}, J=8.0 \mathrm{~Hz}, 2 \mathrm{H}), 3.73(\mathrm{~s}, 2 \mathrm{H}) . \mathrm{LC}-\mathrm{MS}$ (ESI) $m / z:[\mathrm{M}+\mathrm{H}]^{+} 271.0$.

2-(4-((6-(1-methyl-1H-pyrazol-4-yl)thieno[3,2-d]pyrimidin-4-yl)amino)phenyl)acetic acid (14). The title compound was synthesized from 13 and 1-methyl-4-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)-1H-pyrazole following general procedure D. Grey solid (yield, $77 \%)^{1} \mathrm{H}$ NMR ( $400 \mathrm{MHz}, \mathrm{DMSO}-\mathrm{d}_{6}$ ) $\delta 12.32(\mathrm{~s}, 1 \mathrm{H}), 9.58(\mathrm{~s}$, $1 \mathrm{H}), 8.51(\mathrm{~s}, 1 \mathrm{H}), 8.29(\mathrm{~s}, 1 \mathrm{H}), 7.96(\mathrm{~s}, 1 \mathrm{H}), 7.68(\mathrm{~d}, J=8.4 \mathrm{~Hz}, 2 \mathrm{H}), 7.54(\mathrm{~s}, 1 \mathrm{H}), 7.24(\mathrm{~d}, J=8.4 \mathrm{~Hz}, 2 \mathrm{H}), 3.91$ (s, 3H), 3.55 (s, 2H). LC-MS (ESI) m/z: [M + H]+ 366.1.

Synthesis of compound 11. A mixture of $10(4.0 \mathrm{mmol})$ and $\mathrm{LiOH}(16.0 \mathrm{mmol})$ in THF $(5 \mathrm{~mL})$ and $\mathrm{H}_{2} \mathrm{O}(5 \mathrm{~mL})$ was irradiated under microwave at $100^{\circ} \mathrm{C}$ for 0.5 h . The consumption of starting materials was detected by LC-MS analysis. The resulting mixture was adjusted to $\mathrm{pH} 3-4$ with 1 N HCl solution. The resulting suspension was stirred for 5 h then filtered. The residue was dried under vacuum at $45^{\circ} \mathrm{C}$ overnight to get the titled compound ( 980 mg , yield $86 \%$ ). ${ }^{1} \mathrm{H}$ NMR ( 400 MHz , DMSO- $\mathrm{d}_{6}$ ) $\delta 12.31(\mathrm{~s}, 1 \mathrm{H}), 9.68(\mathrm{~s}, 1 \mathrm{H}), 8.56(\mathrm{~s}, 1 \mathrm{H}), 8.22$ (d, $J=5.2 \mathrm{~Hz}, 1 \mathrm{H}), 7.70(\mathrm{~d}, J=7.9 \mathrm{~Hz}, 2 \mathrm{H}), 7.46(\mathrm{~d}, J=5.4 \mathrm{~Hz}, 1 \mathrm{H}), 7.25(\mathrm{~d}, J=8.0 \mathrm{~Hz}, 2 \mathrm{H}), 3.56(\mathrm{~s}, 2 \mathrm{H})$. LCMS (ESI) m/z: $[\mathrm{M}+\mathrm{H}]^{+}$286.3.

## Synthesis of compound 24.

Step 1. 2-(5-amino-3-(tert-butyl)-1 H -pyrazol-1-yl)ethanol[34] ( 27.3 mmol ) and $\mathrm{Et}_{3} \mathrm{~N}(81.8 \mathrm{mmol})$ was dissolved in DCM $(100 \mathrm{~mL})$. A solution of $\mathrm{TsCl}(81.8 \mathrm{mmol})$ in $\mathrm{DCM}(100 \mathrm{~mL})$ was added dropwise. The resulting reaction
mixty = was refluxed overnight. The consumption of starting materials was detected by LC-MS analysis.
0 mL ) was added into the reaction mixture. Separate the organic phase and concentrated in vacuum. resulting residue was purified by silica gel chromatography to get 21 as a beige solid ( 8.8 g , yield $50 \%$ ). ${ }^{1} \mathrm{H}$ NMR (400 MHz, CDCl ${ }_{3}$ ) $\delta 7.77(\mathrm{dd}, J=11.5,8.3 \mathrm{~Hz}, 6 \mathrm{H}), 7.36(\mathrm{~d}, J=8.1 \mathrm{~Hz}, 4 \mathrm{~h}), 7.30(\mathrm{~d}, J=8.1 \mathrm{~Hz}, 2 \mathrm{H})$, $5.66(\mathrm{~s}, 1 \mathrm{H}), 4.35(\mathrm{t}, \mathrm{J}=6.4 \mathrm{~Hz}, 2 \mathrm{H}), 3.91(\mathrm{t}, J=6.4 \mathrm{~Hz}, 2 \mathrm{H}), 2.49(\mathrm{~s}, 6 \mathrm{H}), 2.42(\mathrm{~s}, 3 \mathrm{H}) . \operatorname{LC}-\mathrm{MS}(\mathrm{ESI}) \mathrm{m} / \mathrm{z}$ : $[\mathrm{M}+\mathrm{H}]^{+}$646.2.

Step 2. A mixture of $21(3.0 \mathrm{mmol})$ in morpholine $(30 \mathrm{~mL})$ was heated to $120^{\circ} \mathrm{C}$ overnight. The consumption of starting materials was detected by LC-MS analysis. Water ( 100 mL ) and DCM $(200 \mathrm{~mL})$ was added into the reaction mixture. The organic phase was washed with water and brine twice, and concentrated in vacuum. The resulting residue was purified by silica gel chromatography to get 22 as colorless oil ( 0.8 g , yield $64 \%$ ). ${ }^{1} \mathrm{H}$ NMR ( $400 \mathrm{MHz}, \mathrm{CDCl}_{3}$ ) $\delta 7.63(\mathrm{~d}, J=8.1 \mathrm{~Hz}, 2 \mathrm{H}), 7.23(\mathrm{~d}, J=8.1 \mathrm{~Hz}, 2 \mathrm{H}), 5.99(\mathrm{~s}, 1 \mathrm{H}), 3.91(\mathrm{t}, J=4.6 \mathrm{~Hz}$, 4h), $3.88-3.83$ (m, 2H), 2.68 - 2.62 (m, 2H), 2.58 (s, 4h), 2.39 (s, 3H), 1.22 (s, 9H). LC-MS (ESI) m/z: 646.2 $[\mathrm{M}+\mathrm{H}]^{+}$. Also get 23 as colorless oil ( 0.36 g , yield $30 \%$ ). ${ }^{1} \mathrm{H}$ NMR ( $400 \mathrm{MHz}, \mathrm{CDCl}_{3}$ ) $\delta 7.68(\mathrm{~d}, J=7.9 \mathrm{~Hz}, 2 \mathrm{H})$, $7.26(\mathrm{~d}, J=8.1 \mathrm{~Hz}, 2 \mathrm{H}), 5.91(\mathrm{~s}, 1 \mathrm{H}), 4.11(\mathrm{t}, J=7.6 \mathrm{~Hz}, 2 \mathrm{H}), 3.97(\mathrm{t}, J=7.5 \mathrm{~Hz}, 2 \mathrm{H}), 2.39(\mathrm{~s}, 3 \mathrm{H}), 1.26(\mathrm{~s}$, 9H). LC-MS (ESI) $m / z:[M+H]^{+} 320.1$.

Step 3. A mixture of $\mathbf{2 2}(0.9 \mathrm{mmol})$ in $\mathrm{H}_{2} \mathrm{SO}_{4}$ was stirred at ambient temperature overnight. The consumption of starting materials was detected by LC-MS analysis. The reaction mixture was poured into iced water and adjusted to pH 8-9 with 2 N NaOHaq solution. The resulting mixture was concentrated in vacuum to get crude 24 as a white solid. LC-MS (ESI) $\mathrm{m} / \mathrm{z}:[\mathrm{M}+\mathrm{H}]^{+}$253.1.

## Enzymatic assay:

TRKA-C activity was measured via a mobility shift assay monitoring the separation of the phosphorylated product from substrate. In 96 -well plates, compound stock solution ( 20 mM in DMSO) were diluted with kinase buffer ( 10 mM HEPES, $0.05 \%$ Triton X-100, $5 \%$ glycerol, $5 \mathrm{mM} \mathrm{DTT} ,10 \mathrm{mM} \mathrm{MgCl}_{2}, 10 \mathrm{mM} \mathrm{MnCl} 2$ and $0.1 \mathrm{mg} / \mathrm{mL}$ BSA) in 12 -point $1 / 2 \log$ dilutions ( $2 \mathrm{mM}-6.32 \mathrm{nM}$ ). The TRKA enzyme (Life Technologies, CA, USA) was diluted to $2.35 \mu \mathrm{~g} / \mathrm{ml}$ with the kinase buffer. Then, $4 \mu \mathrm{l}$ of the TRKA solution and $1 \mu \mathrm{l}$ of the diluted compound solution were transferred to a 384 -well assay plate to be incubated for 1 hour. $1 \mu \mathrm{l}$ of kinase buffer was added in the positive control instead of the compound solution and $5 \mu$ l of the kinase buffer were added to the negative control.

After ${ }_{\text {S }}$ e pre-incubation, $5 \mu$ l of the substrate mix containing $10 \mu \mathrm{M} \mathrm{ATP}$ and $3 \mu \mathrm{M} 5$-FAM tagged peptide (erssonivstrate \#22, 5-FAM-EEPLYWSFPAKKK-CONH 2 [PerkinElmer, MA, USA]) in kinase buffer were added to crerso assay plate. Final running concentrations were as follows: ATP ( $5 \mu \mathrm{M}$ ), peptide ( $1.5 \mu \mathrm{M}$ ), compounds ( $0.2 \mathrm{mM}-$ 0.632 nM ). Entrectinib was used as the running control. The plate was run on the Caliper EZ Reader II (PerkinElmer, MA, USA) with separation buffer ( 100 mM HEPES, 10 mM EDTA, $0.015 \%$ Brij-35, $0.1 \%$ CR-3 [PerkinElmer, MA, USA]) until 10-20\% conversion based on the positive control wells. The following separation conditions were utilized: upstream voltage -500 V ; downstream voltage, -1900 V ; chip pressure -0.8 . Doseresponse curves were generated in GraphPad Prisim 8 and $\mathrm{IC}_{50}$ values were obtained from the curves. $\mathrm{IC}_{50}$ values were generated in duplicate and error was calculated from the standard deviation between two values.

## Cellular assay:

KM12 cells were cultured using RPMI-1640 (Gibco, CA, USA) growth medium containing 10\% fetal bovine serum (FBS) (Gibco, Carlsbad, USA), $100 \mathrm{U} / \mathrm{ml}$ penicillin (Gibco, Carlsbad, USA) and $100 \mathrm{U} / \mathrm{ml}$ streptomycin (Gibco, CA, USA). The cells were cultured at $37^{\circ} \mathrm{C}$ in a humidified atmosphere with $5 \% \mathrm{CO}_{2}$. To do the viability assay, cells were seeded in 96 -well cell culture plates. After 24 -hour incubation, cells were exposed to 7 concentrations of compounds ranging from 200 to $0.0002 \mu \mathrm{M}$ for 72 hours. Meantime, DMSO were added to the controls. Resazurin staining viability assay (Biotium, CA, USA) was used to determine the cell viability at the end of compound treatment. Then, $\mathrm{IC}_{50}$ values were calculated from the dose-response curves which were generated in GraphPad Prism 8. Error was determined as the standard deviation between two independent $\mathrm{IC}_{50}$ measurements.

## Western blot:

Cells were harvested in lysis buffer ( 50 mM Hepes, pH 7.5, $150 \mathrm{mM} \mathrm{NaCl}, 10 \%$ glycerol, $1 \%$ Triton X100, 1 mM EGTA, $1.5 \mathrm{mM} \mathrm{MgCl} \mathrm{m}_{2}, 10 \mathrm{mM} \mathrm{NaF}, 10 \mathrm{mM}$ sodium pyrophosphate, $1 \mathrm{mM} \mathrm{Na} \mathrm{VO}_{4}, 10 \mu \mathrm{~g}$ of aprotinin $/ \mathrm{ml}, 10 \mu \mathrm{~g}$ of leupeptin $/ \mathrm{ml}$ ) and clarified by centrifugation at $10,000 \times \mathrm{g}$. Protein concentration was estimated with a modified Bradford assay (Bio-Rad, Munich, Germany). Antigens were revealed by an enhanced chemiluminescence detection kit (Pierce ECL, ThermoFisher Scientific, Waltham, MA, USA) [14]. PhosphoTRKA (Tyr490) (\#9141), and phospho-TRKA (Tyr674/675) (\#4621) antibodies were from Cell Signaling Technology (Danvers, MA, USA). Anti-TRKA (C-14) (\#sc-11) antibody was from Santa Cruz Biotechnology

USA). Secondary antibodies coupled to horseradish peroxidase were purchased from Bio-Rad.

## Molecular modeling:

Molecular modeling studies were performed using AutoDock Tools, AutoDock Vina,[35] and Maestro 2018-2. TRKA crystal structure (PDB ID: 6PL1) was retrieved from RCSB Protein Data Bank. Using AutoDock Tools, the TRK model was prepared as follows: (i) remove all waters; (ii) add all hydrogen as 'Polar Only'; (iii) generate a grid box for the ATP binding site, which includes the hinge region, DFG residues and the residues of the $\alpha$ C-helix facing the ATP pocket. Compounds to be docked were assigned torsions around rotatable bonds using AutoDock Tools and were docked into the grid by using Autodock Vina. The docking results were visualized and analyzed with Maestro 2018-2.

## In vitro human brain-like endothelial cells (BLEC) BBB model:

A human BBB model was then used in order to predict entrectinib and $\mathbf{4 c}$ passage. The human in vitro BBB model derived from hematopoietic stem cells was prepared as described previously.[36] Infants' parents signed a consent form and cells were manipulated following the procedure validated by the local ethical committee and French ministry of Research. Briefly, CD34+ cells were isolated from human umbilical cord blood by ficoll density gradient and using the mini-MACS immunomagnetic separation system (Miltenyi Biotec, Bergisch Gladbach), according to the manufacturer's recommendations. These CD34 positive cells were subsequently were transferred onto $1 \%(\mathrm{w} / \mathrm{v})$ gelatin coated 24 -well plates ( $2 \times 10^{5}$ cells/well) and incubated in endothelial growth medium (EGM-2; Lonza, Gaithersburg, MD, USA) with $20 \%(\mathrm{v} / \mathrm{v})$ fetal bovine serum (FBS; Invitrogen,Carlsbad, USA) and $50 \mathrm{ng} / \mathrm{mL}$ vascular endothelial growth factor (VEGF165; PrepoTech Inc., Rocky Hill, USA), at 5\% $\mathrm{CO}_{2}$, $37^{\circ} \mathrm{C}$. After 2-3 weeks they are differentiated into endothelial cells and endothelial markers are checked by western blot, FACs and quantitative PCR.[37] Then, CD34+derived endothelial cells were co-cultivated with bovine brain pericytes in Transwell ${ }^{\text {TM }}$ Costar polycarbonate inserts, $0.4 \mu \mathrm{~m}$ pore size, coated with matrigel® (Corning) in a complete medium (ScienCell). Brain péricytes were isolated as previously described.[38] After 6 days of co-culture, CD34+-derived endothelial cells have acquired BBB properties induced by brain pericytes as previously described.[39] This model is then named Human brain like-endothelial cells (BLECs) and is routinely used to determine molecules toxicity and delivery into the CNS.[40-42]

Endothelial permeability coefficients were determined as described by Dehouck et al.[43] The inserts, containing an endothelial cell monolayer or only coated with matrigel, were transferred into 12 -well plates containing 1.5 mL of Ringer-HEPES (RH) bufferer saline ( $150 \mathrm{mM} \mathrm{NaCl}, 5.2 \mathrm{mM} \mathrm{KCl}, 2.2 \mathrm{mM} \mathrm{CaCl}$,, 0.2 mM $\mathrm{MgCl}_{2}, 0.6 \mathrm{mM} \mathrm{NaHCO} 3,5 \mathrm{mM}$ HEPES, pH 7.4 ) per well, thus constituting the abluminal compartment. Then, 0.5 mL of RH buffer saline containing $10 \mu \mathrm{M}$ of the compound to be tested (entrectinib or 4 c ) was added to the upper (luminal) compartment, in place of the culture medium in contact with the BLECs. After different timepoints ( 30,60 , and 120 min ), filters were transferred into a new well filled with RH buffer saline. All incubations were performed at $37^{\circ} \mathrm{C}$ and stopped at 120 minutes.

To assess the toxicity of the molecules on the BLECs, Lucifer yellow (LY) (Sigma-Aldrich, L0259) that poorly crosses the BBB, was used as the monolayer integrity marker. Diffusion of $25 \mu \mathrm{M}$ LY was tested alone (control) or incubated with entrectinib or 4c, as described above. At 120 min , a $200 \mu \mathrm{~L}$ aliquot from a lower compartment at each time point and a $20 \mu \mathrm{~L}$ aliquot of the initial solution placed in the upper compartment were measured using a fluorimeter (Synergy H1; BioTek, Winooski, VT, USA) to calculate the LY concentration.

Concentrations of entrectinib or $\mathbf{4 c}$ were determined by LC-MS (Waters ACQUITY UPLC System with ACQUITY QDa mass spectrometer) in the solutions from the upper and lower compartments. The endothelial permeability coefficients ( Pe ) to LY , entrectinib and $\mathbf{4 c}$, were calculated using the clearance principle to generate a concentration-independent parameter as described by Siflinger-Birnboim et al.[44] First, the volumes cleared were plotted versus time ( 30 and 60 min ) and the slopes were estimated by linear regression for both insert only coated with matrigel (PSf) and for insert coated with matrigel and seeded with cells (PSt). PSe was calculated according to the following formula: $1 / \mathrm{PSe}=1 / \mathrm{PSt}-1 / \mathrm{PSf}$ where " S " represented the surface area of the porous membrane of the insert. PSe was divided by the membrane surface ( $1.12 \mathrm{~cm}^{2}$ ) to generate the endothelial Pe to the studied molecules expressed in $\mathrm{cm} / \mathrm{min}$.

To assess possible adsorption, aggregation and/or degradation of molecules during the transport experiments, their mass balances (recovery in \%) were calculated at 120 min from the amount of compound recovered in both compartments at the end of the experiment divided by the total amount added in the donor compartment at the start of the experiment.

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## Author contributions

The idea was conceived by WY, BF and HL. The synthesis was done by WY and FL. The biological assays were performed by LZ, MM, FC and CL. The article was written by WY and edited by all.
${ }^{\text {TW }}$ W and LZ contributed equally.

## Declaration of competing interest

The authors declare no competing financial interest.

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## Appendix A. Supplementary data

The following is the Supplementary data to this article: ${ }^{1} \mathrm{H}$ NMR, ${ }^{3} \mathrm{C}$ NMR and HPLC spectra of final compounds. Supplementary information for endothelial permeability measurement assay.

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31\% TRKA inhibition @ $20 \mu \mathrm{M}$ KM12 IC $_{50}=\mathbf{2 5} \boldsymbol{\mu}$ M


17\% TRKA inhibition @ $20 \mu \mathrm{M}$ KM12 $\mathrm{IC}_{50}=133 \boldsymbol{\mu}$


Lead

A series of novel thieno[3,2-d]pyrimidine type-II pan-TRK inhibitors were designed and synthesized.

- Representative compound $\mathbf{4 c}$ was discovered through structure-based drug design (SBDD) approach
- Compound 4c exhibited an effective blockage of TRKA autophosphorylation.
- Compound $\mathbf{4 g}$ exhibited excellent selectivity for KM12 in the NCI-60 human cancer cell lines screen

