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Nanomolar-Potency Small Molecule Inhibitor of STAT5 Protein

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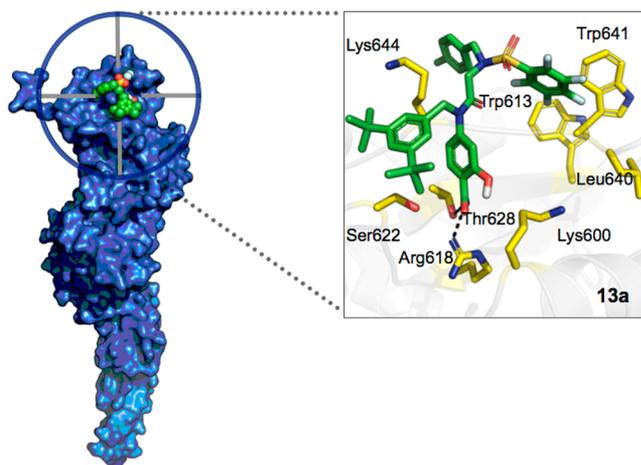
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S Supporting Information

ABSTRACT: We herein report the design and synthesis of the first nanomolar binding inhibitor of STAT5 protein. Lead compound **13a**, possessing a phosphotyrosyl-mimicking salicylic acid group, potently and selectively binds to STAT5 over STAT3, inhibits STAT5–SH2 domain complexation events *in vitro*, silences activated STAT5 in leukemic cells, as well as STAT5's downstream transcriptional targets, including MYC and MCL1, and, as a result, leads to apoptosis. We believe **13a** represents a useful probe for interrogating STAT5 function in cells as well as being a potential candidate for advanced preclinical trials.



KEYWORDS: STAT5, small-molecule inhibitor, protein–protein interactions, anticancer drug, leukemia cells

Numerous inhibitors have been developed to target the JAK-STAT signaling pathway, a major driving force in hematopoietic malignancies. While emphasis has focused on identifying effective upstream kinase inhibitors for suppressing STAT activity, inhibitors have suffered from poor kinase selectivity,¹ cardiovascular toxicity,^{2,3} and, in some clinical cases, acquired resistance.⁴ Multiple kinase inhibitors have been used in combination to try to combat resistance.^{5,6} An alternative strategy to reduce off-target toxicity is to target proteins immediately downstream of the kinases, such as the Signal Transducer and Activator of Transcription (STAT) 5 protein. STAT5 is thus a compelling molecular target for therapeutic intervention.

In normal cells, the activation of STAT5 proteins is tightly regulated by cytokines (IL-2, IL-5, IL-7, GM-CSF, erythropoietin (EPO), thrombopoietin, and prolactin) and growth factors.⁷ Binding of these extracellular ligands to their target receptors induces the activation of receptor-associated JAK kinases that phosphorylate key tyrosine residues within the receptor, providing docking sites for the SRC homology 2 (SH2) domains of the inactive cytoplasmic STAT5 monomers. STAT5 is then phosphorylated at specific tyrosine residues, either Y694 (STAT5A) or Y699 (STAT5B) of the C-terminus. Phosphorylated STAT5 monomers form either homo- or

hetero-^{8,9} STAT5X-STAT5Y dimers through reciprocal phosphotyrosine–SH2 interactions. Activated STAT5 dimers translocate to the nucleus, where they bind to STAT5 DNA response elements inducing transcription of genes involved in proliferation (Bcl-xl, c-Myc, *pim-1*), cell differentiation (p21), cell survival (MCL-1) inflammation (Osm), and apoptosis (JAB).¹⁰ In contrast, mutations within cytosolic kinases (TEL-JAK2, Bcr-Abl, FLT-3) as well as overactive receptor-associated tyrosine kinases (SRC, EGFR) induce constitutive phosphorylation of STAT5 proteins, increasing the production of antiapoptotic genes, which can contribute to driving the cancer phenotype.¹¹

Approaches aimed at directly targeting STAT5 have been limited to a chromone-derived acyl hydrazine inhibitor, identified through a high-throughput fluorescence polarization (FP) screen. While this agent exhibited promising *in vitro* disruption of the STAT5:EPOR phosphopeptide interaction, higher concentrations were required to inhibit STAT5 in cells.¹² A STAT5 function-based screening approach identified

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the FDA approved neuroleptic drug, pimozide, as a potent inhibitor of STAT5 phosphorylation, resulting in the down-regulation of STAT5 target gene expression in Ba/f3 FLT3 ITD cells. The authors believe pimozide is an effector of negative regulators that modulate STAT5 phosphorylation.^{13,14}

We have previously prepared a library of phosphopeptide-mimicking salicylic acid-based compounds to target STAT3's SH2 domain. Screening against STAT5 via FP¹⁵ identified compounds **1** and **2** as STAT5 binders with 2–3-fold selectivity over STAT3 (Figure 1). In whole cells, **1** showed selective

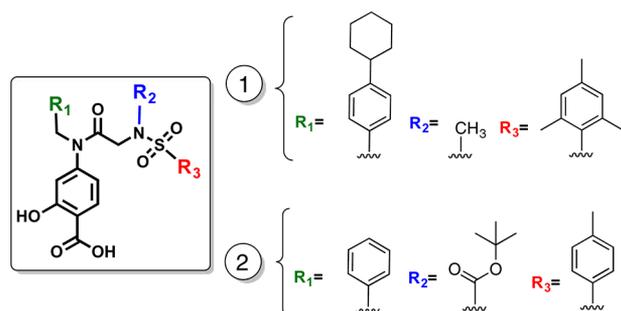


Figure 1. STAT5 inhibitors BP-1-108 (**1**) and SF-1-088 (**2**).

suppression of STAT5 Tyr phosphorylation ($IC_{50} \sim 20 \mu\text{M}$) and STAT5 target genes, *MYC* and *CCND1*, at μM concentrations ($\sim 40 \mu\text{M}$).¹⁶ However, progress toward a potent and selective nanomolar (nM) inhibitor of STAT5 has been slow. Herein, we describe the identification of the first nM, STAT5-selective inhibitor using *in silico*-directed efforts.¹⁷

Scaffolds **1** and **2** were selected as a starting point for structural optimization. Interestingly, both leads possessed opposing lipophilic trends for the substituents found at the R_1 position. Increasing lipophilicity and bulk at R_1 seems to favor selectivity for the STAT3 protein. However, bulky hydrophobic substituents on the R_2 position seem to dictate the largest selectivity for the STAT5 protein, suggesting structural variability within the STAT3 and STAT5 SH2 domains.¹⁶ To exploit these observed differences, we probed the SH2 domain pocket using computational docking simulations.

While the activated STAT5–STAT5 dimer structure has yet to be solved, the pY binding pocket of the SH2 domain was identified by comparing and contrasting the structural architectures and binding sites of 121 SH2 domain-containing proteins.¹⁸ SH2 domains are defined by an antiparallel β -sheet flanked by two α helices. The pY binds in a pocket located within the βB , C, and D strands. Specifically, a conserved R(XXX) residue on the βB strand participates in electrostatic interactions with pY.¹⁹ Notably, STAT3's structure was resolved in both the unphosphorylated (PDB: 3CWB) and phosphorylated (PDB: 1BG1) states.^{20,21} Superimposition of structures revealed that STAT3's SH2 domain structure is not significantly altered upon phosphopeptide binding.²² Thus, we reasoned that the unphosphorylated STAT5 structure (PDB: 1Y1U) is suitable for *in silico*-based drug design. Analogous to the canonical pY-SH2 domain binding, the STAT5 pY likely docks proximal to the conserved R618 (βB strand), making H-bonding/electrostatic interactions with nearby polar residues, K600 (αA), T628 (βC), and S622 (βB and βC) (Figure 2A).

Next, using AutoDock4.2, we performed global searches of the conformational space along with careful local searches to derive the best conformational fit within the STAT5 SH2 domain. Docking simulations were carried out with a rigid

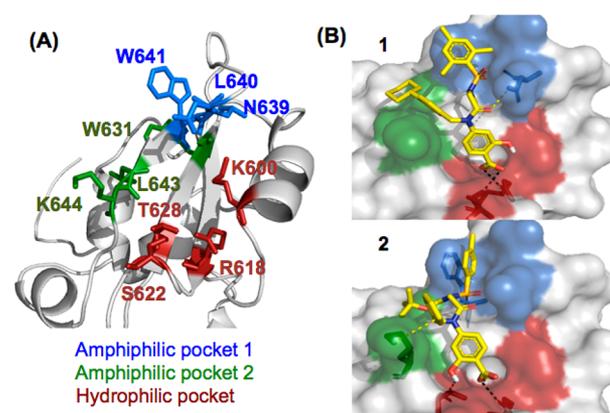


Figure 2. (A) STAT5a's SH2 domain with three binding pockets: hydrophilic, red; amphiphilic, green; amphiphilic, blue; (B) *in silico* docking of **1** interacting with R618, S622, and N639; **2** interacting with R618 and S622, as well as a cationic– π interaction of the R_1 benzyl with K644.

protein structure, allowing for ligand flexibility using a Lamarckian Genetic Algorithm (GA) with the global and adaptive local search parameters through 50 trials of the “long” GA runs. **1** and **2** occupied two amphiphilic pockets. The first pocket contained residues N639, L640, and W641 (blue region, Figure 2A,B), which interact with the sulfonamide mesityl and tolyl of **1** and **2**, respectively. The adjacent pocket containing W631, L643, and K644 (green) was found to interact with the R_1 substituents of both leads. Interestingly, for **2** ($R_1 = \text{Ph}$), docking poses showed favorable cationic– π interactions with K644. In contrast, the binding of **1** ($R_1 = \text{cyclohexylphenyl}$) was dominated by van der Waal's interactions with L643. Since the K644 residue is unique to STAT5, we elected to prepare analogs possessing hydrophobic, electron rich, aromatic R_1 substituents to derive selectivity. The R_3 substituents were carried forward from previous SAR studies, owing to their favorable biological profiles.

To survey the R_1 binding pocket, we explored 24 substituents of varying size and chemical diversity, including furan (**8a,m**), thiophene (**8b,n**), imidazole (**8c,o**), cyclopropyl (**8d,p**), and bicyclo[2.2.1]heptane (**8e,q**) heterocycles, naphthyl, and phenyl; these were chosen for their small size and rich electronic character, instead of the cyclohexylphenyl moiety of **1** (Table 1). The R_3 position was left either as a mesityl or pentafluorobenzyl substituent. The tolyl group of **2** was not retained due to poor solubility.

The library was screened through a previously reported high-throughput STAT5B FP assay which measures the disruption of phosphopeptide–STAT5B SH2 domain interactions.¹⁵ FP measurements were taken every 15 min for an hour against the fluorescein conjugated phosphopeptide–STAT5B complex to verify that the phosphopeptide probe (5-FAM-GpYLVDKW) was not being displaced over time.

Time intervals identified the optimal time point at which the maximum change in fluorescence polarization was observed. In general, the smaller heterocyclic R_1 substituents had no observable activity at inhibitor concentrations up to $100 \mu\text{M}$.

In contrast, only aromatic derivatives **8j**, **8l**, **8v**, and **8x** demonstrated similar potencies relative to the parent leads, with K_i values of 6.62, 4.77, 12.36, and $11.51 \mu\text{M}$, respectively (Supporting Information Figure S4 and Table S1). We noted that the electron rich, hydrophobic di-*tert* butyl benzyl groups of **8l** and **8x** consistently engaged in cation– π interactions with

Table 1. Focused Library of 24 Derivatives with Small Heterocycles and Substituted Phenyl Groups at the R₁ Position, with Corresponding K_i Values Determined through FP

R ₁	K _i (μM)	R ₁	K _i (μM)	R ₁	K _i (μM)
8a,m	>60	8e,q	>60	8i,u	>60
8b,n	>60	8f,r	>60	8j,v	6.62, 12.36
8c,o	>60	8g,s	>60	8k,w	>60
8d,p	>60	8h,t	>60	8l,x	4.77, 11.51

K644, more so than the corresponding heterocyclic derivatives. The naphthyl derivatives, **8j** and **8v**, favored π - π stacking interactions with W641 and not with K644 (Supporting Information Figures S9–12). Introduction of small heterocycles abolished activity. To investigate whether the *N*-boc R₂ substituent of **2** might contribute to STAT5 selectivity, we prepared a new series of tetrapodal inhibitors. Di-*tert*-butyl benzyl and naphthyl substituents at R₁ were retained for SAR analysis. *In silico* experiments showed that the boc group in **2** interacted with residues L643, W631, and W641 (Figure 2B, green region). Since this region contained predominantly electron-rich aromatic residues (W), we hypothesized that agents equipped with electron-deficient R₂ aromatic groups such as a *p*-chlorobenzyl may favor strong π - π stacking interactions. Thus, a focused library **13a–d** was prepared and evaluated via FP against both STAT5 and STAT3 at 1–60,000 nM (Supporting Information Figure S5). Most encouragingly, **13a** was found to potentially disrupt phosphopeptide–STAT5B interactions, K_i = 145 nM. Moreover, **13a** was 1000-fold more selective for STAT5B cf. STAT3, with a STAT3 K_i = 143 μM (Figure 3B). *In silico* docking simulations revealed that **13a**, via the *p*-chlorobenzyl and pentafluorobenzene groups, made favorable contacts with the two adjacent amphipathic pockets (Figure 3A, and Figure S13).

In addition to FP, surface plasmon resonance (SPR) binding experiments were conducted to measure the kinetic association and dissociation using a ProteOn XPR36 (Biorad) with full-length His-tagged, STAT5, and STAT3 (SignalChem) immobilized on a biosensor chip. Most encouragingly, inhibitors exhibited potent nM binding affinities for STAT5, and selectivity for the STAT5 cf. STAT3 (Supporting Information Figure S2). Compound **13a** exhibited the most potent K_D (k_{off}/k_{on}) of 42 ± 4 nM, with 7-fold selectivity for STAT5 cf. STAT3, K_D = 287 ± 29 nM (Figure 3C). As assessed by FP and SPR, **13a** represents the first nM STAT5-binding, STAT5-selective inhibitor.

To determine the selectivity of **13a** for STAT5, we screened for off-target kinase activity, a possible alternative target for an effector of STAT5 phosphorylation. **13a** was assessed against

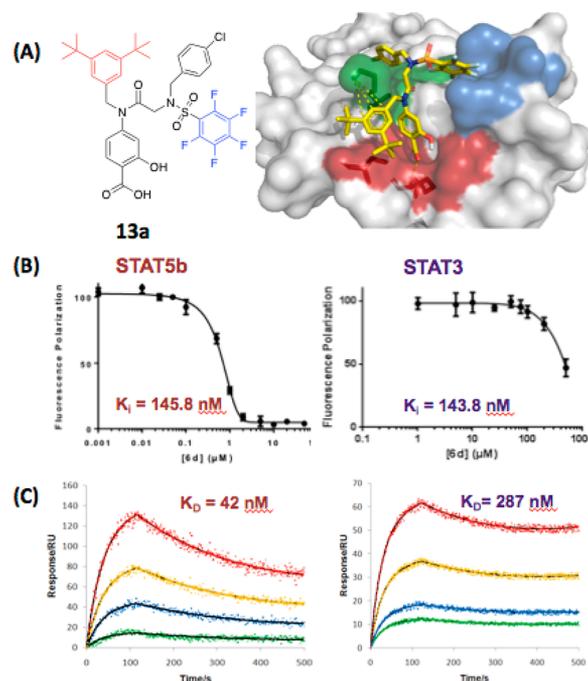


Figure 3. (A) **13a** docked with STAT5b; (B) FP binding traces for **13a** against STAT5b and STAT3 protein; (C) SPR curves for **13a** against STAT5b and STAT3 (S, 1.67, 0.56, 0.19 μM).

120 representative kinases covering the diversity of the kinase families (DiscoverX). Ultrasensitive quantitative PCR (qPCR) was used to measure levels of immobilized kinases after treatment with **13a** at 5 μM. Encouragingly, **13a** showed negligible effects against the bank of kinases and, more notably, against upstream STAT5 activating kinases, JAK1/2, ABL, and FLT-3 (Supporting Information Figure S1). These data suggest that inhibition of pSTAT5 is due to interaction with STAT5's SH2 domain and not through inhibition of upstream kinases.

Next, **13a** was assessed for whole cell potency against STAT5 transformed CML and AML cell lines, K562 (Bcr-Abl) and MV-4;11 (FLT3-ITD), respectively. Cell viability was assessed following treatment at various concentrations of inhibitor (0.78–50 μM) using a CellTiter-Blue cell viability assay (72 h). As compared to first generation STAT5 inhibitors (**8j**, **8l**, **8v**, **8x**), IC₅₀ values for **13a–d** were 2–3-fold higher in potency, with activities ranging from 3 to 20 μM (Supporting Information Figures S14, S15). Encouragingly, **13a** displayed the most potent activity in FLT3-ITD driven MV4;11 cells, IC₅₀ = 3.5 μM. We next evaluated **13a**-mediated inhibition of STAT5 phosphorylation levels. K652 leukemia cells were treated with **13a** for 5 h, the cells were harvested, and the levels of phosphorylated STAT5 (Y694) were determined (Figure 4A). **13a** decreased pSTAT5 in a dose dependent manner and ablated pSTAT5 above 10 μM with no change in the total STAT5 concentration or cleavable PARP-1. However, K562 cells were found to undergo cell death (cleaved PARP-1, Caspase-3) at 24 h at 15 μM, indicating that **13a** might induce apoptosis as a result of STAT5 inactivation (Figure 4B). To investigate selectivity, MDA-MB-231 breast cancer cells, which harbor high pSTAT3 and negligible pSTAT5 activity (Supporting Information, Figure S18), were assessed for differential pSTAT inhibition by **13a** (Figure 4C). Encouragingly, pSTAT3 was not inhibited at doses corresponding to pSTAT5 inhibition within the leukemic cell line and neither

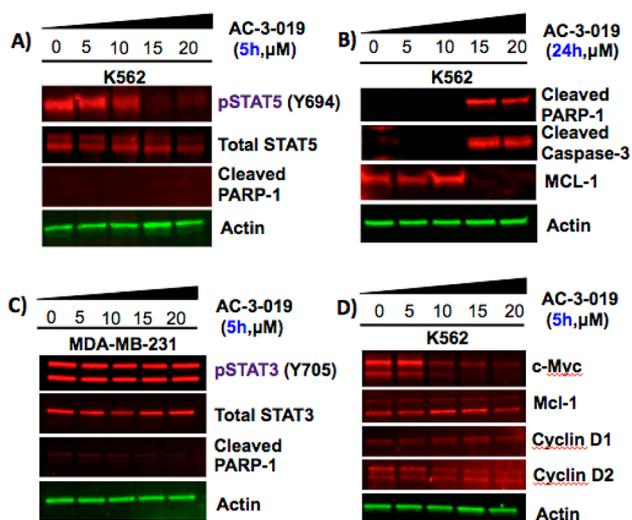


Figure 4. (A) Inhibition of pSTAT5 with 13a (15 μM) in K562 cells; (B) Initiation of apoptosis after 24 h and knockdown of MCL-1 (15 μM); (C) 13a shows no effect on pSTAT3 in MDA-MB-231 cells up to 20 μM ; (D) Knockdown of downstream target c-Myc after 5 h.

total STAT3 or STAT5 protein levels were affected. It was shown that 13a was 3-fold less cytotoxic in MDA-MB-231 ($\text{IC}_{50} = 10 \mu\text{M}$) than in the high pSTAT5 leukemic cell line (Supporting Information Figure S16). Downstream of STAT5, we assessed for modulation of the STAT5 transcriptional targets, MCL-1, CYCLIN D1/D2, and MYC. We reasoned that 13a should decrease gene expression and induce apoptosis by 24 h. K562 cells were dosed with 13a at the same concentrations observed for selective STAT5 inhibition. At 5 h, we observed dose-dependent decreases in MYC and complete knockdown of MCL-1 observed at 24 h at 15 μM (Figure 4D).

We evaluated 13a in healthy human CD34⁺ umbilical cord cells to determine off-target effects and the therapeutic window. Most encouragingly, there existed an approximate order of magnitude difference in sensitivity, with little reduction in cell viability at 10 μM 13a, while MV4-11 cells were abolished at the same concentrations (Figure 5).

In summary, we have identified the first nM binding inhibitor of STAT5 protein. Moreover, lead compound 13a has been shown to potently and selectively disrupt STAT5-phosphopeptide interactions (nM) as compared to STAT3 (μM). With no off-target kinase activity, 13a was shown to suppress pSTAT5

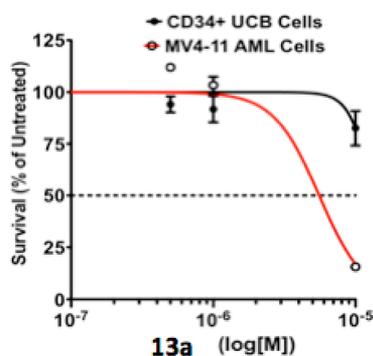


Figure 5. Healthy human CD34⁺ umbilical cord cells treated with compound 13a at 10 μM with limited effect on cell viability in comparison to MV4;11 cells.

while having no effect on pSTAT3. Thus, 13a represents a useful probe for interrogating the STAT5 function in diseased cells as well as being a candidate for advanced preclinical trials.

■ ASSOCIATED CONTENT

Supporting Information

Inhibitor synthesis, characterization, and experimental procedures for FP, SPR, cytotoxicity, and Western Blot studies. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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Author Contributions

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Notes

The authors declare no competing financial interest.

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