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## Targeting Protein Tyrosine Phosphatase SHP2 for the Treatment of *PTPN11*- Associated Malignancies

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### Abstract

Activating mutations in *PTPN11* (encoding SHP2), a protein tyrosine phosphatase (PTP) that plays an overall positive role in growth factor and cytokine signaling, are directly associated with the pathogenesis of Noonan syndrome and childhood leukemias. Identification of SHP2 selective inhibitors could lead to the development of new drugs that ultimately serve as treatments for *PTPN11*-associated diseases. As the catalytic core of SHP2 shares extremely high homology to those of SHP1 and other PTPs that play negative roles in cell signaling, to identify selective inhibitors of SHP2 using computer-aided drug design, we targeted a protein surface pocket that is adjacent to the catalytic site, is predicted to be important for binding to phosphopeptide substrates, and has structural features unique to SHP2. From computationally selected candidate compounds, #220–324 effectively inhibited SHP2 activity with an IC<sub>50</sub> of 14 μM. Fluorescence titration experiments confirmed its direct binding to SHP2. This active compound was further verified for its ability to inhibit SHP2-mediated cell signaling and cellular function with minimal off-target effects. Furthermore, mouse myeloid progenitors with the activating mutation (E76K) in *PTPN11* and patient leukemic cells with the same mutation were more sensitive to this inhibitor than wild-type cells. This study provides evidence that SHP2 is a “druggable” target for the treatment of *PTPN11*-associated diseases. Since the small molecule SHP2 inhibitor identified has a simple chemical structure, it represents an ideal lead compound for the development of novel anti-SHP2 drugs.

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### AUTHOR CONTRIBUTIONS

B.Y., W.L., W.M.Y., S.A., O.G. performed the experiments and summarized the data. A.D.M., L.D.T., and C.K.Q. designed the experiments. M.L. provided critical reagents. W.L., O.G., A.D.M., and C.K.Q. wrote the manuscript.

## Keywords

*PTPN11* (SHP2); Phosphatase; Inhibitor; Drug development; Leukemia

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## Introduction

SHP2 (encoded by *PTPN11*), a ubiquitously expressed SH2 domain-containing protein tyrosine phosphatase (PTP), is involved in multiple cell signaling processes, such as the Ras-Erk, PI3K-Akt, Jak-Stat, NF- $\kappa$ B, and mTOR pathways (1–3). It contains two tandem SH2 domains at the N-terminus and a PTP domain at the C-terminus, with flexible polypeptide linker regions connecting the three domains. The SH2 domains, in particular, the N-terminal SH2 (N-SH2) domain, mediate the binding of SHP2 to other signaling proteins via phosphorylated tyrosine residues in a sequence-specific fashion (4, 5). This directs SHP2 to the appropriate subcellular location and helps determine the specificity of substrate-enzyme interactions. The X-ray crystal structure of SHP2 reveals the formation of an intramolecular interface between the N-SH2 domain and the PTP domain (6–8). This self-interaction blocks substrate access to the catalytic site, and polypeptide ligands with phospho-tyrosine (pY) residues activate SHP2 by binding the SH2 domains (primarily the N-SH2 domain), thereby disrupting the self-inhibitory interaction between N-SH2 and PTP domains and exposing the phosphatase catalytic site (6–8). Intriguingly, despite its direct function in protein dephosphorylation, SHP2 plays an overall positive role in transducing signals initiated from growth factors/cytokines and extracellular matrix proteins (1–3). However, the signaling mechanisms of SHP2 are still not completely understood. In particular, the molecular basis for the positive role of its catalytic activity in the Ras-Erk pathway as well as other pathways remains elusive.

Germline or somatic mutations in *PTPN11* (encoding SHP2) that cause hyperactivation of SHP2 catalytic activity have been identified in the developmental disorder Noonan syndrome and various childhood leukemias. Specifically, heterozygous *PTPN11* mutations have been found in 50% of patients with Noonan syndrome, as well as in 35% of juvenile myelomonocytic leukemia (JMML), 10% of myelodysplastic syndrome, 7% of B cell acute lymphoblastic leukemia/lymphoma, and 4% of acute myeloid leukemia cases (9–13). Additionally, activating mutations of *PTPN11* have been identified in sporadic solid tumors (14). Such mutations disrupt the inhibitory intramolecular interaction of SHP2, leading to gain-of-function by allowing constitutive access to the phosphatase catalytic site on the enzyme (12, 15). *PTPN11* mutations and other JMML-associated *Ras*, *c-CBL*, or Neurofibromatosis1 mutations are mutually exclusive (9–13, 16, 17). Remarkably, recent studies have demonstrated that single *PTPN11* gain-of-function mutations are sufficient to induce Noonan syndrome, JMML-like myeloproliferative disease, and acute leukemias in mice (18–23), suggesting that the *PTPN11* mutations play a causal role in the development of these diseases.

The direct connection between hyperactivation of SHP2 and human diseases points to SHP2 as a potential target for mechanism-based therapeutics. Selective and potent SHP2 inhibitors are needed. Discovery of SHP2-specific inhibitors would not only facilitate research on SHP2 signaling in model systems but could also lead to the development of new drugs for *PTPN11*-associated diseases. However, little progress has been made in this field because the discovery of SHP2-specific inhibitors is complicated by the extremely high homology of the catalytic core of SHP2 with those of SHP1 and other PTPs that play negative roles in growth factor and cytokine signaling. To overcome this we previously applied computer-aided drug design (CADD) to screen a virtual database of compounds targeting a protein surface pocket that is adjacent to the catalytic site of SHP2, is predicted to be important for

binding to phosphopeptide substrates, and has structural features unique to SHP2. This effort led to the identification of several low-molecular weight compounds capable of inhibiting SHP2 (24). In the present study we expand that effort by using CADD similarity searching to identify analogs of the previously discovered active compounds that may have improved activity, leading to the identification of a new compound (#220–324) that is superior to the previously reported parent inhibitors in terms of selectivity and potency. This new compound selectively inhibited SHP2 catalytic activity without significant off-target effects. Moreover, it was efficacious in blocking SHP2-mediated cellular functions, even in *PTPN11* mutated mouse and human leukemia cells, providing evidence that SHP2 is a “druggable” target for the treatment of *PTPN11*-associated diseases.

## Materials and Methods

### Reagents and Chemicals

RPMI 1640, IMDM, DMEM, and the penicillin-streptomycin stock solution were purchased from Thermo Scientific/HyClone Laboratories (Logan, UT). Fetal bovine serum (FBS) was obtained from Invitrogen (Grand Island, NY). The CellTiter 96<sup>®</sup> Aqueous One Solution Cell Proliferation Assay kit was purchased from Promega (Madison, WI). Antibodies against p-Erk (E-4), Erk (C-16), Jak2 (C-20), and SHP2 (C-18) were purchased from Santa Cruz Biotechnology, Inc (Santa Cruz, CA). Antibodies against p-Akt (Ser473), Akt, and p-Stat5 (Y694) were obtained from Cell Signaling Technology (Beverly, MA). Antibodies against p-Jak2 (Tyr1007/1008) and phospho-tyrosine were purchased from Millipore Corporation (Temecula, CA). Antibody against Stat5 was purchased from BD Biosciences (San Jose, CA). Goat-anti-rabbit and anti-mouse IgG horseradish peroxidase (HRP) conjugates were purchased from Jackson ImmunoResearch Laboratories, Inc. (West Grove, PA). DMSO and other chemicals used in buffer solutions were purchased from Fisher Scientific (Pittsburgh, PA). #220–324 was obtained from Specs (Wakefield, RI) and dissolved in DMSO to prepare stock solutions (20 mM) for subsequent experiments.

### CADD Similarity Searching

Identification of compounds similar to the published lead compounds used the program MOE (Chemical Computing Group, Inc., Montreal, Canada). An in house database of 1.3 million commercially-available compounds was screened using the BIT MACCS structural fingerprints, with the extent of similarity quantified using the Tanimoto coefficient (25). For each active compound the top 50 scoring compounds were considered for experimental evaluation.

### Computational Modeling of the Binding Pose of #220–324

#220–324 was built into the previously-predicted binding site (24) using coordinates of analogous atoms from the –previously-docked pose of the parent compound #220 (24) and maintaining SHP2 coordinates as per this previous docked complex. The protein-ligand system was represented using the CHARMM36 protein force field (26) and the CHARMM General Force Field (CGenFF) (27) for the ligand in combination with the GBMV Generalized Born implicit solvent model (28, 29), with automatic assignment of atom types and missing parameters for #220–324 generated via the CGenFF engine (30, 31) as implemented in the ParamChem web portal. Molecular dynamics (MD) simulations were done with a 1.5 fs integration timestep, the SHAKE algorithm (32) to constrain bonds to hydrogen atoms to their equilibrium values, an 18-Å spherical cutoff for non-bonded interactions, with a switching function (33) applied in the 16–18 Å interval, and Langevin thermostating (34) at 298 K. The CHARMM software (35) was used for all system construction and simulation. Images were generated using VMD (36).

### **In Vitro Phosphatase Activity Assay**

Glutathione S-transferase (GST) fusion proteins of SHP2 purified in-house were used as the enzyme and a phosphopeptide corresponding to the surrounding sequence of pTyr<sup>1146</sup> in the insulin receptor (Thr-Arg-Asp-Ile-Tyr[PO<sub>3</sub>H<sub>2</sub>]-Glu-Thr-Asp-Tyr-Tyr) was used as the substrate. The assay determines free phosphate generated by dephosphorylation of the substrate using the Malachite Green reagent (Sigma, St. Louis, MO). Briefly, 0.5 µg of GST-SHP2 PTP was incubated in 40 µL assay buffer (25 mM Tris-HCl, pH 7.4, 50 mM NaCl, 5 mM DTT, and 2.5 mM EDTA) with test compounds at various concentrations at room temperature for 30 min. The substrate was then added to a final concentration of 0.2 mM. The system was incubated at 30°C for 30 min. Finally, 50 µL of Malachite Green solution was added and OD<sub>620</sub> was measured after 10 min. The protocols for the phosphatase assays for SHP1, CD45, LAR, MEG2, and TC-PTP were similar, with the exception that GST-SHP1, GST-CD45 cytoplasmic domain, GST-LAR, GST-MEG2, and GST-TC-PTP enzymes purchased from Biomol International, L.P. (Plymouth Meeting, PA), were used in the respective assays. For IC<sub>50</sub> determinations, 5 concentrations of #220–324 were tested. Each experiment was performed in triplicate.

### **Fluorescence Titrations**

For all experiments, purified SHP2 PTP domain GST-fusion protein was diluted into 20 mM Tris-HCl, pH 7.5. Fluorescence spectra were recorded with a Luminescence Spectrometer LS50 (Perkin-Elmer, Boston, MA). Titrations were performed by increasing the test compound concentrations while maintaining the SHP2 protein concentration at 3 µM. Contributions from background fluorescence of the inhibitor were accounted for by subtracting the fluorescence of the inhibitor alone from the protein-inhibitor solution. The excitation wavelength was 295 nm and fluorescence was monitored from 360 to 500 nm. All reported fluorescence intensities were relative values and were not corrected for wavelength variations in detector response.

### **Cell Proliferation Assay**

Ba/F3 cells and mouse embryonic fibroblasts (MEFs) were seeded into 96-well plates at a density of 1×10<sup>4</sup> cells/well (Ba/F3) and 5×10<sup>3</sup> cells/well (MEFs) in RPMI-1640 containing 10% FBS plus recombinant mouse IL-3 (1.0 ng/mL) and DMEM containing 10% FBS, respectively. Cells were grown overnight and then treated with either test compounds or the same concentrations of DMSO. Three days later, the number of viable cells was determined using a CellTiter 96<sup>®</sup> Aqueous One Solution Cell Proliferation Assay kit.

### **Western Blotting**

Ba/F3 cells were starved in serum and cytokine-free RPMI1640 overnight. Cells were then treated with #220–324 for 3 hrs prior to IL-3 stimulation. Stimulated cells were harvested and lysed on ice with RIPA buffer containing 50 mM Tris-HCl, pH 7.4; 1% NP-40; 0.25% Na-deoxycholate; 150 mM NaCl; 1 mM EDTA; 1 mM NaF; 1 mM Na<sub>3</sub>VO<sub>4</sub>; 1 mM PMSF and protease inhibitor cocktail (Roche, Indianapolis, IN). Equivalent amounts of protein (50 µg) were resolved on 10% SDS-PAGE and transferred to nitrocellulose membranes (Millipore, Bedford, MA). Membranes were blocked with 2% BSA in TBS-T [20 mM Tris-HCl (pH 7.4), 150 mM NaCl, and 0.1% Tween 20] for 1 hr at room temperature and probed with primary antibodies overnight at 4°C. Blots were washed with TBS-T and exposed to HRP conjugated goat-anti-mouse or goat-anti-rabbit secondary antibodies for 1 hr at room temperature. Immunoreactive bands were detected by using ECL Plus Reagents (GE Healthcare, Piscataway, NJ).

## Colony-Forming Unit Assay

Freshly harvested mouse bone marrow cells or patient splenocytes ( $2 \times 10^4$  cells/mL) were assayed for colony forming units (CFUs) in 0.9% methylcellulose IMDM containing 30% FBS, glutamine ( $10^{-4}$  M),  $\beta$ -mercaptoethanol ( $3.3 \times 10^{-5}$  M), GM-CSF (1.0 ng/mL), and varying doses of #220–324 or the same concentrations of DMSO. After 7 days (mouse bone marrow cells) or 14 days (patient cells) of culture at 37°C in a humidified 5% CO<sub>2</sub> incubator, myeloid colonies (CFU-GM and CFU-M) were counted under an inverted microscope.

## Results

### Identification of #220–324 as a novel selective SHP2 inhibitor

As reported in our previous study, by targeting a putative phosphopeptide binding site adjacent to the catalytic site that has structural features unique to SHP2, nine active compounds were identified by CADD chemical database screening combined with experimental assays (24). To identify more potent and selective SHP2 inhibitors, in the present work we extend those efforts by performing a similarity search of a chemical database of 1.3 million compounds based on the structural characteristics of the 9 active compounds following the procedures previously described (37). From the top scoring compounds sixty-two candidate compounds were obtained and subjected to biological screening. Twenty of these compounds inhibited SHP2 catalytic activity with various efficiencies in biochemical assays (Supplementary Fig. 1A and 1B). Among these inhibitors, #220–324 was found to be the most potent and most selective in subsequent cell-based screenings that test the effects of the compounds on IL-3-induced cell proliferation (data not shown). This compound inhibited SHP2 catalytic activity with an IC<sub>50</sub> of 13.88  $\mu$ M (Fig. 1A). To determine functional specificities of #220–324, its inhibitory activity was assessed against several mammalian PTPs, in particular those phosphatases that are highly expressed in hematopoietic cells. As shown in Fig. 1B, #220–324 was approximately 3-fold more selective for SHP2 compared to SHP1, and 13-, 10-, 13-, 3-fold more selective for SHP2 compared to CD45, LAR, MEG2, and TC-PTP, respectively.

Molecular dynamics (MD) simulations were performed on the SHP2:#220–324 complex to test the plausibility of a non-catalytic site binding pose for this inhibitor. Using the previously-docked coordinates of the parent compound #220 as the basis for a starting conformation, the SHP2:#220–324 complex was first minimized for 1000 steps and then allowed to evolve unrestrained for 70,000 steps (105 ps) of generalized Born MD at 298 K. Like the parent compound in its docked pose (24), after the MD refinement #220–324 makes extensive hydrophobic contact with SHP2 (Supplementary Fig. 2). Additionally, the predicted binding site of SHP2 undergoes conformational changes to accommodate some relaxation of this compound relative to its initial conformation, such that after MD refinement, the new SHP2 conformation results in unfavorable steric clashes with the original docked pose of the parent compound #220. These results suggest a conformationally-flexible binding pocket adjacent to and distinct from the catalytic site that is capable of binding drug-like small molecules with substantial hydrophobic moieties, like #220–324. The results also provide a potential explanation for the increased activity of #220–324 over the parent compound #220.

### #220–324 preferentially binds to the activated form of SHP2

In the basal state, SHP2 is self-inhibited by hydrogen bonding of the N-SH2 domain loop to the deep pocket of the PTP domain. Tumor-associated mutations of *PTPN11* result in amino acid changes at the interface formed by the N-SH2 and PTP domains, disrupting the inhibitory intramolecular interaction, leading to hyperactivation of SHP2 catalytic activity

(12, 15). The E76K mutation, localized in the N-SH2 domain is the most common and most active mutation found in leukemias (38, 39). To further characterize the mechanisms of action of #220–324, we purified GST fusion proteins of the SHP2 PTP domain, wild-type (WT) full-length SHP2 and the full-length SHP2 E76K mutant, and compared the sensitivities of these SHP2 recombinant proteins to #220–324. #220–324 inhibited the SHP2 E76K mutant in a dose-dependent manner, with an  $IC_{50}$  of 23.81  $\mu$ M that is 1.5-fold higher than that for the SHP2 PTP domain (13.88  $\mu$ M). However, the  $IC_{50}$  for SHP2 E76K is 2.7 fold lower than that for full-length WT SHP2 (64.74  $\mu$ M) (Fig. 1C). The observation that full-length SHP2 is less sensitive to #220–324 than the SHP2 PTP domain and that the SHP2 E76K mutant is more sensitive than WT SHP2 to this inhibitor indicates that the N-SH2-PTP domain interaction interferes with #220–324 binding to SHP2 and that this compound preferentially inhibits the activated form of SHP2. The activity of #220–324 was further characterized by testing its effect on SHP2 catalytic activity using *para*-Nitrophenylphosphate (*p*NPP), a general small compound substrate for all PTPs. As demonstrated in Fig. 1D, #220–324 did not inhibit the SHP2 dephosphorylation of this small substrate regardless of which form of the SHP2 protein was used as the enzyme. This result suggests that #220–324 inhibits SHP2 dephosphorylation of the phosphopeptide substrate by disrupting binding of the phosphopeptide to SHP2 at a location other than the catalytic core.

To validate that #220–324 was binding directly to the SHP2 protein, we determined whether binding of the inhibitor altered the fluorescence of the SHP2 PTP domain using the fluorescence quenching assay taking advantage of the four tryptophans in the PTP domain. Of these residues, Trp248 is located 15 Å from the drug docking pocket and Trp423 located 8 Å from the phosphatase active site. As shown in Fig. 1E, #220–324 exhibited strong quenching of SHP2 fluorescence in a dose-dependent manner. These fluorescence quenching experiments indicate that #220–324 binds directly to SHP2, thereby attenuating its activity toward the phosphopeptide substrate.

### Effects of #220–324 on SHP2-mediated cellular function

Given the potency and selectivity of #220–324 toward SHP2 *in vitro*, we proceeded to evaluate its ability to inhibit SHP2-mediated signaling and cellular function. *PTPN11*-associated leukemia is characterized by hypersensitivity of myeloid progenitors to GM-CSF and IL-3 (18, 20–22). Our previous studies have demonstrated that SHP2, in contrast to other PTPs, especially SHP1, promotes IL-3-triggered cell proliferation and activation of Erk, Akt, and Jak2-Stat5 pathways (40). To verify the selectivity of #220–324 in cells, it was critical to assess the cellular activities of this compound in GM-CSF or IL-3-induced cellular responses. IL-3-dependent Ba/F3 cells were therefore used. As shown in Fig. 2A, treatment of Ba/F3 cells with the compound for 72 hrs greatly decreased cell growth in a concentration-dependent manner, with an  $IC_{50}$  of 37.26  $\mu$ M, consistent with the overall positive role of SHP2 catalytic activity in cellular response to IL-3 (40). Furthermore, as the SHP2 signaling is known to play a positive role in the cell cycle progression (41), we performed cell cycle analysis in #220–324 treated Ba/F3 cells. We found that this compound inhibited cell cycle progression, specifically the G<sub>1</sub>-S transition, in a dose-dependent manner (Fig. 2B). To evaluate whether #220–324 inhibited SHP2-dependent cell signaling, we examined the effects of this inhibitor on IL-3-induced Erk, Akt, Jak2, and Stat5 activation. Ba/F3 cells were serum-starved overnight followed by pretreatment with #220–324 or DMSO for 3 hrs, and then stimulated with IL-3. As shown in Fig. 2C, IL-3 induced activation of Erk, Akt, Jak2, and Stat5 determined by their phosphorylation levels was greatly suppressed by #220–324 at all time points. Furthermore, Ba/F3 cells were preincubated with #220–324 at various concentrations, followed by IL-3 stimulation for 10 min. We observed that #220–324 inhibited IL-3 induced Erk, Akt, and Stat5 activation in a dose-dependent manner (Supplementary Fig. 3A), in agreement with previous findings that

SHP2 catalytic activity is required for optimal activation of these signaling pathways elicited by IL-3 (40). In addition, we found that tyrosyl phosphorylation of Gab2 (carried out by Jak2 kinase), an important intermediary of the PI3K-Akt pathway (42, 43), and the interaction between SHP2 and phosphorylated Gab2 was markedly decreased by #220–324 (Supplementary Fig. 3B). Taken together, these results imply that IL-3 signaling, including Ras-Erk, PI3K-Akt, and Jak2-Stat5 pathways, was down-regulated through a mechanism involving the inhibition of SHP2 activity, and point to the possibility that #220–324 mainly inhibits the catalytic activity of SHP2, not those of other PTPs.

#### #220–324 has minimal off-target effects

To validate that the effects of #220–324 on cell growth were mediated through inhibiting SHP2, we tested #220–324 in *PTPN11* knockout cells. *PTPN11* conditional knockout (*PTPN11<sup>Fl/Fl</sup>/Ade-Cre+*) and WT control (*PTPN11<sup>+/+</sup>/Ade-Cre+*) mouse embryonic fibroblasts (MEFs) were treated with #220–324 at various concentrations or control DMSO for 72 hrs, and cell growth rates were determined. As shown in Fig. 3A, proliferation of WT cells was significantly decreased by the SHP2 inhibitor, consistent with the positive role of SHP2 in growth factor signaling (1–3). However, SHP2-depleted MEFs showed only minimal inhibitory effects. We also treated knockout and WT control cells with #220–324 at 10  $\mu$ M or DMSO for various times and obtained similar results (Fig. 3B). The observation that *PTPN11* deficient cells were insensitive to #220–324 strongly suggests that this compound is an agent capable of inhibiting SHP2 in cells without significant off-target effects.

#### Mutant cells harboring hyperactivated SHP2 are more sensitive to #220–324

We next assessed effects of #220–324 on mutant cells with leukemia-associated *PTPN11<sup>E76K</sup>* mutation. As illustrated in Fig. 4A, treatment of both WT and *PTPN11<sup>E76K/+</sup>* MEFs with the compound for 72 hrs decreased cell growth in a dose-dependent manner. Intriguingly, *PTPN11<sup>E76K/+</sup>* mutant cells were more sensitive to SHP2 inhibition. We also compared the effects of #220–324 in WT and *PTPN11<sup>E76K/+</sup>* myeloid progenitors using CFU assays. In the presence of GM-CSF, the colony-forming capabilities of *PTPN11<sup>E76K/+</sup>* myeloid progenitors were more sensitive to #220–324 compared to those of WT counterparts (Fig. 4B). Additionally, we found that the cytokine-independent colony formation of *PTPN11<sup>E76K</sup>* mutant myeloid progenitors, a characteristic of *PTPN11*-associated JMML and other myeloid malignancies, was significantly suppressed by this inhibitor (Fig. 4C). Additionally, we tested for growth inhibitory function of #220–324 in human lung cancer cell lines. Both H661 and H596 cell lines are WT in *K-Ras* and epidermal growth factor receptor. However, H661 cells carry an activating mutation (N58S) in *PTPN11* while H596 cells have WT *PTPN11*. As demonstrated in Fig. 5A and 5B, #220–324 showed a marked inhibitory effect in H661 cells, but it was less effective in H596 cells. Together, these results further support that #220–324 is capable of decreasing proliferative advantages associated with gain-of-functions in SHP2 and that *PTPN11* mutated cells are more sensitive to SHP2 inhibition.

#### #220–324 corrects the hypersensitive growth pattern of JMML myeloid progenitors with the *PTPN11<sup>E76K</sup>* mutation

Bone marrow cells from patients with JMML display a characteristic hypersensitive growth pattern, resulting in greater production of granulocyte-macrophage colonies in response to GM-CSF (44, 45). As shown in Fig. 6A, in the presence of GM-CSF, colony formation of myeloid blast cells from JMML patients with the *PTPN11<sup>E76K/+</sup>* mutation was sensitive to #220–324 (IC<sub>50</sub> = 7.24  $\mu$ M). Consistent with this data, the total cell number of JMML cells cultured in GM-CSF-containing liquid medium was greatly decreased by the treatment of this inhibitor, with an IC<sub>50</sub> = 3.09  $\mu$ M (Fig. 6B). By contrast, we did not observe significant

inhibition of healthy donor myeloid progenitors by #220–324 under 100  $\mu$ M (Fig. 6C). The finding that #220–324 is able to selectively overcome the dominant effects of oncogenic SHP2 E76K in primary human leukemia cells provides strong support for the notion that SHP2 is a useful therapeutic target for the treatment of *PTPN11* mutation-associated JMML.

## Discussion

Identification of a lead inhibitor of SHP2 would greatly facilitate the development of anti-SHP2 therapeutic drugs. Mutations in *PTPN11* (encoding SHP2) that cause enhanced catalytic activity of SHP2 have been found in the patients with Noonan syndrome, childhood leukemias, and solid tumors (9, 12–14), and more importantly, these mutations appear to play a causative role in the development of JMML and acute leukemias (18–23). Selective and potent SHP2 inhibitors have the potential to be further developed into therapeutic drugs for these diseases. Previous studies have reported several inhibitors of SHP2 (46–49). However, none of those compounds have been well characterized, for example, their potential off-target effects and their functions in suppressing GM-CSF/IL-3 –induced signaling and cellular responses (that are known to be associated with the pathogenesis of *PTPN11*-associated JMML) have not been determined. More importantly, it is not clear whether those compounds have any potential therapeutic benefits in *PTPN11* mutated human or mouse leukemia cells. In this study, we have now identified and extensively characterized a novel small molecule SHP2 inhibitor.

Targeting the phospho-peptide binding site in SHP2 appears to be a feasible approach to developing SHP2 selective inhibitors. Despite a great need, little progress has been made in the development of SHP2 inhibitors. One of the challenges is the specificity of potential inhibitors. The active site of SHP2, which catalyzes de-phosphorylation of phospho-tyrosine (pY) in pY-containing peptides, is almost identical to that of the SHP1 phosphatase and shares high homology to those of other PTPs, such that compounds that target the catalytic core of SHP2 often inhibit other phosphatases. Therefore, our initial structure-based approach was based on targeting a non-homologous pocket in the pY-peptide binding cleft in SHP2 adjacent to the homologous catalytic site (24). This non-homologous pocket has unique features in both amino acid composition and 3D structure. Using CADD database screening targeting this pocket in combination with experimental assays, we previously identified 9 SHP2 selective inhibitors (24) although the activities of these inhibitors were low. In the present study, we have used CADD to identify compounds that are structurally similar to those 9 active compounds. Of these, 20 were shown to be active, of which #220–324 was shown to have a much improved potency and selectivity over the previously identified compounds. This small-molecule inhibitor selectively inhibited SHP2 over SHP1 and other PTPs. Most notably, it does not seem to have significant off-target effects as it only had minimal effects in *PTPN11* knockout cells. Additional biological analyses validated that it was effective in blocking SHP2-mediated signaling and cellular function. This inhibitor was verified to directly bind to the SHP2 protein by the fluorescence quenching assay. Since it only inhibited SHP2 enzymatic activity toward the pY-peptide substrate but not the small molecule substrate pNPP, it is likely that it functions as designed by disrupting the interaction between pY-peptide substrates and SHP2 at a location distinct from the catalytic core.

This study also provides proof of concept that SHP2 is a “druggable” target for the treatment of *PTPN11*-associated malignancies. SHP2 phosphatase plays an overall positive role in cell signaling and cellular function (1–3). Loss of SHP2 function leads to severe consequences in embryogenesis and hematopoietic cell development (50–52). A prevalent question is whether SHP2 can be used as a drug target for therapeutic purposes. Our studies have now provided evidence that mutant MEFs or lung cancer cells with activating mutations of



*PTPN11* were more sensitive to SHP2 inhibition than control cells with WT *PTPN11*, and the sensitivity of mutant mouse myeloid progenitors with the *PTPN11*<sup>E76K</sup> mutation was greater to SHP2 inhibition than WT counterparts. The same is true in human JMML cells. Leukemic cells from JMML patients with the *PTPN11*<sup>E76K</sup> mutation were more sensitive to the SHP2 inhibitor than healthy donor cells. These observations suggest that SHP2 can be targeted by inhibitory drugs for the control of *PTPN11* mutation-associated malignancies.

While #220–324 is a drug-like small molecule that is bioavailable in cells, its affinity is not high, with an IC<sub>50</sub> in the range of 10–20 μM. We tested the activity of #220–324 in the presence of the detergent Triton X-100. We found that the inhibitory function of #220–324 was decreased by ~30% in the presence of Triton X-100 at the concentration of 0.001% (Supplementary Fig. 4), raising the possibility that the interaction between #220–324 and SHP2 is weak and that this binding may be partially mediated by a promiscuous effect. Accordingly, further studies are required to optimize this compound to improve the binding to SHP2 in order to develop new SHP2 inhibitors with increased activities and selectivity suitable for pre-clinical and clinical studies. Such efforts would be greatly facilitated by co-crystallization of this compound with SHP2 that would reveal structural details of the binding interaction that would be used to guide optimization of the lead compound.

## Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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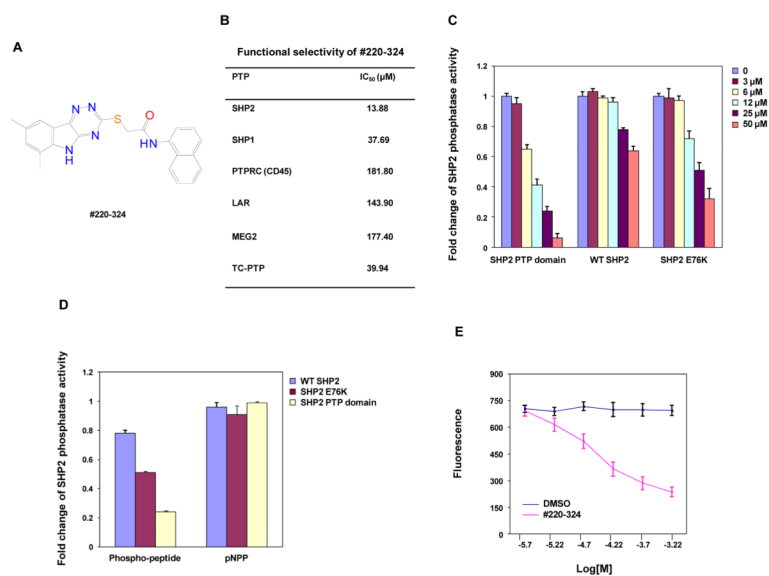
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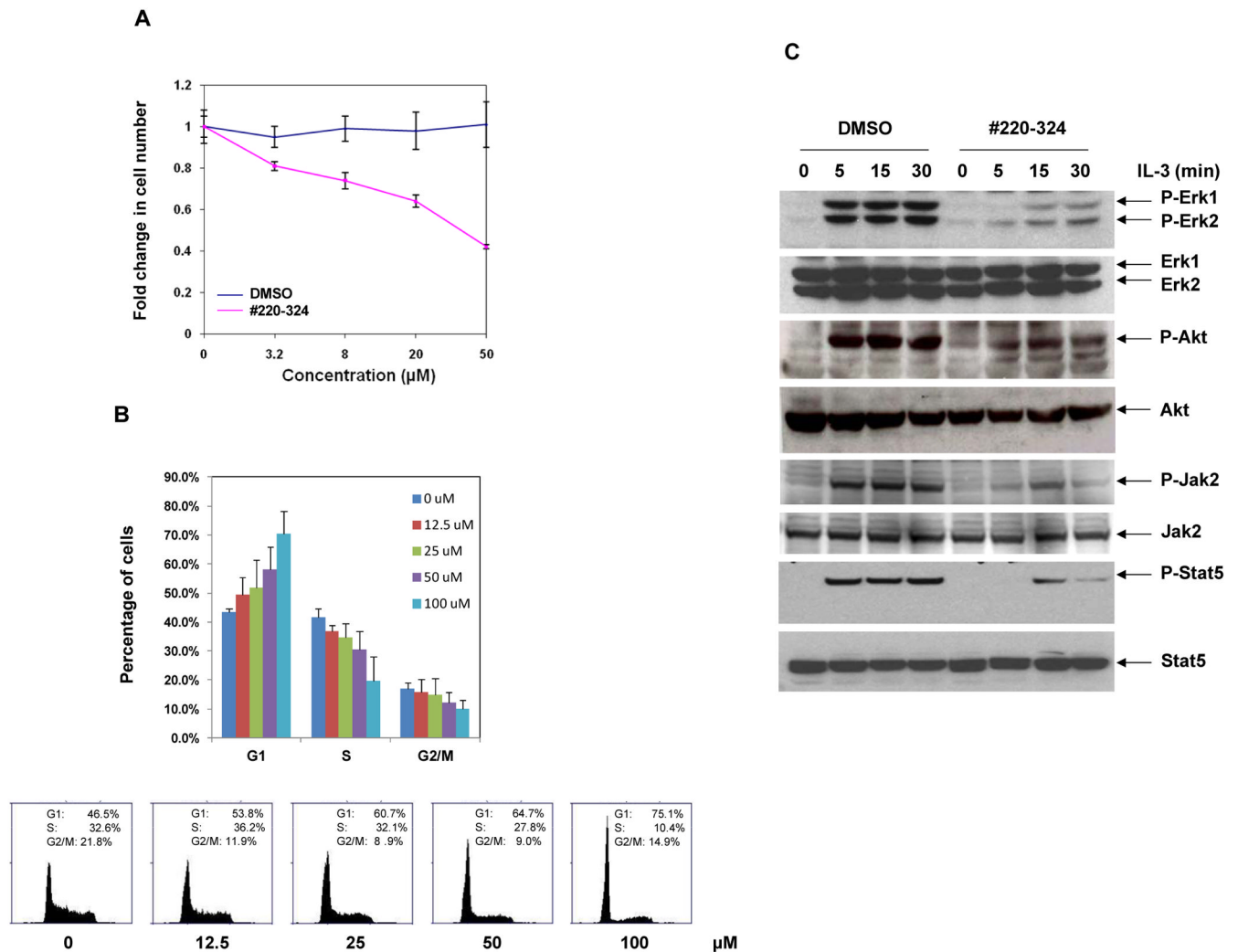
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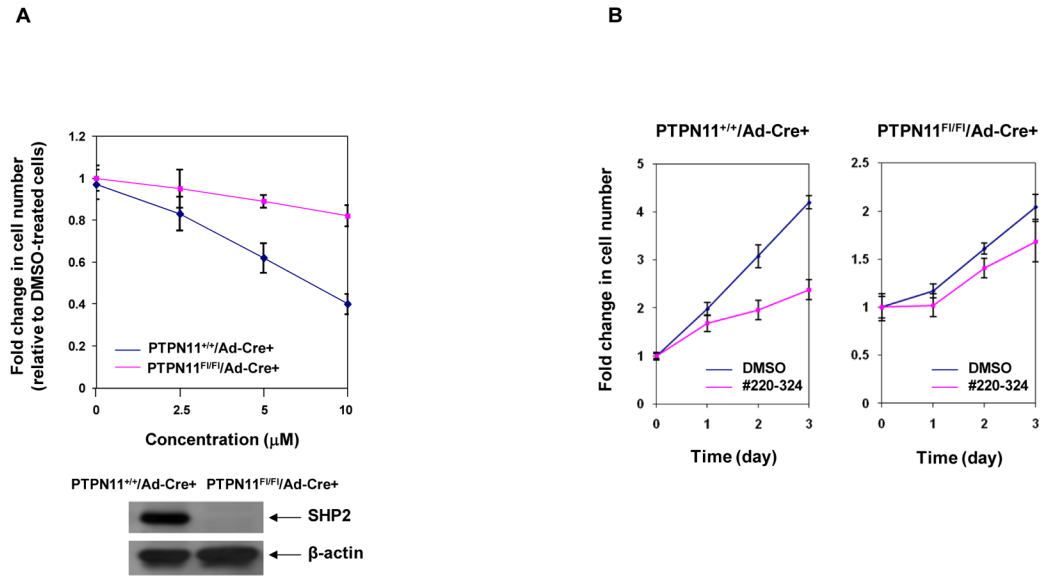


**Figure 1. #220–324 selectively inhibited the phosphatase activity of SHP2**

(A) Chemical structure of #220–324. (B) Phosphatase assays were carried out using the indicated phosphatases as enzymes and the phospho-insulin receptor peptide as a substrate in the presence of various concentrations of #220–324, as described in Materials and Methods. Experiments were performed three times with similar results obtained. Representative results from one experiment are shown. (C) PTP activities of the SHP2 PTP domain, WT full length SHP2, and full length SHP2 E76K were determined in the absence or presence of #220–324 at the indicated concentrations using the *in vitro* phosphatase assay. Experiments were repeated three times. Similar results were obtained in each. Data shown are mean±S.E.M. from one experiment. (D) Inhibitory effects of #220–324 (25 μM) on the enzymatic activities of the SHP2 PTP domain, WT SHP2, and SHP2 E76K were determined using the phospho-insulin receptor peptide or *p*NPP as the substrates. Experiments were repeated three times. Similar results were obtained in each. Data shown are mean±S.E.M. from one experiment. (E) Fluorescence titration of SHP2 was performed by increasing the concentrations of #220–324 while maintaining the SHP2 protein concentration. The fluorescence is plotted against the log concentration in mol/L (Log [M]) for the test compound. Experiments were repeated twice. Similar results were obtained in each. Data shown are mean±S.E.M. from one experiment.

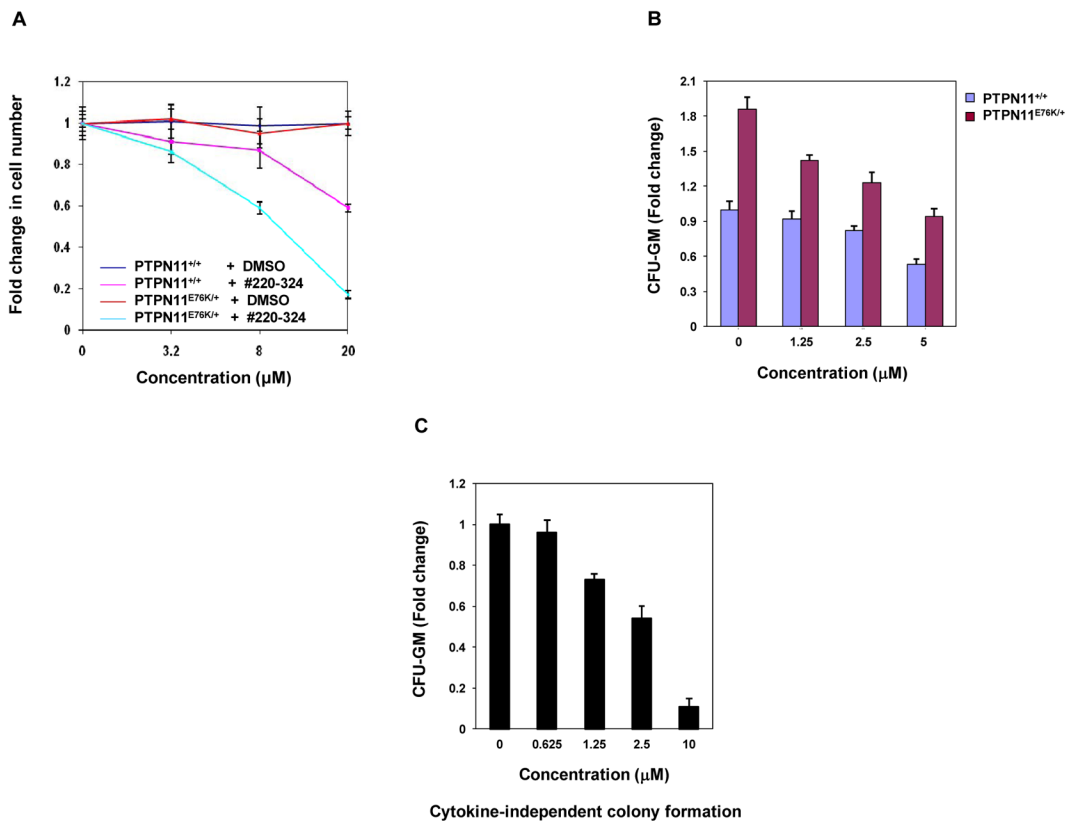


**Figure 2. #220–324 inhibited SHP2-mediated cell signaling and cellular proliferation** (A) Ba/F3 cells were cultured in IL-3 (1.0 ng/mL) containing medium supplemented with #220–324 at the indicated concentrations or control DMSO. Cell numbers were determined 72 hrs later using the One Solution Cell Proliferation Assay kit. Experiments were performed three times with similar results obtained. Data shown are mean $\pm$ S.E.M. from one experiment. (B) The cell cycle of Ba/F3 cells was analyzed 24 hrs after the treatment of #220–324 at the indicated concentrations. Percentages of the cells at G1, S, and G2/M phases were quantified by flow cytometric analyses. Experiments were repeated three times. Results shown are mean $\pm$ S.E.M. from one experiment. (C) Ba/F3 cells were deprived of IL-3 overnight. Cells were treated with #220–324 (30  $\mu\text{M}$ ) for 3 hrs and then stimulated with IL-3 (2.0 ng/mL) for the indicated times. Whole cell lysates were prepared. The levels of p-Erk, p-Akt, p-Stat5, p-Jak2 were determined by immunoblotting analysis. Blots were striped and reprobbed with anti-Erk, anti-Akt, and anti-Jak2, and anti-Stat5 antibodies to check for protein loading. Experiments were repeated three times. Representative results from one experiment are shown.



**Figure 3. #220–324 had minimal effects in *PTPN11* knockout cells**

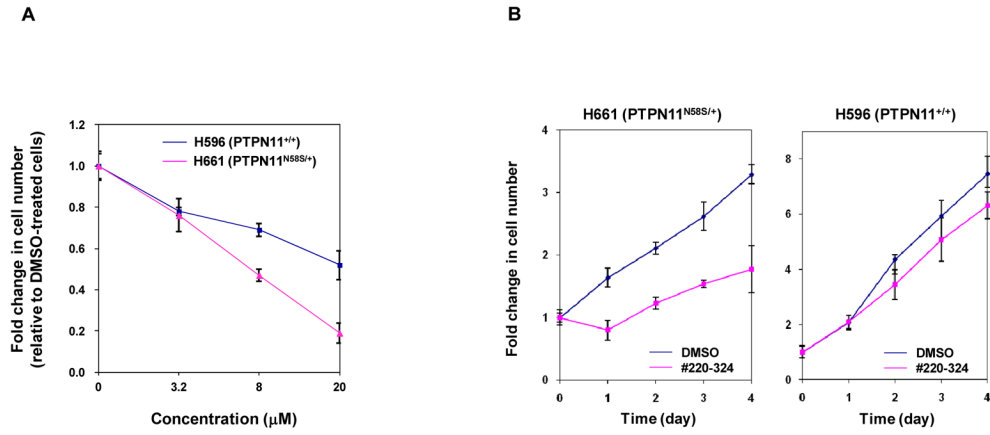
*PTPN11* knockout (*PTPN11<sup>F1/F1</sup>/Ad-Cre+*) and WT (*PTPN11<sup>+/+</sup>/Ad-Cre+*) MEFs were cultured in DMEM with 10% FBS and #220–324 at the indicated concentrations for 72 hrs (A) or at 20 μM for the indicated periods of time (B). DMSO-treated cells were included as negative controls. Cell numbers were determined using the One Solution Cell Proliferation Assay kit. Experiments were performed twice with similar results. Data shown are mean ±S.E.M. from one experiment.



**Figure 4. Mutant MEFs and myeloid progenitors with the *PTPN11*<sup>E76K/+</sup> mutation were more sensitive to #220–324 than WT cells**

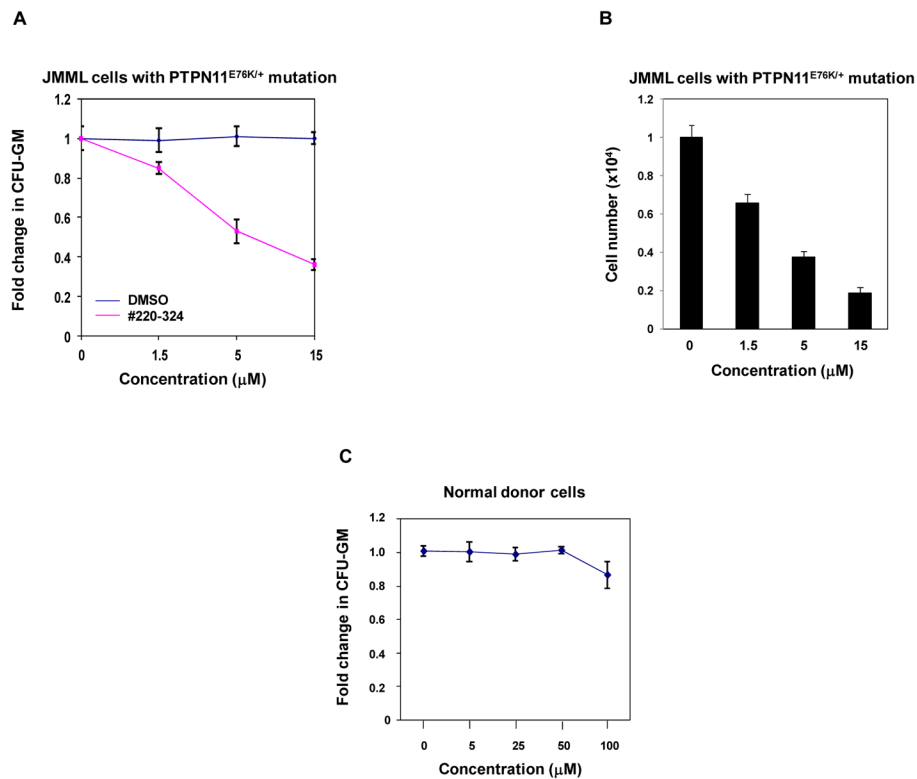
(A) WT and *PTPN11*<sup>E76K/+</sup> MEFs were cultured in 10% FBS-containing DMEM supplemented with #220–324 at the indicated concentrations. DMSO-treated cells were included as negative controls. Cell numbers were determined using the One Solution Cell Proliferation Assay kit 72 hrs later. Experiments were repeated three times. Similar results were obtained in each. Data shown are mean±S.E.M. from one experiment. (B) Bone marrow cells ( $2 \times 10^4$  cells/mL) harvested from *PTPN11*<sup>E76K/+</sup>/*Mx1-Cre*<sup>+</sup> and *PTPN11*<sup>+/+</sup>/*Mx1-Cre*<sup>+</sup> mice were plated in methylcellulose medium containing GM-CSF (1.0 ng/mL) and #220–324 at the indicated concentrations or control DMSO. Colonies were enumerated 7 days later and normalized against the number of colonies derived from WT control cells without #220–324 treatment. Representative results from two independent experiments are shown. Data are presented as mean±S.E.M. (C) Bone marrow cells ( $5 \times 10^4$  cells/mL) harvested from *PTPN11*<sup>E76K/+</sup>/*Mx1-Cre*<sup>+</sup> mice were plated in methylcellulose medium (without GM-CSF) supplemented with #220–324 at the indicated concentrations or control DMSO. Colonies derived in the cytokine-free medium were enumerated 7 days later and normalized against the number of colonies derived from the cells without #220–324 treatment. Representative results from two independent experiments are shown. Data are presented as mean±S.E.M.





**Figure 5. Lung cancer cells with the *PTPN11*<sup>N58S/+</sup> mutation were more sensitive to #220–324 than those without the *PTPN11*<sup>N58S/+</sup> mutation**

Human lung cancer cell lines H596 and H661 were cultured in DMEM supplemented with 10% FBS and #220–324 at the indicated concentrations for 72 hrs (A) or the cells were treated with #220–324 at 20 μM for the indicate times (B). DMSO-treated cells were included as negative controls. Cell numbers were determined using the One Solution Cell Proliferation Assay kit. Experiments were repeated three times. Similar results were obtained in each. Data are presented as mean±S.E.M. from one representative experiment.



**Figure 6. #220–324 abrogated the growth of JMML myeloid progenitors with *PTPN11*<sup>E76K/+</sup> mutation**

(A) Splenocytes ( $2 \times 10^4$  cells/mL) from a JMML patient with the *PTPN11*<sup>E76K/+</sup> mutation were plated in methylcellulose medium containing GM-CSF (1.0 ng/mL) and #220–324 at the indicated concentrations or control DMSO. Colonies were enumerated 14 days later and normalized against the number of colonies derived from the cells without #220–324 treatment. Three patient samples were tested in three independent experiments. Similar results were obtained in each. Data are presented as mean  $\pm$  S.E.M. from one patient sample. (B) Splenocytes from a JMML patient with the *PTPN11*<sup>E76K/+</sup> mutation were cultured in RPMI 1640 medium containing GM-CSF (1.0 ng/mL) and #220–324 at the indicated concentrations or control DMSO. Total cell numbers were determined and normalized against the number of the cells in the untreated group. Three patient samples were tested in three independent experiments. Similar results were obtained in each. Data are presented as mean  $\pm$  S.E.M. from one patient sample. (C) Apheresis peripheral blood cells ( $2 \times 10^4$  cells/mL) from normal donors were plated in methylcellulose medium containing GM-CSF (1.0 ng/mL) and #220–324 at the indicated concentrations. Colonies were enumerated 14 days later and normalized against the number of colonies derived from the cells treated with control DMSO. Three samples were tested in three independent experiments. Similar results were obtained in each. Data are presented as mean  $\pm$  S.E.M. from one sample.