Short Report

Discovery and biological evaluation of a potent small molecule CRM1 inhibitor for its selective ablation of extranodal NK/T cell lymphoma

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Abstract

Background

The overactivation of NF-κB signaling is a key hallmark for the pathogenesis of extranodal natural killer/T cell lymphoma (ENKTL), a very aggressive subtype of non-Hodgkin’s lymphoma yet with rather limited control strategies. Previously, we found that the dysregulated exportin-1 (also known as CRM1) is mainly responsible
for tumor cells to evade apoptosis and promote tumor-associated pathways such as NF-κB signaling.

Methods

Herein we reported the discovery and biological evaluation of a potent small molecule CRM1 inhibitor, LFS-1107. We validated that CRM1 is a major cellular target of LFS-1107 by biolayer interferometry assay (BLI) and the knockdown of CRM1 conferred tumor cells with resistance to LFS-1107.

Results

We found that LFS-1107 can strongly suppresses the growth of ENKTL cells at low-range nanomolar concentration yet with minimal effects on human platelets and healthy peripheral blood mononuclear cells. Treatment of ENKTL cells with LFS-1107 resulted in the nuclear retention of IkBα and consequent strong suppression of NF-κB transcriptional activities, NF-κB target genes downregulation and attenuated tumor cell growth and proliferation. Furthermore, LFS-1107 exhibited potent activities when administered to immunodeficient mice engrafted with human ENKTL cells.

Conclusions

Therefore, LFS-1107 holds great promise for the treatment of ENKTL and may warrant translation for use in clinical trials.

Funding
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**Key words:** CRM1; ENKTL; sulforaphene; NF-κB; drug discovery

**Introduction**

ENKTL represents as a rare yet highly aggressive subtype of non-Hodgkin’s lymphoma (NHL)\(^1\) with rather poor prognosis and short median survival time\(^2\). Previously, we and colleagues uncovered that the overactivation of NF-κB signaling and its downstream cytokines is a key hallmark for the pathogenesis of ENKTL\(^3\). Nevertheless, to date, targeted therapy towards ENKTL is still lacking and new therapeutic strategies are urgently needed to combat this rare yet deadly tumor\(^4\).

Exportin-1, also known as CRM1, has been implicated in the aggressive behavior of various malignances by nuclear exporting critical tumor suppressor proteins and transcription factors\(^5\). Moreover, CRM1 has been found to be highly upregulated in a panel of tumor types\(^6\). Indeed, it is believed that tumor cells strive to evade powerful negative regulation of apoptosis and cell proliferation through CRM1-mediated nuclear export machinery. Previously, we reported that a natural and covalent CRM1 inhibitor sulforaphene and its synthetic analogues\(^7,8,9,10\), can selectively kill a panel of tumor cells. Here, we presented the development and evaluation of a novel
sulforaphene analogue as a potent CRM1 inhibitor with superior antitumor activities towards ENKTL while sparing human platelets.

Results & Discussion

In this work, we want to find aromatic fragments that can be installed with sulforaphene parent structure as CRM1 inhibitors with strong potency. Previously, we developed a CRM1 inhibitor LFS-829 via traditional virtual screening and structure-based design approach. Herein, we strive to design a CRM1 inhibitor with strong potency against ENKTL cells through the recently developed AIDD (Artificial Intelligence Drug Discovery) approaches\textsuperscript{11}. Briefly, we adopted the deep reinforcement learning molecular de novo design model developed by Olivecrona et al.\textsuperscript{12} to facilitate the discovery of novel aromatic fragments. This model prioritizes those structures with modest similarity to the molecules in the positive dataset of known CRM1 inhibitors rather than very close analogues. We obtained an initial output of 3,000 moiety structures and the top 50 candidate moieties generated from this step were used as seed structures to search for commercial-available aromatic fragments from Sigma-Aldrich compound library (similarity ~ 97%). Next, we selected and purchased 10 commercial-accessible aromatic fragments (Figure 1A) which were experimentally tested via biolayer interferometry assay (BLI). Among the ten experimentally tested fragments, tetrazole moieties (S5 and S8 fragment) obtained the second highest binding constant with CRM1 via BLI assay experiments (Figure 1C and Figure 1 - Figure supplement 1). Yet, given that tetrazole fragments are
more synthetic accessible as compared to the fragment with the highest binding constant (S4 moiety in Figure 1A, benzo[d]oxazol-2-yl)-N, N-dimethylaniline), the tetrazole moiety was chosen to install with sulforaphene parent structure.

Subsequently, we prepared a synthetic analogue of sulforaphene with the tetrazole aromatic moiety, named as LFS-1107 (Figure 1B), which was further evaluated through cell lines and *in vivo* animal models.

First, we sought to evaluate the binding specificity of LFS-1107 with CRM1 as compared to other possible protein targets with reactive cysteine in the binding pocket through the Octet® K2 biolayer interferometry assay (BLI). Here, we chose IkBα and Keap1 as two control probes to compare because both of them are reactive for covalent compounds and thereby frequently used for testing selectivity. Our BLI results revealed that LFS-1107 binds strongly with CRM1 ($K_d \sim 1.25 \times 10^{-11}$ M, Figure 1D) as compared to KPT-330 ($K_d \sim 5.29 \times 10^{-9}$ M), a known CRM1 inhibitor approved by FDA. In contrast, LFS-1107 doesn’t bind with either two control probes IkBα or Keap1 in BLI assay experiments (Figure 1 - Figure supplement 2). Furthermore, the results revealed that LFS-1107 is a reversible CRM1 inhibitor with clear dissociation process which may implicate a low toxicity profile. This is consistent with the results of our previous studies that sulforaphene synthetic analogues are reversible CRM1 inhibitors. Moreover, we used GSH/GSSG detection assay kit to show that the GSH:GSSG ratio keeps constant upon the treatment of LFS-1107 and therefore we ruled out the possibility that LFS-1107 is a general redox modulator (Figure 2B).
Collectively, our results revealed that CRM1 is a major cellular target of LFS-1107 and responsible for its cellular activities.

Next, we assessed LFS-1107 for its activity and specificity in human ENKTL cell lines. Our data show that LFS-1107 achieves IC\textsubscript{50} value of 26 nM in SNK6 cell line and 36 nM in HANK-1 cell line (Figure 2A). To further study the selectivity of LFS-1107 towards ENKTL cells, we evaluated the toxicity of LFS-1107 in normal PBMCs isolated from peripheral blood of healthy donors (Figure 2C). Interestingly, LFS-1107 barely showed any toxicity at concentration of 4μM and the toxicity towards normal human PBMC is minimal even at a high dose of 9μM. This implicates that LFS-1107 can selectively eliminate ENKTL cells while sparing normal human PBMC with good safety profile. Moreover, drug-induced toxicity towards platelets (also called as thrombocytopenia) is a common side effect in clinical trials for cancer therapeutics. We assessed the toxicity effects of LFS-1107 towards platelets and we found that LFS-1107 barely exhibits any effects on human platelets even at very high concentrations of 500μM (Figure 2D).

Subsequently, we want to investigate whether LFS-1107 could inhibit CRM1-mediated nuclear export of IκB\textgreek{a} and reduce constitutive NF-κB activity in ENKTL cells. First, we demonstrated that LFS-1107 could suppress the expression of CRM1 in a dose-dependent manner (Figure 2E). We revealed that the cellular activities by LFS-1107 was significantly abrogated when CRM1 was knocked down by siCRM1, implicating that CRM1 is a main target responsible for the cellular
activities of LFS-1107 (Figure 2F). Our immunofluorescent results by confocal microscopy revealed that LFS-1107 can lead to nuclear accumulation of IκBα after 3h treatment in a dose dependent manner (Figure 2G-H). Next, we employed Western blot to further confirm nuclear localization of IκBα upon the treatment of LFS-1107 (Figure 2J). The results ascertained that IκBα was trapped in the nucleus as the protein level of nuclear IκBα was significantly upregulated after LFS-1107 treatment (Figure 2J). Moreover, it is well known that the constitutive activity of NF-κB/p65 was accompanied by the increased production of a few proinflammatory cytokines. Consequently, we found the expression of proinflammatory and proliferative proteins p65, COX-2, c-Myc and Survivin were downregulated in a dose dependent manner after treatment with LFS-1107 (Figure 2I). Furthermore, when SNK6 cells were treated with LFS-1107, we observed a significant reduction of the proinflammatory cytokines including TNF-α, IFN-γ, NF-κB/p65, IL-1α, IL-1β, IL-6, IL-8, and MCP-1 as measured by ELISA assay (Figure 2K). Moreover, proteomics analysis suggests that Biological Process (regulation of cellular component organization) and Molecular Function (protein binding) related to CRM1 were modulated upon the treatment of LFS-1107 (Figure 2- Supplement 1). These results suggested that the inhibition of CRM1 by LFS-1107 could lead to the downregulation of NF-κB transcriptional activity, proinflammatory cytokines and oncogenic signatures of ENKTL.

Lastly, we established a xenograft mouse model to test our findings by injecting SNK6 cells intraperitoneally into NCG mice. We found that LFS-1107 treatment
(10mg/kg/week) was able to extend mouse survival (Figure 3A-B) and eliminate tumor cells considerably as evidenced by flow cytometric analysis (Figure 3C-D-E), demonstrating the efficacy of LFS-1107 in controlling ENKTL. Moreover, remarkably, mice injected with LFS-1107 reduced splenomegaly and restored spleen weight and spleen volume as compared with the normal group (Figure 3F-G) with a better overall survival rate. Noteworthy, splenomegaly is a conventional symptom for extranodal natural NK/T cell lymphoma patients\textsuperscript{13}. Hence, our results implicate that LFS-1107 can ameliorate the symptoms of ENKTL and might be a potentially promising compound for ENKTL treatment.

**Conclusions**

In summary, we report the discovery and biological evaluation of a synthetic sulforaphene analogue as potent CRM1 inhibitor towards the treatment of ENKTL. We demonstrated that treatment of LFS-1107 holds great promise as an effective remedy for ENKTL, acting through the nuclear retention of \( \text{IkB} \alpha \) and subsequent attenuation of NF-\( \kappa\)B signaling. We want to remind the reader that we don’t rule out the possibility that LFS-1107 may kill ENKTL cells via other mechanisms due to the broad-spectrum cargo proteins of CRM1. Yet, we want to argue that our study has captured the most prominent mechanism of actions for LFS-1107 towards ENKTL. Our study may represent a novel route for eliminating ENKTL tumor cells with a distinct mode of actions.
Acknowledgments

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Conflict of interest statement

None declared.

Materials and methods

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The purity of the compound

Compound LFS-1107 was ascertained by $^1$H NMR, LC-MS and HPLC analysis (≥95% purity). The organic synthesis was conducted following standard procedures. All the chemicals and solvents were purchased from Sigma.

Organic synthesis

The NMR spectra of small molecule compounds were recorded on Bruker Avance 400 MHz for $^1$H NMR and 100 MHz for $^{13}$C NMR. The LCMS were taken on a quadrupole Mass Spectrometer on Shimadzu LCMS 2010 (Column: sepax ODS 50 × 2.0 mm, 5µm) or Agilent 1200 HPLC, 1956 MSD (Column: Shim-pack XR-ODS 30×3.0, 2.2µm) operating in ES (+) ionization mode. Chromatographic purifications were by flash chromatography using 100~200 mesh silica gel. Anhydrous solvents were pre-treated with 3A MS column before used. All commercially available reagents were used as received unless otherwise stated.

Cell lines, cell cultures and antibodies
Human lymphoma cell lines SNK6 and HANK-1 were purchased from ATCC. SNK6 and HANK-1 cell lines were maintained in RPMI 1640 medium supplemented with 10% FBS and 293T cell line in DMEM medium with 10% FBS. RPMI 1640 medium, DMEM medium and FBS (fetal bovine serum) were purchased from Hyclone (Thermo Scientific). Antibodies against Survivin (ab76424) was obtained from Abcam. Other antibodies were purchased from Cell Signaling Technology: anti-β-actin (#3700), anti-Cox2 (#4842), anti-p65 (#8424), anti-C-Myc (#13987) and anti-IκBα (#4814). Alexa 488-conjugated goat anti-rabbit antibody and Alexa 594-conjugated goat anti-mouse antibody were obtained from Invitrogen Life Technology (Invitrogen, CA, USA). Human lymphoma cell lines SNK6, HANK-1, 293T and Hela were purchased from the American Type Culture Collection (ATCC). The identity of these cell lines was confirmed by STR and the cells were tested negative for mycoplasma contamination throughout the experimental period.

**Deep reinforcement learning model**

In the present study, we adopted the deep reinforcement learning based molecular *de novo* design method developed by Olivecrona etc. from AstraZeneca. In brief, the deep reinforcement learning model consists of two main modules, the Design module (D) and the Evaluate module (E). The Design module (D) is used to produce novel chemical structures whereas the Evaluate module (E) is used to assess the feasibility and properties of the novel structures by assigning a numerical award or penalty to each new structure. The model employs the simple representation of chemical


structures by the simplified molecular-input line-entry system (SMILES) strings in both the Design module (D) and the Evaluate module (E). The Design module was first pre-trained with about ~2 million structures from the PubChem database to learn basic rules of organic chemistry that define SMILES strings within the context of Recurrent neural network (RNN). To check the validity of the approach, we used the module to generate about 0.2 million compounds which were assessed by the structure checker ChemAxon and 97% of the generated structures were chemical sensible structures. Finally, the deep reinforcement learning based molecular design model was able to design ~3,000 chemical structures from which we chose the top 50 candidate moieties for further consideration. In this work, we want to find novel fragments that can be installed with sulforaphene parent structure as CRM1 inhibitors. Indeed, our model prioritizes those structures with modest similarity to the molecules in the positive dataset rather than very close analogues. We used the values of molecule descriptors and root mean square deviation (RMSD) calculated as the reward function in the reinforcement learning stage in the model. We obtained an initial output of 3,000 structures, which were then automatically filtered to remove molecules bearing structure alerts. The model prioritizes the synthetic feasibility of a small-molecule compound, and the distinctiveness of the compound from other molecules in the literatures and in the patent space. The top 50 candidate moieties generated from the previous step were used as seed structures to search for commercial-available aromatic fragments (similarity ~ 97%). Lastly, we purchased 10
commercial-available aromatic fragments which were experimentally tested via biolayer interferometry assay (BLI).

**Bio-layer interferometry (BLI) assay**

The Octet® K2 system (Molecular Device, ForteBIO, USA) is suitable for the characterization of protein-protein or protein-ligand binding kinetics and binding affinity. The assay consists of the following steps: 1. Protein and BLI sensor preparation. The recombinant protein CRM1 (50 μg/mL) was biotinylated in the presence of biotin at room temperature (RT) for 1 h. Then, the excess biotin was removed through spin desalting columns. The recombinant CRM1 protein was then immobilized on super streptavidin sensors (ForteBIO, USA). The sensors were then blocked, washed, and moved into wells containing various concentrations of the test compounds in kinetic buffer (ForteBIO); 2. BLI experimental process. Automated detection was performed using an Octet® K2 (Molecular Devices ForteBIO, USA). Buffer was added to a 96-well plate, and the plate was transferred to the K2 instrument for analysis by running serially at 25°C with a shaking speed of 1000 rpm. Baseline readings were obtained in buffer (120 s), associations in wells containing compound (180 s), and dissociation in buffer (180 s). The signals from the following buffer were detected over time. First, each of these concentrations was applied to four sets of experiments: (a) CRM1-immobilized sensor in drug-containing kinetics buffer; (b) Blanked-sensor in drug-containing kinetics buffer; (c) CRM1-immobilized sensor in PBS-kinetics buffer; and (d) Blanked-sensor in PBS-kinetics buffer. The final value
was calculated using the equation: (a-c)-(b-d). This method removes interference from the sensor and drug-containing buffers. The binding signals were identified and the results were analyzed using the Octet®HT V10.0 software.

**Cell viability assay**

Cell viability assays were performed as previously described. Human lymphoma cell lines SNK6 and HANK-1 were seeded into 96-well plates and treated with LFS-1107 in concentrations of 0-800 nM for 72 h. Cell viability was evaluated using the WST-8-based Cell Counting kit-8 (Beyotime), which was added to the wells and incubated for 3 h. The absorbance of wells at 450 nm (reference wavelength 610 nm) was measured with a microplate reader (Infinite F50, Tecan).

**Cytotoxicity assay**

The Research Ethics Committee (REC) of Dalian University of Technology approved conduct of *ex vivo* assays with donated human cells (approval number: 2018-023). Normal human peripheral blood mononuclear cells (PBMCs) were obtained from blood samples collected from healthy volunteers. Approval was obtained from The Second Affiliated Hospital of Dalian Medical University institutional review board for these studies. Peripheral blood mononuclear cells were isolated by density gradient centrifugation over Histopaque-1077 (Sigma Diagnostics, St. Louis, MO, USA) at 400g for 30 min. Isolated mononuclear cells were washed and assayed for total number and viability using Trypan blue exclusion. PBMCs were suspended at
8×10^5/mL and incubated in RPMI 1640 medium containing 10% FBS in 24-well plates. Platelets suspended in plasma were collected by apheresis from volunteer donors after obtaining written consent (The Second Affiliated Hospital of Dalian Medical University). After dilution 1:10 in Tyrode's buffer, platelets were incubated for 24h at 37°C with graduated concentrations of test compounds (LFS-1107 or KPT-330) dissolved in DMSO. The EC50 values was calculated using GraphPad Prism software.

**Western blot**

The cytoplasmic and nucleic protein were extracted separately using the protein extraction kit (Boster) according to the manufacture’s introduction. Protein (30-45μg) was fractionated on a 10%-15% acrylamide denaturing gel and transferred onto a PVDF membrane. The membrane was blocked with 5% nonfat dry milk in TBST for 1 h at room temperature and washed in TBST three times, five minutes each time. The membrane was then incubated with primary antibodies at 1:500 to 1:1000 dilutions overnight at 4 °C. After washing with TBST for 15 min, the membrane was incubated with horseradish peroxidase (HRP) -conjugated secondary antibody at a 1:5000 dilutions for 1 h at room temperature. After further washing in TBST, the proteins were detected by enhanced chemiluminescence on X-ray film with ECL Western blotting detection kit (Thermo Fisher Scientific).

**Nuclear export assays and confocal fluorescence microscopy**
293T cells were seeded onto glass bottom cell culture dish at a density of 3,000-5,000 cells in 1000μL complete media. After incubation at 37 °C, 5% CO₂ for 24 h, cells were treated with CRM1 inhibitors for 3h. Small molecule compounds (KPT-330 and LFS-1107) were serially diluted 1:2 starting from 1μM in RPMI 1640 medium supplemented with 10% FBS. Following the indicated treatments, cells were fixed for 20 minutes with 4% paraformaldehyde in PBS. Next, cell membranes were permeabilized by 0.3% Triton X-100 in PBS for 20 minutes. After blocking with 5% bovine serum albumin (BSA) in PBS for 1 h at 37°C, cells were treated with IkBα antibodies in blocking buffer for 24h at 4°C. Anti-mouse Alexa 594 were used as secondary antibodies. Cell nuclei were stained with DAPI for 20 minutes. After washing, photomicrographic images were recorded with confocal laser scanning microscope Fluoview FV10i. For cell counts, at least 200 cells exhibiting nuclear, nuclear and cytoplasmic, or cytoplasmic staining were counted from three separate images. Percentages of N(Nuclear), N/C(Nuclear/Cytoplasmic) and C(Cytoplasmic) cells were calculated and standard deviations were determined.

**ELISA Assay**

Cell culture media were collected and centrifuged at for 20 min at 1000g at 4°C to remove cell debris. The concentrations of serum cytokines were assayed by the ELISA kit (Elabscience Biotechnology Co., Ltd) according to the manufacturer’s instructions. Three experimental replicates were performed for each sample.

**RNA Interference experiment**
The siRNA targets were as follows: XPO1-siRNA-1 (5'-CCAGCAAAGAAUGGCUCAATT-3'),
XPO1-siRNA-2(5'-GGAAGAUUCUCCAAGGAATT-3'),
XPO1-siRNA-3(5'-CCAGGAGACAGCUAUUUTT-3'), and the control target was 5'-UUCUCCGAACGUGUCACGUTT-3', all of which were obtained from Future Biotherapeutics. 293T cells were seeded in 96-well plates at a density of 5,000 cells per well in DMEM medium with 10% FBS the day before transfection. 293T cells were transfected with control siRNA or XPO1-siRNA according to the manufacturer’s instructions. The transfected cells were then collected for experiments 48h after transfection.

GSH/GSSG ratio detection

SNK6 cells were seeded onto four plates at a density of 100,000 cells/well in 3 mL complete media. After incubation at 37 °C, 5% CO2 for 24 h, cells were treated with LFS-1107 for 48h. Compound LFS-1107 was serially diluted 1:2 starting from 10mM in RPMI 1640 medium supplemented with 10% FBS. Following the indicated treatments, cells were tested by GSH and GSSG Assay Kit (Beyotime), which can detect the content of GSH and GSSG in the sample separately. We then calculated the GSH:GSSG ratio of each sample. The results were displayed by GraphPad Prism.

In vivo efficacy studies

NOD/SCID mice were purchased from the Nanjing Biomedical Research Institute of
Nanjing University (NBRI). The mice were randomly divided into three groups (n = 6–13 per group, day 0). In brief, SNK6 cells were intravenous injected into SCID mice and the growth of ENKTL cells was monitored every week for two consecutive weeks until the intrasplenic accumulation and proliferation of ENKTL cells. Subsequently, LFS-1107 was intraperitoneal injected to this ENTKL disease model per week (dose: 10mg/kg). In some experiments, SNK6 cells were pretreated with LFS-1107 (1μmol/L) or DMSO for 48hrs. Cells were then harvested and resuspended and the cell number was counted using a MuseTM cell analyzer. Animal studies in the present work have been conducted in accordance with the ethical standards and the Declaration of Helsinki. The investigation has been approved by the animal care and use committee of our institution.

Flow cytometry analysis.

Monocytes collected from mouse tissue were stained with FITC anti-human CD45 antibody and PE anti-mouse CD45 antibody. Isotype control antibody was used as a negative control. The Cytomics FC500 flow cytometry (Beckman, USA) was used for analysis.

Statistical analysis

All continuous variables were compared using one-way ANOVA, followed by Dunnett’s test or Tukey’s test for multiple comparisons whereas error bars in all
figures represent the SEM. * P < 0.05, **P < 0.01, ***P < 0.001. Survival curves were compared using the log-rank test with GraphPad software.

**Ethnics statement**

The protocol for the animal experiments and *ex vivo* assays with donated human cells was approved by the Research Ethics Committee (REC) of Dalian University of Technology (approval number: 2018-023). All the animal experiments were strictly followed ethnical rules under the guidance of Declaration of Helsinki.
Figure 1. The discovery of sulforaphene synthetic analogue LFS-1107. A. The identification of ten commercial-accessible aromatic fragments aided by deep reinforcement learning model; B. Synthesis of LFS-1107 via the installation of aromatic tetrazole moiety selected from the previous step to the sulforaphene parent structure; C. Assessment of protein-ligand binding kinetics and binding affinity of tetrazole aromatic fragments via Bio-layer interferometry (BLI) assay; D. Binding affinity of LFS-1107 and KPT-330 determined via BLI assay: LFS-1107, $K_d \sim 1.25 \times 10^{-11}$ M; KPT-330: $K_d \sim 5.29 \times 10^{-9}$ M.
Figure 1-source data 1 The chemical structure of ten commercial-accessible aromatic fragments.

Figure 1-source data 2 The synthesis of compound LFS-1107.

Figure 1-source data 3 The data of affinities and binding kinetics of CRM1 to S5 and S8.

Figure 1-source data 4 The data of affinities and binding kinetics of CRM1 to LFS-1107 and KPT-330.
Figure 1 - Figure supplement 1. Binding affinities and binding kinetics of ten commercial-accessible fragments with CRM1 were determined using Bio-layer interferometry (BLI) assay.

Figure 1 - Figure supplement 1 - source data The data of affinities and binding kinetics of CRM1 to ten commercial-accessible aromatic fragments.

Figure 1 - Figure supplement 2. The BLI results of two control proteins Keap1 (A) and IκBα (B). No binding affinities were detected for the two control proteins during BLI assay with compound LFS-1107.

Figure 1 - Figure supplement 2 - source data The data of affinities and binding kinetics of Keap1 and IκBα to LFS-1107.
**Figure 1 - Figure supplement 3.** Organic synthesis scheme of compound LFS-1107.

**Figure 1 - Figure supplement 3 - source data** Organic synthesis scheme of compound LFS-1107.
**Figure 1 - Figure supplement 4.** Expression of CRM1 mRNA in different tumor types. It shows that CRM1 was significantly elevated in patients with Lymphoma.

**Figure 1 - Figure supplement 4 - source data** The expression of CRM1 mRNA in different tumor types.

**Figure 2.** LFS-1107 strongly suppresses the growth of ENKTL cells acting through the nuclear retention of IkBα and subsequent attenuation of NF-κB signaling. A. Suppression of different human NK/T cell lymphoma cells; B. GSH/GSSG ratio detection upon the treatment of LFS-1107; C. The effect of...
LFS-1107 on normal PBMC cell lines; **D.** The viability of platelets treated with LFS-1107; **E.** Western blot result of the CRM1 with β-actin as loading control; **F.** The cellular activities of LFS-1107 on siCRM1-293T and the wild-type 293T cell line with different concentrations of treatment for 48h; **G.** Nuclear retention of IκBα by confocal microscopy; **H.** Quantification of the nuclear IKB-α ratio by fluorescence intensity per cell; **I.** Representative western blots of p65, Cox-2, c-Myc and Survivin; **J.** Western blots showing the protein level of IκBα in nucleus and cytoplasm; **K.** ELISA detection of cytokine production after treated with LFS-1107.

**Figure 2-source data 1** Inhibition of the cell growth of SNK6 and Hank-1 cells by LFS-1107.

**Figure 2-source data 2** GSH/GSSG ratio detection upon the treatment of LFS-1107.

**Figure 2-source data 3** Suppression of the cell growth of PBMC cells by LFS-1107.

**Figure 2-source data 4** Suppression of the cell growth of platelets by LFS-1107.

**Figure 2-source data 5** Immunoblot of CRM1 expression after LFS-1107 treatment.

**Figure 2-source data 6** The cellular activities of LFS-1107 on siCRM1-293T and the wild-type 293T cell line.

**Figure 2-source data 7** Nuclear accumulation of IκBα induced by treatment with LFS-1107 for 3 hours.

**Figure 2-source data 8** Quantification of the nuclear IκBα ratio by fluorescence intensity per cell.

**Figure 2-source data 9** Immunoblot of expression p65, Cox-2, c-Myc and Survivin after LFS-1107 treatment.

**Figure 2-source data 10** Immunoblot of IκBα in nucleus and cytoplasm expression after LFS-1107.

**Figure 2-source data 11** ELISA detection of TNF-α, IFN-1γ, p65, IL-1α, IL-1β, IL-6, IL-8 and MCP-1 after treated with LFS-1107.
Proteomics analysis of SNK6 cells (control vs. LFS-1107 treatment) indicates that CRM1 was downregulated upon LFS-1107 treatment (FC~1.3). Moreover, the Gene Ontology (GO) analysis suggests that Biological Process (regulation of cellular component organization) and Molecular Function (protein binding) related to CRM1 were modulated upon the treatment of LFS-1107.
Figure 2 – Figure supplement 1 - source data. Proteomics analysis of SNK6 cells.

Figure 2 – Figure supplement 2 - source data. Cellular activities of KPT-330 towards two ENKTL cell lines.
**Figure 2 - Figure supplement 3 – source data.** Nuclear accumulation of IκBα induced by treatment with 500 nM LFS-1107 for 3 hours. The medium (containing LFS-1107) was removed and replaced with new medium in the Wash group. Fixed cells were stained for IκBα (orange) and DAPI (blue).

**Figure 3.** LFS-1107 can ameliorate the symptoms of ENKTL in xenograft mouse model. **A.** Scheme of the xenograft mouse model. SNK6 cells were injected i.v. in female NOD SCID mice, and mice were injected per week with LFS-1107; **B.** Survival rate of different animal groups; **C-E.** Flow cytometry of human ENKTL cell lines in mouse bone marrow with the use of FITC anti-human CD45 antibody (P1, red)
or no primary antibody control (black); **F-G.** The symptoms of splenomegaly in each group of euthanized mice. From left to right: normal group, control ENKTL xenograft model group(splenomegaly), LFS-1107 treatment group (10 mg/kg).

**Figure 3 - source data 1** The process for the in the xenograft mouse model study.

**Figure 3 - source data 2** Survival rate of normal group, control group and LFS-1107 treatment group.

**Figure 3 - source data 3** Flow cytometry of human ENKTL cell lines in mouse bone marrow.

**Figure 3 - source data 4** The symptoms of splenomegaly in normal group, control group and LFS-1107 treatment group mice.

**Figure 3 - source data 5** The spleen weight of normal group, control group and LFS-1107 treatment group mice.
References


