Abstract. Treatment of resistant or recurrent acute lymphoblastic leukemia (ALL) remains a challenge. It was previously demonstrated that the adhesion molecule integrin α4, referred to hereafter as α4, mediates the cell adhesion-mediated drug resistance (CAM-DR) of B-cell ALL by binding to vascular cell adhesion molecule-1 (VCAM-1) on bone marrow stroma. In addition, it was previously observed that the blockade of α4 with natalizumab or inhibition using the small molecule antagonist TBC3486 sensitized relapsed ALL cells to chemotherapy. However, α4-targeted therapy is not clinically available for the treatment of leukemia to date. In the present study, the use of a novel non-peptidic small molecule integrin α4 antagonist, AVA4746, as a potential new approach to combat drug-resistant B-ALL was explored. An in vitro co-culture model of primary B-ALL cells and an in vivo xenograft model of patient-derived B-ALL cells were utilized for evaluation of AVA4746. VLA-4 conformation activation, cell adhesion/de-adhesion, endothelial tube formation, in vivo leukemia cell mobilization and survival assays were performed. AVA4746 exhibited high affinity for binding to B-ALL cells, where it also efficiently blocked ligand-binding to VCAM-1. In addition, AVA4746 caused the functional de-adhesion of primary B-ALL cells from VCAM-1. Inhibition of α4 using AVA4746 also prevented angiogenesis in vitro and when applied in combination with chemotherapy consisting of Vincristine, Dexamethasone and L-asparaginase, it prolonged the survival of ~33% of the mice in an in vivo xenograft model of B-ALL. These data implicate the potential of targeting the α4-VCAM-1 interaction using AVA4746 for the treatment of drug-resistant B-lineage ALL.

Introduction

Although rapid progress has been made in the development of treatment strategies for acute lymphoblastic leukemia (ALL) over the past decades, prognosis for patients with relapsed or treatment-refractory ALL remains poor (1,2). The bone marrow microenvironment serves a critical role in protecting leukemic cells against chemotherapy (3-8), with cell adhesion-mediated drug resistance (CAM-DR) being one of the underlying mechanisms (9-12). Integrins belong to a family of glycoprotein cell surface receptors that are comprised of two subunits, α and β (13), which primarily mediates cell adhesion in the extracellular matrix (ECM). The integrin α4 chain, also known as CD49d, non-covalently associates with the β1 integrin chain, also known as CD29, to form very-late-antigen 4 (VLA-4) (14). VLA-4 in turn binds to counter receptors, including vascular cell adhesion molecule-1 (VCAM-1), fibronectin and osteopontin (14). During normal development and hematopoiesis, α4 is crucial for homeostasis, regeneration and the homing of hematopoietic stem and progenitor cells in the bone marrow (15,16). In particular, integrin α4 is expressed in B-ALL cells (17) and has been previously implicated in CAM-DR which serves a key role in tumor microenvironment (7), since the binding of α4 to its ligand VCAM-1

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mediates signaling to maintain the survival of leukemic cells in the presence of chemotherapy (8,18,19).

In previous study, its was shown that natalizumab (NZM), a humanized anti-integrin α4 monoclonal antibody, can cause the de-adhesion of B-ALL cells from VCAM-1, thus sensitizing ALL cells to chemotherapy (20). However, to date there have been no clinically approved agents that target α4 for the treatment of B-ALL. Previously, a non-peptidic small molecule α4 inhibitor, TBC3486, which is a small molecule mimetic of VCAM-1, was tested in mice with primary ALL, which prolonged the survival of the mice (21). However, TBC3486 remain unavailable in an oral formulation and no in-depth preclinical studies have been previously conducted.AVA4746 is an analog of TBC3486 that is also a small molecule mimetic of VCAM-1 (22,23). Notably, AVA4746 was found to be well tolerated as an oral formulation in previous human clinical trials conducted (22,23). AVA4746 is ~five times more potent in dislodging VLA-4 from VCAM-1 compared with TBC3486, according to in vitro cell assays (unpublished data from Aviara Pharmaceuticals, Inc.). In the present study, the in vitro competitive ligand-binding and anti-adhesive properties of AVA4746 were evaluated. In addition, its chemotherapy sensitization effects in vivo by reversing CAM-DR in B-ALL cells were also assessed.

Materials and methods

Patient samples and cell lines. As described previously (20,24-26) bone marrow cells were obtained from patients with B-ALL after informed consent was obtained under Institutional Review Board of Children's Hospital Los Angeles approved protocols. Primary B-ALL blasts were obtained from the bone marrow aspirates by separation using Ficoll (Cytiva) gradient centrifugation of 450 x g at 20°C for 30 min then pellets were harvested. Harvested mononuclear cells (0.01-1x10^6/200 µL PBS/mouse) were intravenously injected into NOD.Cg-Prkdc^scid Il2rg^tm1Wjl/ (Jackson Laboratories; total of 129 mice; age, 5-7 weeks; weight, 17-24 g), which were conditioned with a single sub-lethal temperature, and expanded. Mice had free access to food and water and were housed at 21-23°C -room temperature, and expanded. Mice had free access to food and water and were housed at 21-23°C room temperature, at 30-70% humidity and with a 6:00 am-7:00 pm light cycle. The mice were sacrificed using CO 2 at 30% air displacement rate. Engrafted human leukemia cells were harvested from the bone marrow and spleen 2-26 weeks after injection, before they were cultured at 37°C with 5% CO 2 further in the presence of murine stromal OP-9 cells (American Type Culture Collection, ATCC) in vitro. Primary B-ALL cells (LAX7, LAX7R, LAX53, TXL3 and ICN24 from the Lab of Dr. Yong-Mi Kim) were co-cultured with OP-9 stromal cells in MEM-α (Invitrogen; Thermo Fisher Scientific, Inc.) supplemented with 20% fetal bovine serum (FBS; Invitrogen; Thermo Fisher Scientific, Inc.), 100 U/ml penicillin and 100 µg/ml streptomycin. The B cell lineage-ALL (B-ALL) cell lines were all incubated at 37°C with 5% CO 2, TOM-1 (DSMZ) and BV173 (DSMZ) were cultured in 80% RPMI-1640 (Invitrogen; Thermo Fisher Scientific, Inc.) + 20% FBS. SupB15 (ATCC) was cultured in 80% IMDM (Invitrogen; Thermo Fisher Scientific, Inc.) + 20% FBS. REH (ATCC), RS4;11 (ATCC), Kasumi-2 (DSMZ), 697 (DSMZ), BEL1 (DSMZ), RCH (DSMZ) were in cultured in 90% RPMI 1640 + 10% FBS. Cytogenetic information and α-integrin expression percentages of all B-ALL cell lines used for the present study are shown in Table S1 (25,26). Human umbilical vein endothelial cells (HUVECs; Thermo Fisher Scientific, Inc.) were cultured in Medium 200 supplemented with 0.2% LSGS (Invitrogen; Thermo Fisher Scientific, Inc.) at 37°C with 5% CO 2.

Compounds. AVA4746 and TBC3486 were obtained from Aviara Pharmaceuticals, Inc. for research use. MG132 was obtained from Selleck Chemicals. Vincristine, Dexamethasone and L-asparaginase were obtained from Children Hospital of Los Angeles.

Flow cytometry. After harvesting, B-ALL cells or HUVECs (0.1-1x10^6) were centrifuged at 500 g for 5 min at room temperature and resuspended in 100 µl PBS containing anti-human CD19 (#302206, 1:2 dilution, Biolegend, Inc.), anti-human CD49d (#304304, 1:20 dilution, Biolegend, Inc.) and anti-human CD29 (#303008, 1:2 dilution, Biolegend, Inc.) antibodies, alongside the isotype control mouse IgG1 (Biolegend, Inc.). Following incubation at 4°C for 30 min, cells were washed with 1 ml PBS and resuspended in 100 µl DAPI/PBS (0.1µg/ml) for 15 min at 4°C for assessment using a flow cytometer (BD FACSCanto II, BD Biosciences). Flow cytometry data were analyzed from the viable cells in the DAPI-negative population, based on an isotype gating strategy using Flow Jo (version 10, BD Biosciences).

For apoptosis assay, B-ALL cells were treated for 48 h with either DMSO or AVA4746 25 µM, with or without VDL chemotherapy (vincristine 5 nM, dexamethasone 0.05 nM, L-asparaginase 2.5x10^3 IU/ml) in the presence or absence of OP9. Annexin V (#640947, 1:20; Biolegend, Inc.) and DAPI were used to stain the 1x10^6 cells for 15 min at room temperature (25°C), in the dark.

VLA-4 conformation activation assay. As previously described (27,28), LAX7R or TXL3 cells were suspended in the HEPES buffer (110 mM NaCl, 10 mM KCl, 10 mM glucose, 1 mM MgCl_2, 1.5 mM CaCl_2 and 30 mM HEPES, pH 7.4 containing 0.1% HSA Sigma-Aldrich; Merck Kga) at a density of 1x10^6 cells/ml. B-ALL cells were then incubated with either AVA4746 of 0.00, 0.01 µM, 0.1 µM, 1 µM, 10 µM, or 100 µM or human VCAM-1 (PeptoTech, Inc.) of 0 µM, 1.35 µM, 13.5 µM, 135 µM, or 1350 µM in the presence of 10% phycoerythrin (PE)-conjugated HUTS-21 antibodies (#556049, BD Biosciences) with or without 5mM Mn^2+ for 30 min at 37°C. Subsequently, 0.5 µl DAPI (200 µg/ml) was added to each reaction 5 min before the end of the incubation time. The PE-HUTS-21 mean fluorescence intensity (MFI) was analyzed by flow cytometry (BD FACSCanto II, Flow Jo version 10, BD Biosciences). Receptor occupancy (RO) assays are designed to quantify the binding of therapeutics to their targets on a cell's surface (29). The protocol used for calculating the normalized VLA-4 RO was as follows: The maximal MFI value from highest concentration of AVA4746 or VCAM-1 was normalized as 100%, whilst the minimum MFI value from lowest concentration of AVA4746 or
VCAM-I was normalized as 0%. EC50 of RO (half maximal occupancy of VLA-4 receptor concentration) was calculated using GraphPad Prism 5 software. To determine a level of non-specific binding, cells were stained in parallel with the 10% isotype control antibody (PE Mouse IgG2a, k Isotype Control, #555574, BD Biosciences).

Cell adhesion and de-adhesion assay. For the cell adhesion assay, non-tissue culture 96-well plates were coated with 10 µg/ml human VCAM-1 (PeproTech, Inc.) overnight at 4°C, washed and blocked with 2% BSA (Sigma-Aldrich; Merck KGaA) for 30 min at room temperature. LAX7R or TXL3 cells (1x10^5/well) were incubated with 0.00, 0.01 µM, 0.1 µM, 1 µM, 10.00 or 100 µM AVA4746 at room temperature for 1 h, added into the plates pre-coated with VCAM-1 and were allowed to adhere for 1 h at 37°C. Non-adherent cells were removed before being gently washed once by PBS, whilst adherent cells were counted using a hemocytometer following Trypan Blue (Invitrogen; Thermo Fisher Scientific, Inc.) exclusion.

Cell de-adhesion assays were performed as previously described (20,21). Similarly, B-ALL cells (1x10^5/well) were seeded into the plates pre-coated with 10 µg/ml human VCAM-1 or OP9 cells for 4 h at 37°C in 5% CO2. Next, either AVA4746 (25 µM) or DMSO was added as a vehicle control and the cells were incubated at 37°C with 5% CO2 overnight. The non-adherent cells in the supernatant were then removed, whereas alive adherent cells were detached by pipetting with PBS prior to counting by using Trypan Blue exclusion of dead cells with a hemocytometer. Finally, the percentage of adhesion was calculated by dividing the number of live adherent cells by the total number of cells.

Western blot analysis. B-ALL cells were treated with AVA4746 (0.1 µM, 5 µM, or 25 µM) for 24 h or 96 h for on-target effect experiments. Furthermore, following 2 h pre-treated with MG132 0 µM or 1 µM B-ALL cells were treated with AVA4746 either 0 µM or 25 µM for 96 h as separate experiments. B-ALL cells were harvested and lysed in M-PER™ Mammalian Protein Extraction Reagent (Invitrogen; Thermo Fisher Scientific, Inc.) containing a 1% protease inhibitor cocktail (VWR International, LLC). Protein concentration was measured using Bradford protein assay. Protein lysates (20 µg/lane) were separated in 4-12% Bis-Tris protein gels by SDS-PAGE and transferred onto PVDF membranes. Then the membranes were blocked by 5% dry fat milk for 1 h at room temperature. The following primary antibodies were incubated overnight at 4°C: Anti-integrin α4 (#4600, 1:1,000; Cell Signaling Technology, Inc.), anti-phosphorylated (p-)AKTSer473 (#9271, 1:1000; Cell Signaling Technology, Inc.), anti-AKT (#9272, 1:1,000; Cell Signaling Technology, Inc.), anti-phosphoantigen (#9525, 1:1000 dilution, EMD Millipore) and anti-β-actin (#sc-47778, 1:2000 dilution, Santa Cruz Biotechnology, Inc.). Secondary antibody solution Alk-Phos conjugated (anti-rabbit #WP20007 or anti-mouse #WP20006; Thermo Fisher Scientific, Inc.) was applied at room temperature for 1 h. To visualize bands, chemiluminescent substrate (#WP20002, Thermo Fisher Scientific, Inc.) was used. Western blots quantity was analyzed via Image J (version 1.53e, National Institutions of Health).

Starvation and activation assay for the detection of phosphorylated proteins. B-ALL cells were first serum-starved by being washed twice with Dulbecco’s PBS (DPBS) and cultured in MEM-α media at 37°C and 5% CO2 overnight. Following another wash with DPBS, B-ALL cells were treated with either the vehicle control 0.1% DMSO or AVA4746 (5 or 25 µM) for 30 min at room temperature. Subsequently, FBS was added to a final concentration of 20% to all cells except for those in the no-activation control groups. In either the human VCAM-1 or OP9 group, 1x10^6 cells were seeded into plates pre-coated with either human VCAM-1 (10 µg/ml) or OP9 (0.2x10^6 cells/well) on 12-well plate were pre-coated at 37°C for 1 day before experiment. Whole cell lysates were isolated after incubation for either 1 h or 24 h at 37°C for use in western blot analysis for either p-AKT or phosphotyrosine detection.

In vivo leukemia cell mobilization study. Mobilization of leukemia cells was tested according to a previously described protocol (20). Primary B-ALL LAX7R cells (0.05x10^6 in 200 µl PBS per mouse) were intravenously injected into total of 14 NSG mice (7 mice in PBS group while 7 mice in AVA4746 treatment group). In total, 80 µl peripheral blood was collected each mouse by retroorbital exsanguination following 2% isoflurane anesthesia once a week from weeks 2 to 4 after the injection of LAX7R cells. The total duration of the blood sampling protocol was typically 3-6 min. When 0.5 or ~2.5% hCD45^+ hCD19^+ LAX7R cells were detected by flow cytometry in the peripheral blood (PB) of NSG mice, the mice were treated with either PBS or AVA4746 (60 mg/kg) by intraperitoneal (i.p.) injection. Following 100 µg/ml purified human IgG (#I2511, Sigma-Aldrich, Inc.) as blocking reagent for elimination of non-specific binding, 0.1x10^6 cells were stained with APC-hCD45 (#304012, 1:20 dilution, BioLegend, Inc.), PE-hCD19 (#302254, 1:20 dilution, BioLegend, Inc.), FITC-mCD45 (#103108, 1:20 dilution, BioLegend, Inc.) and anti-β-actin (#sc-47778, 1:2000 dilution, Santa Cruz Biotechnology, Inc.) at either 8 or 36 h after treatment. Alive mononuclear cells were isolated from the PB, bone marrow (BM) and spleen (SPC) using the ACK lysing buffer (Invitrogen; Thermo Fisher Scientific, Inc.) at either 8 or 36 h after treatment. Alive mononuclear cell (MNC) numbers were counted with a hemocytometer using trypan blue exclusion, whilst the percentages of either mCD45^+ or hCD45^+hCD19^+ cells were determined by flow cytometry (BD FACSCanto II, Flow Jo version 10; BD Biosciences). Therefore, number of mouse cells or leukemia cells was calculated as MNC number x percentage of mCD45^+ or hCD45^+hCD19^+ for mouse and human leukemia cells, respectively. This animal study was performed in compliance with a research protocol approved by the Institutional Animal Care and Use Committee at the Saban Research Institute of Children's Hospital Los Angeles.

Endothelial tube formation assay. A Matrigel-coated 48-well plate (#354508, Biocoat™; BD Biosciences) was first pre-warmed at 37°C for 30 min. HUVECs (4-6x10^4/well) that were treated with AYA4746 (0, 5 or 25 µM) or TBC3486 (0, or 25 µM) at 37°C for 30 min were seeded into each well and incubated for 4-6 h at 37°C in 5% CO2. After incubation, images
of three representative fields of view per well were obtained using phase contrast light microscopy (x100 magnification), before being analyzed using the ‘angiogenesis analyzer’ tool (Gilles Carpentier) in ImageJ (30) (version 1.53e, National Institutes of Health). Tube formation was assessed by analyzing the number of nodes (pixels with ≥ 3 neighboring elements corresponding to a bifurcation), segments (elements delimited by two junctions), meshes (areas enclosed by segments or master segments and made by tube-like structures) and total area, before being quantified.

**In vivo survival study.** Primary B-ALL cells (0.05x10⁶) were intravenously injected into each NSG mouse. In total of 91 mice were divided into four treatment groups: PBS (5 mice in LAX7R exp#1, 4 mice in LAX7R exp#2, 7 mice in RS4;11, 5 mice in TXL3), AVA4746 group (5 mice in LAX7R exp#1, 5 mice in LAX7R exp#2, 7 mice in RS4;11, 6 mice in TXL3), VDL group (6 mice in LAX7R exp#1, 5 mice in LAX7R exp#2, 6 mice in RS4;11, 6 mice in TXL3), and VDL+AVA4746 group (6 mice in LAX7R exp#1, 5 mice in LAX7R exp#2, 8 mice in RS4;11, 5 mice in TXL3). AVA4746 (15 or 30 mg/kg dissolved in PBS) was then administered by oral gavage twice a day continuously for 14, 21 or 28 days, with treatment starting 3 days after B-ALL cell injection, vincristine (0.5 mg/kg) was administered (i.p.) once a week for 2 or 4 weeks, whilst 20% FBS. Therefore, VCAM-1 cell adhesion assays were used to determine the half maximal inhibitory concentration (IC₅₀) of AVA4746. The IC₅₀ of AVA4746 was 8.12 (7.07-9.32 µM) and 13.95 µM (11.15-17.45 µM) in LAX7R and TXL3 cells, respectively (Figs. 1C, D and S1D). There was little conformation changes corresponding to VLA-4 activation by human VCAM-1 without 5 mM Mn²⁺ (Figs. 1C, D and S1B). The MFI of PE-HUTS-21 shown in Table SII are from two independent experiments performed in triplicate. Taken together, AVA4746 exhibits nanomolar levels of affinity for the active conformation of VLA-4, which was ~3,500-fold higher compared with that of VCAM-1 in the presence of Mn²⁺.

Other soluble integrin ligands present in the serum may interfere with VLA-4 ligand binding (31), since the primary B-ALL cells were cultured in MEM-α in the presence of 20% FBS. Therefore, VCAM-1 cell adhesion assays were used to determine the half maximal inhibitory concentration (IC₅₀) of AVA4746. The IC₅₀ of AVA4746 was 8.12 (7.07-9.32 µM) and 13.95 µM (11.15-17.45 µM) in LAX7R and TXL3 cells, respectively, in one out of the two independent experimental repeats (Fig. 1E and F). The percentages of adhesion from the two independent experiments are shown in Table SIII. To achieve a saturating competitive effect, 5 and 25 µM AVA4746 were used for subsequent experiments.

**AVA4746 decreases cell surface and total intracellular expression of integrin α4.** Primary B-ALL cells derived from four human patients (LAX7R, LAX7, LAX53 and LAX56) and the B-ALL cell line REH were used to evaluate the on-target effect of AVA4746 on B-ALL in vitro. Cells were treated with either AVA4746 (1, 5 and 25 µM) or 0.1% DMSO as a control for either 24 or 96 h, before they were analyzed by flow cytometry to detect human integrin α4 expression. The MFI of integrin α4 on the B-ALL cells was decreased by AVA4746 in a dose-dependent manner, whilst the MFIs of integrins α5 and α6 were not affected (Figs. 2A-E and S2). AVA4746 also caused a reduction in the protein expression levels of α4 as determined by western blot analysis, after 24 and 96 h of treatment (Fig. 2F). Subsequently, the effects of proteasomal degradation on this decrease in integrin α4 protein expression were subsequently tested. B-ALL cells were treated for 96h with either AVA4746 (25 µM), MG132 (1 µM) or DMSO as a vehicle control, alone or in combination. α4 was partially restored by MG132 in all cases apart from LAX56 (Fig. S3.). These results suggest that this decrease in α4 expression induced by AVA4746 may also be independent of the ubiquitination-proteasome pathway. In addition, AVA4746 slightly downregulated the phosphorylation levels of AKT following stimulation with human VCAM-1 in serum-starved LAX7R cells, but no obvious changes were observed following AVA4746 treatment after stimulation with serum (Fig. S4A). Similar tendency was found in LAX56 (Fig. S4A). These results suggest that other cell signaling pathways may be involved (Fig. S4A), since AVA4746 may also affect phosphotyrosine cell signaling in LAX53 cells (Fig. S4B). Clear decreased band densities were

**Results**

AVA4746 binds VLA-4 on B-ALL cells with high affinity and efficiently blocks ligand-binding with VCAM-1. Since AVA4746 is a small molecule mimetic compound of VCAM-1 (23), conformation activation experiments were performed to determine the half maximal effective concentration (EC₅₀) of AVA4746. HUTS-21 is a specific monoclonal antibody that targets an epitope corresponding to the active conformation of integrin β1 (31). Anti-HUTS-21 antibody binding would occur when VLA-4 binds to certain ligands, such as VCAM-1 (31). The HUTS-21 antibody binding curve represents the VLA-4 RO, which is used as an indicator of binding affinity (31). In LAX7R cells, the EC₅₀ of AVA4746 was 38.52 nM (26.84-55.28 nM) and 1357.00 nM (1295.00-1422.00 nM) in the presence (Fig. SIC) or absence of 5 mM Mn²⁺ (Fig. S1A), respectively (Fig. 1A). In TXL3 cells, the EC₅₀ of AVA4746 was 32.68 nM (24.56-43.49 nM) and 973.00 nM (914.00-1036.00 nM) in the presence (Fig. SIC) or absence of 5 mM Mn²⁺ (Fig. S1A), respectively (Fig. 1B). By contrast, the EC₅₀ of human VCAM-1 was 102.20 µM (67.41-154.90 µM) and 110.60 µM (63.81-191.90 µM) in the presence of 5 mM Mn²⁺ in LAX7R and TXL3 cells, respectively (Figs. 1C, D and S1B). The MFI of PE-HUTS-21 shown in Table SII are from two independent experiments performed in triplicate. Taken together, AVA4746 exhibits nanomolar levels of affinity for the active conformation of VLA-4, which was ~3,500-fold higher compared with that of VCAM-1 in the presence of Mn²⁺.
found in both 5 µM and 25 µM A VA4746 compared with 0 µM in 1 h treatment group around 50‑60 kDa.

**AVA4746 detaches B-ALL cells from VCAM-1.** To investigate the competing ligand effect, the de-adhesion assays were performed as previously described (21). Five types of primary B-ALL cells, namely LAX7R, LAX53, LAX56, TXL3 and ICN24 cells (Fig. 3A–E), along with seven B-ALL cell lines, namely RS4;11, SupB15, Kasumi-2, 697, BEL-1, BV173 and RCH cells (Fig. 3F–L; Table SI), were plated onto either human VCAM-1 (10 µg/ml) or 2% bovine serum albumin (BSA)-coated plates, before being treated with either DMSO or 25 µM A VA4746 overnight. All 12 types of B-ALL cells showed significant de-adhesion from the VLA-4 ligand, human VCAM-1, after A VA4746 treatment compared with that in the DMSO control (Fig. 3). In addition, A VA4746 was found to induce the detachment of eight B-ALL cell lines from the OP9 stromal cell line (Fig. S5A‑H). Since the murine calvaria-derived stromal cell line OP9 has been found to facilitate survival and drug resistance in B-ALL cells (20), Annexin V assays were subsequently performed to determine the apoptotic effects of A VA4746 in combination with the chemotherapeutic regimen of 0.5 mg/kg vincristine, 10.5 mg/kg dexamethasone and 1500 IU/kg L-asparaginase (VDL). The small but significant induced more apoptosis were found in VDL+A VA4746 compared with VDL alone in TXL3 (without OP9, Fig. S6A), LAX56 (without OP9, Fig. S6B) and LAX7R (with OP9, Fig. S6C). However, neither A VA4746 alone nor A VA4746 in combination with VDL induced noTable apoptotic effects in B-ALL cells compared with that in the corresponding groups that were not treated with A VA4746, regardless of the presence of OP9 cells (Fig. S6A-C).

![Figure 1. AVA4746 has high affinity for VLA4 and efficiently competes with VCAM-1 on B-ALL cells.](image-url)
Figure 2. AVA4746 decreases cell surface and total integrin α4 expression. MFI representing the expression levels of integrin α4 on the (A) LAX7R, (B) LAX7, (C) LAX53, (D) LAX56 and (E) REH cell surfaces 24 and 96 h after treatment with 1, 5 and 25 µM AVA4746, with 0 representing the DMSO control. *P<0.05 vs. 0. Experiments were performed in triplicates. (F) B-ALL cell lines LAX7R, LAX7, LAX53, LAX56 and REH were treated for 24 and 96 h with 1, 5 and 25 µM AVA4746, with 0 representing the DMSO control. Following 24 and 96 h treatment with AVA4746, the integrin α4 protein levels at 140 kDa in LAX7R, LAX7, LAX53, LAX56 and REH cells were analyzed by western blotting, with β-actin being used as the loading control. MFI, mean fluorescence intensity; α4, α4 integrin.
AVA4746 moderately inhibits the migration of LAX7R cells from BM to the peripheral blood. To determine the role of integrin α4 in B-ALL cell metastasis, mobilization assays were performed. In brief, AVA4746 treatment was initiated when detectable B-ALL cells could be observed in PB. Subsequently, mice were sacrificed before the PB, BM and SPC were harvested for analysis. When ~2.5% LAX7R cells was detected in the PB (Fig. S7A), mice were treated with either AVA4746 (60 mg/kg; n=6), PBS as a control (n=6), VDL (n=6) or VDL + AVA4746 (n=6).

After 8 h of treatment, PB, BM and SPCs were harvested for a human leukemic population analysis using flow cytometry. The timeframe of the initial treatment was subsequently optimized, so that when the leukemia cell (hCD45+hCD19+) percentage in the PB was 0.5%, the duration of AVA4746 treatment was prolonged to 36 h (Fig. 4A). Consequently, AVA4746 significantly decreased the percentage of B-ALL cells (hCD45+hCD19+) in the PB compared with that in the PBS control (Figs. 4B and S8A) whilst significantly increasing that of B-ALL cells (hCD45+hCD19+) in the BM (Figs. 4C and S8B). No changes were found in the SPC samples (Figs. 4D and S8C).

AVA4746 inhibits in vitro angiogenesis. Results from the in vivo mobilization assays suggest that AVA4746 can cause the retention of B-ALL cells in the BM. Since integrin α4 and β1 were found to be expressed on endothelial cells (Fig. 5A), it was hypothesized that the underlying mechanism may be due to the AVA4746-mediated inhibition of α4 on endothelial cells. This may in turn reduce vessel formation and reduce the release of B-ALL cells into the peripheral circulation. To test this hypothesis, the potential effects of AVA4746 on HUVECs were evaluated in vitro. AVA4746 significantly attenuated the in vitro angiogenic potential of HUVECs. Representative images of HUVECs in Fig. 5B demonstrate that the number of nodes, segments, meshes and total meshes was all significantly decreased by both 5 and 25 µM AVA4746 (Fig. 5C).
Figure 4. AVA4746 decreases the number of human B-ALL cells in the mouse peripheral blood samples whilst increasing the number of ALL cells in the bone marrow. (A) Experimental design of the in vivo leukemic cell mobilization assay. When 0.5% of human CD45/CD19 double-positive LAX7R cells were detected by flow cytometry in the PB of the NSG mice, mice were treated with either PBS (n=7) or 60 mg/kg AVA4746 (n=7). Mononuclear cells were isolated from the PB, BM and SPC 36 h after treatment. Percentages of hCD45+hCD19+ cells in the (B) PB, (C) BM and (D) SPC are shown. *P<0.05. PB, peripheral blood; BM, bone marrow; SPC, spleen; NSG, NOD.Cg-PrkdcscidIl2rgtm1Wjl/SzJ; NS, no significant; B-ALL, B-lineage acute lymphoblastic leukemia.

Figure 5. AVA4746 attenuates tube formation by HUVECs. HUVECs treated with either 0, 5 or 25 µM AVA4746 for 30 min were seeded into Matrigel-coated plates for 4-6 h. (A) Representative flow cytometry dot plot of human integrin α4 and β1 expression in HUVECs before seeding into the Matrigel wells. (B) Representative images of the HUVECs are shown. (C) Tube formation was assessed by analyzing the number of nodes (pixels with ≥ three neighboring elements corresponding to a bifurcation), segments (elements delimited by two junctions), meshes (areas enclosed by segments or master segments and made by tube-like structures) and total area, before being subsequently quantified. Images were analyzed with using the ImageJ software and the ‘angiogenesis analyzer’ plugin. *P<0.05, **P<0.01 and ***P<0.001. HUVEC, Human umbilical vein endothelial cells; α4, α4 integrin; β1, β1 integrin; Nb, number.
Similar results were found using TBC3486, an analog of AVA4746 (Fig. S9). However, there were no statistical differences in angiogenesis inhibition using NZM (data not shown).

Combination of AVA4746 and chemotherapy treatment prolongs the survival of mice engrafted with primary relapsed B-ALL. To further determine the effects of AVA4746 on the survival of human B-ALL cell mouse xenograft models, LAX7R cells (0.05 x 10⁶ cells per mouse) were injected into the NSG mice intravenously. Subsequently, 2 days post-injection, either AVA4746 or PBS was administered by oral gavage at 15 mg/kg twice a day continuously for 14 days (Fig. 6A). A significant prolongation of overall survival in the group treated with a combination of AVA4746 and VDL (n=6; MST, 78.5 days) compared with that in the VDL-only group (n=6; MST, 68 days; Fig. 6B). *P=0.01. (C) The same protocol as that performed in (A) was conducted, except for the dose of AVA4746 was higher (30 mg/kg) and the treatment time was longer (twice a day for 28 days). (D) Kaplan-Meier survival analysis for the experimental design shown in (C). The MST was 46.5 days for the PBS group (n=4), whilst the MST was 46 days for the AVA4746 group (n=5). The 77-day MST of the VDL-treated group (n=5) was significantly different compared with the 87-day MST in the group treated with the combination of both AVA4746 and VDL (n=5). *P=0.023. NSG, NOD.Cg-PrkdcscidIl2rgtm1Wjl/SzJ; MST, mean survival time; VDL, vincristine, dexamethasone and L-asparaginase; p.o., per os; i.p., intraperitoneal.

In addition, mouse xenograft models were established using cell lines derived from two other cases B-ALL (RS4;11 and TXL3). AVA4746 in combination with VDL (n=8; MST, 105 days) did not lead to a statistically significant improvement in the survival of mice injected with RS4;11 cells in the VDL-alone group (n=6; MST, 95 days; Fig. S10A and B). By contrast, AVA4746 alone (n=6; MST, 66 days) prolonged the survival of the TXL3 xenograft mice compared with that in the PBS control (n=5; MST, 59 days; Fig. S10C and D). However, there were no changes in the survival of mice in the group treated with AVA4746 + VDL compared with VDL alone in the TXL3 model.

Discussion

Integrin α4 has long been of interest in the hematology research field because it has been frequently reported to be a key mediator of adhesion by leukemia cells to the chemoprotective niche (18,20,21,32-34). NZM is a humanized mouse monoclonal IgG4κ antibody that recognizes the human α4 chain of both α4β1 and α4β7 integrin (35,36). Although the Food and Drug Administration has approved NZM for the treatment multiple sclerosis (MS), progressive multifocal leukoencephalopathy can arise in patients treated with NZM (37). In clinical trials, antibodies against natalizumab
have been reported in 11% patients with MS and 7% patients with Crohn’s disease (38,39). This issue can be averted using small molecules. Previously, the beneficial effects of a non-clinical grade small molecule inhibitor, TBC3486 (21), was reported in a xenograft mouse model of primary B-ALL. Therefore, in the present study, the effects of AVA4746, a small molecule antagonist of VLA-4, was evaluated in primary B-ALL.

The epitopes that can be targeted by the HUTS-21 antibody, termed ligand-induced binding site epitopes, becomes exposed as a result of the conformational change that occur following activation, which can be exploited to discover novel allosteric VLA-4 antagonists (27). Therefore, the conformation-sensitive anti-CD29 HUTS-21 antibody was used to measure the VLA-4 RO, which indicates ligand-binding affinity measured in a scale ranging from 0 to 100%, as previously described (31). VLA-4 RO is upregulated by the divalent cation Mn²⁺ (31). AVA4746 was found to have a low EC₅₀ for VLA-4 RO compared with that of VCAM-1 without Mn²⁺, suggesting that AVA4746 binds to integrin α4 with higher affinity compared with VCAM-1. In addition, the prominent detachment effect on B-ALL from VCAM-1 mediated by AVA4746 was consistent with the effects of TBC3486 previously observed (21). Therefore, AVA4746 may be a useful tool for the studying VLA-4/VCAM-1 interactions, which are important for cell-cell adhesion between B-ALL and stromal cells (20).

Subsequently, it was demonstrated that AVA4746 can specifically target human integrin α4 on the cell surface whilst also downregulating the protein expression level of α4 in general. Previous studies have demonstrated that integrins are predominantly degraded through the endosome-lysosomal pathway (40,41). However, data from the present study showed that there was little or no degradative effects from the proteasome pathway, since inhibiting the proteasome using MG132 only partially reversed the downregulation of α4. In addition, there appeared to be no effect mediated by AVA4746 on AKT signaling, which is one of the pathway previously reported to be required stromal cell-mediated chemotherapy protection (42). However, changes in the phototyrosine levels were observed following AVA4746 treatment, which warrants further study. According to previous studies, VCAM-1 alone could not confer resistance to chemotherapy, suggesting the requirement for stromal cells capable of responding to activated leukemic cells (18,34). Similar results were found in the present study, where AVA4746 treatment did not result in any additional effects on B-ALL apoptosis induced by with VDL in vitro.

In addition, it was observed that AVA4746 treatment increased the percentage of CD45⁺/CD19⁺ leukemia cells in the BM, whilst the percentage of CD45⁺/CD19⁺ leukemia cells was decreased in the PB. In endothelial cells, VCAM-1 clustering, either by antibody crosslinking or integrin binding, triggers the activation of Rac1, a Rho-like GTPase (43). The activation of Rac1 results in the rearrangement of the cytoskeletal network, which then remodels the tight junctions among the vascular endothelial cells and mediate transendothelial migration (TEM) (44). Previously, it was demonstrated that overexpression of macrophage inflammatory protein-1α upregulated the expression of the endothelial adhesion molecule VCAM-1 to enhance TEM by ALL cells (45). Therefore, in the present study the effects of AVA4746 on endothelial cell physiology and TEM were investigated. Kim et al (46) previously showed that a specific monoclonal antibody against VCAM-1 was able to block VCAM-1-mediated cell-cell contacts and suppress angiogenesis (46). Similarly, the present study demonstrated that both AVA4746 and TBC3486 attenuated the angiogenesis of HUVECs in vitro. Therefore, one possible hypothesis from this observation is that the B-ALL cells remained in the BM despite their detachment from the stromal cells due to the anti-angiogenesis effects of AVA4746.

According to a previously established in vivo B-ALL xenograft model and chemotherapy regimen (20), AVA4746 in combination with VDL markedly prolonged the survival of mice engrafted with LAX7R cells, which were derived from a patient with primary relapsed B-ALL, but not in mice engrafted with RS4;11 and TXL3 cells. Unlike NZM, which is a pan-α4-integrin antibody that can not only inhibit α4/VCAM-1 but can also inhibit the α4/fibronectin (47) and α4/osteopontin (14) interactions, the VCAM-1 small molecule mimic AVA4746 did not prolong the survival of mice for as long as NZM (20). However, it was found in the present study that interfering with the VLA-4/VCAM-1 interaction can enhance the sensitivity of B-ALL to chemotherapy. One explanation for the contradictory observations of the effects of AVA4746 in vitro and in vivo is the possibility that oral bioavailability is not as robust in the NSG mice used in the present study. High receptor occupancy by integrin antagonists is required for the sufficient inhibition of adhesion (31). Therefore, it remains possible that sufficiently high levels of AVA4746 were not maintained throughout the course of the in vivo study in NSG mice. A follow-up study is required to confirm this possibility, which should include additional, in-depth cell adhesion studies involving different extracellular matrix components for α4 integrin, including fibronectin and osteopontin (14).

Taken together, AVA4746 is a potent antagonist of VLA-4 that can efficiently induce the detachment of B-ALL cells from VCAM-1. It is possible that AVA4746 can exert effects on normal hematopoietic stem and progenitor cells in the bone marrow, since these progenitor cells also express integrin α4 (48). However, this possibility is beyond the scope of the present study. The present study showed that the interruption of VLA-4/VCAM-1 by AVA4746, an orally applicable small molecule inhibitor, can sensitize LAX7R cells derived from a patient with relapsed B-ALL to chemotherapy. Therefore, AVA4746 hold promise a favorable tool for the investigation of VLA-4/VCAM-1 interactions in the CAM-DR mechanism of B-ALL.

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Availability of data and materials

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Authors’ contributions

DB, ASW and YMK were responsible for the conceptualization of the present study. YR, EJG, MY and YMK designed the present study. YR, EJG, HNK, SL and HAO performed the research, analyzed and interpreted the data. HCL and MY analyzed the data. All the authors reviewed the draft manuscript, edited the final draft and approved the final version for submission. YMK, YR, EJG, HNK, SL and HAO confirm the authenticity of all the raw data. All authors have read and approved the final manuscript.

Ethics approval and consent to participate

Bone marrow and peripheral blood samples from patients with B lineages acute lymphoblastic leukemia were acquired in compliance with the Institutional Review Board of Children's Hospital Los Angeles regulations. The human studies were conducted in accordance with the Declaration of Helsinki. Informed consent was obtained from the patients under the approved IRB protocol (approval no. CCI‑08‑00102). All studies were approved by the Institutional Animal Care and Use Committee of Children’s Hospital Los Angeles (Los Angeles, USA).

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

References


