Tetraspanin CD82 drives Acute Myeloid Leukemia chemoresistance by modulating Protein Kinase C alpha and β1 integrin activation

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Abstract

A principal challenge in treating acute myeloid leukemia (AML) is chemotherapy refractory disease. As such, there remains a critical need to identify key regulators of chemotherapy resistance in AML. In this study, we demonstrate that the membrane scaffold, CD82, contributes to the chemoresistant phenotype of AML. Using an RNA-seq approach, we identified the increased expression of the tetraspanin family member, CD82, in response to the chemotherapeutic, daunorubicin. Analysis of the TARGET and BEAT AML databases identifies a correlation between CD82 expression and overall survival of AML patients. Moreover, using a combination of cell lines and patient samples, we find that CD82 overexpression results in significantly reduced cell death in response to chemotherapy. Investigation of the mechanism by which CD82 promotes AML survival in response to chemotherapy identified a crucial role for enhanced protein kinase c alpha (PKCα) signaling and downstream activation of the β1 integrin. Additionally, analysis of β1 integrin clustering by super-resolution imaging demonstrates that CD82 expression promotes the formation of dense β1 integrin membrane clusters. Lastly, evaluation of survival signaling following daunorubicin treatment identified robust activation of p38 mitogen-activated protein kinase (MAPK) downstream of PKCα and β1 integrin signaling when CD82 is overexpressed. Together, these data propose a mechanism where CD82 promotes chemoresistance by increasing PKCα activation and downstream activation/clustering of β1 integrin, leading to AML cell survival via activation of p38 MAPK. These observations suggest that the CD82-PKCα signaling axis may be a potential therapeutic target for attenuating chemoresistance signaling in AML.
**Introduction**

Acute myeloid leukemia (AML) is characterized by the disruption of myeloid differentiation and accumulation of blast cells in the bone marrow. In 2019, AML is estimated to be the most prevalent newly diagnosed leukemia and the leading cause of leukemia-related mortality (1). Response to therapy differs widely between AML subclasses due to the highly heterogeneous characteristics of the disease at both the phenotypic and molecular level. The standard induction therapy for AML: cytarabine for seven consecutive days and daunorubicin on the first three days, is effective in a small subset of AML patients, but most patients are not cured and long-term disease-free survival remains poor (2). With a chemotherapy-only treatment regimen, less than 40% of adult patients achieve disease-free survival greater than 5 years; the disease free-survival rate is further reduced in younger patients (3). Chemoresistance is largely responsible for treatment failure and overall reduced survival in AML. Therefore, there is a critical need to identify key regulators of chemotherapy resistance in AML.

While specific mechanisms of drug resistance in AML remain poorly defined, many studies suggest that resistance may be a result of multiple factors. Some of the most recognized mechanisms of drug resistance in AML include genetic alterations, drug resistance-related proteins, miRNAs alterations, and aberrant activation of drug resistance-related signaling pathways (4, 5). For example, Fms-like tyrosine kinase 3 (FLT3) mutations found in one-third of patients with AML constitutively activates the kinase resulting in aberrant proliferation, which supports AML cell survival upon treatment with conventional chemotherapeutics (6, 7). The activation of additional signaling pathways including PI3K/AKT, JAK/STAT and NF-κB have also been reported as survival mechanisms for drug resistance in AML (8–11).

Tetraspanins, a family of membrane scaffolding proteins, function as regulators of cellular signaling through the organization of membrane microdomains consisting of membrane receptors and effector signaling molecules (12). Tetraspanins were shown to be involved in tumor progression at many stages including: angiogenesis, metastasis, and therapy resistance (13). With respect to AML, Tetraspanin 3 was recently demonstrated to play a role in the development of de novo AML (14), whereas tetraspanin CD81 was identified as an unfavorable prognostic marker in AML (15).

The tetraspanin CD82 was also found to be upregulated in several leukemias, including AML (16), with aberrant CD82 expression detected specifically in chemotherapy-resistant CD34+/CD38− AML cells (17, 18). Additionally, work from our group identified CD82 as a key regulator of AML cell homing to the bone marrow (19) and demonstrated that CD82 modulates the spatial and temporal dynamics of Protein Kinase C α (PKCα) signaling in AML (20). Interestingly, PKCα expression was shown to correlate with poor survival in AML patients and contribute to cancer progression (21). Moreover, targeting PKC-mediated signaling stimulated an increase in apoptosis of AML cells (22). In the present study, we identify CD82 as a tetraspanin upregulated in response to chemotherapy in AML. Moreover, we demonstrate that tetraspanin CD82 promotes chemoresistance in AML cells and delineate a mechanism involving integrin β1 and p38 MAPK activation downstream of
PKCa signaling, providing valuable targets for further development of therapies to treat chemotherapy-resistant AML.

**Results**

**Chemotherapy increases the expression of tetraspanins CD82 and CD53 in AML.**

To evaluate how tetraspanin expression is modulated by AML cells in response to chemotherapy, we completed a RNA-sequencing (RNA-seq) study using the AML cell line (KG1a) treated with the conventional chemotherapeutic, daunorubicin. Analysis of tetraspanin family members identified transcriptional alterations following chemotherapy (Fig. 1A). Most notably, daunorubicin treatment stimulated the expression of tetraspanins CD82 and CD53 (Fig. 1A). Next, we subjected multiple AML cell lines (KG1a, Kasumi-1 and HL-60) representing the most prevalent and undifferentiated stem-like M1 and M2 subtypes (Fig. 1B, Supp. Table 1), to treatment with daunorubicin and measured the surface expression changes of both CD82 and CD53 (23–26). Flow cytometry analysis established that CD82 expression is significantly increased on the surface of all cell lines analyzed in response to chemotherapy (Fig. 1C,D). In contrast, CD53 expression is increased in the KG1a cells only, with no differences detected in the Kasumi-1 or HL-60 cells (Fig. 1E,F). Lastly, two AML patient samples were analyzed for CD82 expression before and after daunorubicin treatment, again identifying a significant upregulation of surface CD82 expression following chemotherapy (Fig. 1G). Collectively, these data suggest that the tetraspanin CD82 is upregulated in response to chemotherapy and thus may stimulate a critical survival signaling cascade.

**Increased CD82 expression in AML patient samples correlates with reduced overall survival.**

To determine whether CD82 overexpression could be an indicator of patient outcomes, we analyzed the Therapeutically Applicable Research to Generate Effective Treatment (TARGET) AML database, which included RNA-seq analysis of 152 primary AML patient samples, of which 125 samples were collected from bone marrow aspirates and categorized into even-sized groups based on tertiles. Examination of survival outcomes based on CD82 (Fig. 2A) or CD53 (Fig. 2B) expression identifies poor overall survival in patients aged 0-24 with increased CD82 expression, but not CD53. To investigate how CD82 expression impacts adult AML patient outcomes, we analyzed overall survival data from the BEAT AML trial (310 patients, ages 25-74). Patients were further classified into two age groups of 25-49 (n=86) and 50-74 (n=224). Increased CD82 expression predicted an increased hazard ratio (HR) of 1.4 (CI: 0.72-2.89) for overall survival ages 25-49 (Fig. 2C) and a smaller ratio of 1.1 (CI: 0.76-1.52) in the 50-74 group (Fig. 2E). In contrast, comparison of survival outcomes between CD53 high to low expression predicted a HR of 1.2 (CI: 0.68-2.50) for ages 25-49 (Fig. 2D) and 1.0 (CI: 0.76-1.48) for ages 50-74 (Fig. 2F). In combination, our analyses indicate a correlation between increased CD82 expression and reduced overall survival in both pediatric and adult AML patients, further implicating a role for CD82 in AML survival.
CD82 overexpression promotes AML chemoresistance.

Recognizing the critical role of chemoresistance in overall AML survival and relapse, we went on to determine the impact of CD82 expression on AML cell response to chemotherapy in vitro. AML cell lines (KG1a and Kasumi-1) were generated with CD82 overexpression (CD82OE), CD82 knock down (CD82KD), and vector control cells (control). Cells were treated with standard induction therapy agents daunorubicin (1.7µM) or cytarabine (0.5µM) for 72 hours. Caspase 3/7 activation analysis by flow cytometry indicates that CD82OE cells display reduced apoptosis in response to chemotherapy when compared to control cells (Fig. 3A–D). Moreover, CD82KD cells display enhanced cell death following chemotherapy treatment, further implicating an important role for CD82 expression in AML response to chemotherapy (Fig. 3A–C). CD82OE cells also show a decrease in apoptosis as detected by confocal imaging of annexin V positive cells (green) following daunorubicin treatment (Fig. 3E). Taken together, these data support the findings that CD82 expression modulates chemotherapy-mediated apoptosis in AML cells.

PKCα is required for CD82-mediated chemoresistance.

To investigate the mechanism by which CD82 promotes AML chemoresistance and cell survival, we completed RNA-seq experiments with the CD82OE cells treated with and without daunorubicin. When compared to control cells, the combination of daunorubicin treatment and CD82 overexpression led to significant transcriptomic alterations, with over 6000 genes differentially expressed with both P-value and false discovery rate (FDR) <0.05. Principal component analysis displays differential clustering between control and CD82OE cells following vehicle or daunorubicin treatment (Fig. 4A). Of particular interest, PRKCA gene expression was increased upon CD82 overexpression and further increased following daunorubicin treatment as indicated on the heat map (Fig. 4A). Our previous work identified a critical role for CD82 in the regulation of PKCα activation (20), thus we went on to measure potential changes in PKCα expression and activation in response to chemotherapy treatment. Western blot analysis indicates that CD82 overexpression significantly increases phospho-PKCα expression following daunorubicin treatment (Fig. 4B–E). To determine whether the expression and/or activation of PKCα contributes to the chemoresistant phenotype of CD82OE cells, we utilized both pharmacologic (Fig. 4F) and genetic inhibition (Fig. 4G,H) of PKCα and evaluated cell death response to daunorubicin. Caspase activity assays demonstrate that while PKCα inhibition has no effect on cell death of control cells or CD82KD cells (Supp. Fig 1), PKCα inhibition in combination with daunorubicin increases cell death of CD82OE cells, restoring chemosensitivity to that of control cells. Therefore, PKCα is a critical signaling component of CD82-mediated chemoresistance in AML.

Chemotherapy activates β1 integrin downstream of PKCα.

PKCα regulates a number of key biological events including proliferation and differentiation (27–30). Moreover, PKCα interacts with a number of downstream targets that support cell survival. Further analysis of the transcriptional patterns of CD82OE cells following daunorubicin treatment identified enrichment of the MAPK signaling for integrins biological pathway (Fig. 5A). This transcriptional signature led us to investigate the impact of CD82
expression on the integrin signaling activity of daunorubicin-treated AML cell lines. Flow cytometry analyses of total surface β1 integrin show minimal expression changes between control or CD82OE cells with or without daunorubicin treatment (Fig. 5B). In contrast, CD82 overexpression and daunorubicin treatment both increase the activation of β1 integrin in the KG1a and Kasumi-1 cells with the greatest increase in β1 activity detected in the CD82OE cells treated with daunorubicin (Fig. 5C–E). We also analyzed the β1 activity of AML patient samples following daunorubicin treatment, identifying a 1.4-fold and 1.8-fold increase in β1 activity in patients 39 and 75, respectively (Fig. 5F). Taken together, daunorubicin treatment and CD82 overexpression both stimulate the activation of β1 integrin. Recognizing the critical role PKCα activity plays in both upstream (inside-out) and downstream (outside-in) integrin signaling, we repeated the β1 activity assay in the presence of PKCα inhibition. Both pharmacologic (Fig. 5C) and genetic (Fig. 5D) inhibition of PKCα attenuated the β1 activation observed upon CD82 overexpression and daunorubicin treatment, suggesting that PKCα signaling is upstream of chemotherapy-induced β1 activation. PKCα promotes talin-1 activation and binding to the β1 integrin tail, which is a critical final step in inside-out integrin activation stimulating β1 activation (31). Next, we quantified talin-1 phosphorylation, finding a significant increase in talin-1 activation in CD82OE cells that is further increased upon daunorubicin treatment (Fig. 5G). Lastly, to confirm that β1 activation is a critical player in CD82-mediated chemoresistance, we incorporated BIO 5192, a previously characterized selective small molecule inhibitor of both the activated and inactivated forms of α4β1 integrin (32). Figure 5H displays increased cell death in the BIO 5192-treated CD82OE cells upon daunorubicin treatment, consistent with a recovery of chemosensitivity. Collectively, these data suggest that PKCα-stimulated inside-out signaling of the β1 integrin is the primary signaling pathway responsible for CD82-mediated AML chemoresistance.

CD82 expression alters β1 integrin membrane clustering.

Inside-out signaling can also stimulate the clustering of integrins in the membrane, which can modulate integrin avidity, promoting cell adhesion and signaling (33, 34). Therefore, we next used the super-resolution imaging (SRI) technique, direct stochastic optical reconstruction microscopy (dSTORM), to measure changes in β1 integrin membrane organization following CD82 overexpression and/or chemotherapy treatment. Figures 6A,B illustrate representative dSTORM images that were analyzed by the density-based spatial clustering of applications with noise (DBSCAN) algorithm to quantify β1 clustering (35). DBSCAN analysis identified significant changes in the organization of β1 clusters, finding that CD82 overexpression promotes an increased number of smaller-sized clusters when compared to control cells (Fig. 6C,E). Moreover, nearest neighbor analysis suggests that β1 clusters are in closer proximity with each other (Fig. 6D) and more densely packed (Fig. 6F) upon CD82 overexpression. In contrast, daunorubicin treatment appears to have minimal impact on β1 integrin membrane distribution (Fig. 6B–F). Therefore, when taken together, SRI analyses suggest that CD82 overexpression significantly contributes to β1 integrin organization, which can modulate downstream survival signaling.
**p38 MAPK is activated upon chemotherapy treatment in AML.**

Chemotherapy agents stimulate DNA damage and are also strong activators of the p38 MAPK pathway (36, 37). Activation of p38 can enhance cancer cell survival, with increasing evidence suggesting that p38α facilitates tumor chemoresistance (38). Integrin signaling is frequently linked to p38 activation (39, 40), thus, we evaluated p38 signaling downstream of PKCα and β1 integrin. Using western blot analysis, we measured total and phospho-p38 following chemotherapy treatment alone or in combination with PKCα or β1 integrin inhibition. We find no changes in total p38 expression (Fig. 7A,B), but detect a significant increase in phospho-p38 expression upon chemotherapy treatment in control cells that is enhanced by CD82 overexpression (Fig. 7A,C). Inhibition of PKCα in combination with chemotherapy treatment significantly reduced the activation of p38 detected, supporting a role for p38 activation downstream of PKCα signaling (Fig. 7A,C). Additionally, inhibition of α4β1 combined with chemotherapy also shows a reduction in phospho-p38 when compared to chemotherapy treatment alone (Fig. 7D,F), with no change in total p38 expression (Fig. 7D,E). Lastly, we went on to pharmacologically inhibit p38 activity and analyze response to chemotherapy. While we detect no change in response to chemotherapy in control cells, we measure an overall increase in chemosensitivity in CD82 overexpressing cells upon p38 inhibition (Fig. 7G). Collectively, these data demonstrate that CD82 overexpression promotes chemoresistance in AML cells via p38 activation downstream of PKCα and β1 integrin mediated signaling (Fig. 7H).

**Discussion**

A major challenge in treating AML is the development of resistant disease resulting in relapse. Numerous studies have focused on identifying important molecules and pathways associated with chemoresistance mechanisms in aggressive hematologic malignancies with the goal of eliminating disease relapse. Aberrant activation of drug resistance-related signaling proteins represents a critical mechanism in AML chemoresistance currently being explored (5). In this study, we investigate AML cell signaling response to chemotherapeutics with a focus on the tetraspanin family of membrane scaffold proteins.

Tetraspanins are widely and abundantly expressed in multiple malignancies and have been described to promote various cancer stages including initiation (13), progression (14), and metastasis (12, 16, 19). Additionally, evidence suggests supporting roles for tetraspanins in cancer drug resistance. Specifically, CD151 (41), Tspan12 (42), CD9 and CD81 (43, 44) have all been shown to modulate cancer cell response to various chemotherapy agents. Using RNA-seq to pursue an unbiased assessment of potential tetraspanins involved in chemoresistance, we identify the upregulation of two specific tetraspanins, CD53 and CD82. Further analysis of cell lines and patient samples illustrates that CD82 expression is significantly enhanced following chemotherapy across all samples analyzed. Our subsequent analysis of the TARGET and BEAT AML databases identified a significant correlation between CD82 expression and overall AML patient survival. These aggregate findings led us to focus our study on CD82, which has been shown to have key signaling roles in AML (18, 45–48).
CD82 was originally identified as a suppressor of tumor migration, invasion and metastasis in malignancies including prostate, breast and lung (49–51). In hematology, CD82 was identified on early hematopoietic progenitor cells and found to be upregulated in distinct human leukemias including AML (16). CD82 was shown to regulate adhesion, survival and bone marrow homing of AML (52) and identified in a proteomics screen to be enriched in the plasma membrane of leukemia stem cells (48). In the current study, we find that CD82 overexpression in AML reduces apoptosis in response to conventional chemotherapeutics. Furthermore, we find that chemosensitivity is restored following CD82 knock down. Thus, CD82 expression modulates AML response to chemotherapy. Our findings are supported by a recent study investigating pediatric AML, which identified an upregulation of CD82 gene expression in AML cells isolated from the bone marrow and went on to find that increased CD82 expression promoted adriamycin chemotherapy (46). Collectively, these data suggest that CD82 expression has the capacity to regulate AML survival signaling in response to chemotherapy.

Previously, STAT5 (45, 47) and Wnt/β-catenin signaling (46) were described to impact CD82-mediated survival signaling of AML. Our RNA-seq analysis of CD82OE cells identified significant expression changes in 6810 genes, including upregulation of PKRCA following daunorubicin treatment. Increased PKCα activation was correlated with poor survival rates in AML patients (21) and increased AML cell viability (27, 53). Work from our lab discovered that the CD82 membrane scaffold stabilizes activated PKCα at the plasma membrane in response to stimulation, resulting in aberrant sustained ERK activation (20). Here, we find that CD82 overexpression contributes to enhanced PKCα activation following chemotherapeutic treatment, and that inhibition of PKCα can restore chemosensitivity of CD82OE cells. These data suggest that PKCα activation is a critical regulator of CD82-mediated chemoresistance. The classical PKC isofoms alpha and beta are known to support survival signaling, and thus have been explored as potential targets for anti-cancer therapies. Enzastaurin, the PKCβ inhibitor, was shown to stimulate apoptosis in AML-derived cell lines and in blast cells from AML patients at concentrations that also inhibit PKCα (53). Clinical trials involving PKC inhibitors as single therapies have been inconsistent (54–56), however, the results of our study suggest that stratification of patients based on CD82 expression could potentially improve response to PKC specific therapies.

PKC activation functions in both inside-out and outside-in integrin signaling, which can contribute to chemoresistance (57–60). Pathway analysis of the transcriptional signature of CD82OE cells following daunorubicin treatment identified the upregulation of MAPK signaling for integrins. In the presence of environmental and therapeutic stresses, integrins can function in the absence of ligand binding to promote survival and a “stemness” phenotype (61). We find that chemotherapy treatment and CD82 overexpression increase β1 activation in AML cell lines and patient samples. Interestingly, β1 activation is reduced back to basal signaling upon PKCα inhibition by pharmacologic or genetic knock-down, suggesting that PKCα is signaling upstream of β1 integrin activation. Also consistent with inside-out signaling, we detect talin-1 activation in CD82OE treated with daunorubicin. Additionally, the restored chemosensitivity of CD82OE cells following treatment with the α4β1 inhibitor is supported by recent findings in T-Acute Lymphoblastic Leukemia where blockade of β1 integrin sensitized xenografted leukemic cells to doxorubicin and diminished

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leukemic burden in the bone marrow (62). Collectively, our work supports a previously proposed concept where a tetraspanin-PKC-integrin complex is essential for signaling (63) and goes on to suggest that the level of CD82 expression serves as a modulator of signal intensity and perhaps signal duration critical for chemoresistance.

Membrane clustering of integrins is fundamental for integrin signaling where interactions with scaffold proteins, including tetraspanins, have been shown previously to modulate the formation of integrin clusters (64). Using SRI, we identify significant changes in β1 cluster organization and density when CD82 is overexpressed in AML cells. Specifically, we find an increase in β1 integrin packing into smaller scale clusters upon CD82 overexpression, which is likely to contribute to the measured increase in β1 activation. Previous work from our group identified similar density changes with integrin α4 upon CD82 overexpression, which significantly altered cell adhesion (64) and we went on to show that enhanced membrane clustering of PKCα, promoted by CD82 resulted in a sustained signaling response (20). Therefore, when taken together, CD82 not only promotes the stabilized signaling of PKCα, but also enhances the membrane clustering of integrins, which collectively can have significant impacts on downstream survival signaling.

The p38 MAPK pathway is activated in response to a host of cellular stressors. The well-defined role of p38 in cell growth inhibition and apoptosis has led to its characterization as a tumor suppressor protein, but p38 also functions in the less well-characterized function of cell survival (36). In fact, p38 often facilitates tumor cell survival in response to chemotherapeutic treatments (37, 38). Our data demonstrate that following daunorubicin treatment the p38 signaling pathway is activated in AML cells. More importantly, CD82 overexpression leads to a significant increase in p38 activation in response to daunorubicin. Inhibition of PKCα and α4β1 activation in combination with daunorubicin treatment resulted in attenuated p38 signaling. Moreover, p38 inhibition restored chemosensitivity of CD82 overexpressing cells. Thus, p38 inhibitors, which have been investigated in the treatment of myelodysplastic syndrome and AML (65, 66) may also provide benefit to AML patients with increased CD82 expression.

A better understanding of the cellular mechanisms that can be exploited to restore AML sensitivity to chemotherapy is a critical requirement to improving long-term treatments for patients. In this study we identify the CD82 scaffold as a modulator of survival signaling in response to chemotherapeutic agents, resulting in chemoresistance. Tetraspanins are currently being used in clinical trials for the treatment of other hematological malignancies (67) and our data suggest that targeting the CD82 scaffold may provide a mechanism to disrupt the aberrant signaling that contributes to chemotherapeutic response. Therefore, the ability to disrupt the CD82 membrane scaffold, which we show serves to stimulate PKCα, β1 integrin and p38 activation (three critical drivers of cell survival signaling) may represent a novel approach to treat chemoresistant AML.
Materials and Methods

Cell culture
KG1a, Kasumi-1 and HL-60 cells (American Type Culture Collection) were cultured in RPMI 1640 or IMDM supplemented with 10%FBS, 2 mM l-glutamine, 100 u/ml penicillin, and 100 μg/ml streptomycin. Nucleofections were performed using Amaxa Cell Line Nucleofector Kit V (Lonza) according to the manufacturer’s directions. G418 was used to generate stable cell lines. Stable CD82 knockdown was established using KG1a cells transfected with the CD82 shRNA plasmid (Santa Cruz Biotechnology, sc-35734-SH); cells were put under puromycin selection and sorted for negative CD82 surface expression. PKCα shRNA plasmid (Santa Cruz Biotechnology; sc-36243-SH) was used for PKCα knockdown.

RNA-Sequencing
Total RNA was extracted from KG1a WT and CD82OE cell lines treated with 1.7uM Daunorubicin or vehicle control using the RNeasy Mini Kit (QIAGEN). Synthesis of cDNA and library preparation were performed using the SMARTer Universal Low Input RNA Kit for Sequencing (Clontech) and the Ion Plus Fragment Library Kit (ThermoFisher) as previously described (68–70). Library was quality checked on an Agilent DNA High Sensitivity Chip and quantified using qPCR. Samples are diluted to 50-100 pM and pooled in equalmolar concentrations prior to loading into the Ion Chef for emulsion, enrichment, and chip loading. All libraries were sequenced on 540 chips using the Ion Proton S5/XL systems (Life Technologies) in the Analytical and Translational Genomics Shared Resource at the University of New Mexico Comprehensive Cancer Center. All samples were aligned with Torrent Mapping Alignment Program (TMAP, v5.2.25), to a BED file containing non-overlapping exons from UCSC genome hg19. Exon feature counts were generated with HTSeq-count (71). Normalized data was used for analysis of differentially expressed genes (DEGs) between the CD82OE and wild-type cells were identified by using R packages DESeq2 and edgeR (72, 73) using the following criteria: |log2 fold change (FC)| ≥2, for both P-value and false discovery rate (FDR) <0.05, heatmaps was generated using JavaTreeview and pathway analysis was conducted using Metascape.(74, 75). RNA sequencing data is available for download from the NCBI BioProject database using study accession number PRJNA601161

Patient bone marrow aspirate RNA sequencing data was available through the TARGET website (https://ocg.cancer.gov/programs/target) and available under database of Genotypes and Phenotypes (dbGaP) accession number phs000465 (n= 125). Beat AML trial data was available through the (http://vizome.org/aml/) (n=310).

Clinical Samples
AML patient samples were deidentified and obtained from the Human Tissue Repository and Cell Analysis Shared Resource at the University of New Mexico Health Sciences Center. All samples were collected with informed consent of the donors and studies were conducted in accordance with the principles of the Declaration of Helsinki. All studies were performed under Institutional Review Board-approved protocols.
Caspase-3/−7 activity assay

Cells treatments were as follows: 1.7μM Daunorubicin (chemotherapy, Sigma-Aldrich), 1μM Cytarabine (chemotherapy, Tocris), 1μM GO6976 (PKC inhibitor, Tocris), 20nM BIO5192 (VLA-4 inhibitor, Tocris) or 25nM BMS 582949 (p38 MAP kinase inhibitor, Cayman Chemical) individually or in combination as stated. Caspase-3/−7 activity was measured with the CellEvent™ Caspase-3/7 Green Flow Cytometry Assay Kit as indicated by manufacturer (Thermo Fisher). Cells were analyzed using an Accuri C6 flow cytometer (BD Bioscience) and percent caspase positive cells were normalized to unstained controls.

Flow Cytometry

Cells were labeled with antibody or the corresponding isotype control in 1%BSA/PBS for 30 mins on ice. Cells were washed and analyzed using an Accuri C6 flow cytometer (BD Bioscience); histograms were generated using FlowJo software. Mean fluorescence values were normalized to the control cell line level. Antibodies used were CD82-Alexa-647 (Biolegend, ASL-24), CD53-PE (BioLegend, HI29), and Integrin β1-Alexa-647 (BioLegend, TS2/16). For active β1 integrin expression, cells were labeled with Ligand-induced binding site (LIBS) Anti-Integrin β1 Antibody (Millipore, HUTS-4) for 30 mins. Cells were washed and labeled with goat anti-mouse Alexa-647 secondary antibody (Invitrogen).

Immunofluorescence

Control or CD82OE cells were treated as indicated for 48 hours. Cells were blocked in 3%BSA in PBS followed by incubation with primary antibodies (AnnexinV-FITC, 1:200, Biolegend; CD82-Alexa647, 1:200, Biolegend) and DAPI (Invitrogen). Cells were washed and imaged with a Zeiss airyscan (LSM 800) system (Carl Zeiss) using excitation wavelengths of 405, 488 or 633 nm and a 63X/1.2 numerical aperture oil immersion objective.

Western Blotting

Western blots were performed as previously described (64). Antibodies used for Western blotting were purchased from Cell Signaling Technology as follows: calnexin (C5C9), PKCa (2056), phospho-PKCa/β (Thr638/641, 9375), Talin-1 (C4SF1), Phospho-Talin-1 (Ser425, D2P2M), p38 (D13E1), phospho-p38 (Thr180/Tyr182, D3F9), or β-Actin (Sigma, AC-74); all antibodies were used at a 1:1000 dilution. Horseradish peroxidase conjugate enzymes were stimulated with SuperSignal West Pico Chemiluminescent Substrate or Femto Maximum Sensitivity Substrate (Life Technologies). Blots were imaged using the ChemiDoc XRS Imager (Bio-Rad) and analyzed using ImageJ (National Institutes of Health) densitometry software.

Super-resolution imaging

Cells were plated on fibronectin coated eight-well chamber slides and treated with 1.7 μM Daunorubicin or vehicle overnight at 37°C. Cells were fixed with 4% PFA, blocked with 3% BSA/PBS and labeled with Alexa Fluor 647 anti–human CD29 antibody (BioLegend, TS2/16, 1:200). Cells were washed and fixed again with 4% PFA. Labeled cells were
imaged in a reducing buffer composed of 50 mM Tris, 10 mM NaCl, 10% w/v glucose, 168.8 u/ml glucose oxidase (Sigma #G2133), 1404.0 U/ml catalase (Sigma #C9332), and 20 mM MEA, pH8.5. Cells were imaged using a custom TIRF microscope system as previously described (20, 64, 76). 40,000 frames were collected per cell, with a brightfield image acquired every 2,000 frames for registration and drift correction. Each frame was 256 × 256 pixels with a pixel size of 0.107 µm and an acquisition time of 10ms per frame. Instrumentation was controlled by custom-written software in Matlab.

Collected data were analyzed as previously described (77). DBSCAN algorithm was applied to the reconstructed images to assess clustering β1 integrin (parameters: ε = 50nm and n = 20). Analyses performed using other parameters (ε = 100nm and n = 30 ε = 50nm and n = 30) yielded similar clustering and the same trends as our final parameters. Three 5µm x 5µm ROIs were selected per cell for analysis. Clusters identified by DBSCAN were subjected to the ROUT method of outlier detection (Prism 8, GraphPad) with Q = 1%, and identified outliers were removed from further analysis. Cluster radii were calculated using a convex hull around all points identified as a cluster. Equivalent area of a cluster was calculated as area = pi*R^2. Density of clustered points was calculated as the number of localizations in a cluster divided by the cluster area.

Statistics

Data were analyzed by GraphPad Prism 6 or 8 software. Data are presented as the means ± SD/SEM from three independent experiments. At least three biological replicates were performed for each experiment, unless otherwise stated. Two-tailed Student’s t test was used to compare differences between two groups. For comparisons in 3 or more groups One-way ANOVA followed by Tukey’s post-hoc test for multiple comparisons. The Kaplan–Meier curves for survival analyses were analyzed using the Log-rank (Mantel-Cox) or Hazard Ratio (Mantel-Haenszel) tests. The Kolmogorov–Smirnov test was used for comparison of cumulative distributions. A P value <0.05 was considered statistically significant and significance was labeled as *p < 0.05, **p < 0.01, ***p < 0.001, and ****p < 0.0001.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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Figure 1: Chemotherapy increases the expression of tetraspanin CD82 in AML.
(A) Tetraspanin gene expression profile based on RNA-seq analysis of vehicle or daunorubicin (1.7μM) treated KG1a control cells following 24 hr. (B) Analysis of AML prevalence by French-American-British (FAB) classification subtypes. Flow cytometry histograms measuring surface expression of (C) CD82 or (E) CD53 following vehicle or daunorubicin (1.7μM) treatment for 24 hr. (D) Fold Change in protein expression of (D) CD82 or (F) CD53 in different AML cell lines treated with vehicle or daunorubicin (1.7μM) for 24 hr measured by flow cytometry (n=3). (G) Flow cytometry analysis of CD82 surface...
expression in AML patient samples following treatment with vehicle or daunorubicin (1.7μM) for 24 hr. Means ± SD (n = 2 in A, 3 in C-F) *p < 0.05, and ****p < 0.0001.
Figure 2: Increased CD82 expression in AML patient samples correlates with reduced overall survival.
Kaplan-Myer survival curve analysis for overall survival of 125 pediatric AML patients (Ages:0-24) from the TARGET database by (A) CD82 and (B) CD53 expression. Kaplan-Myer survival curve analysis for 310 adult AML patients (Ages:25-74) included in the BEAT AML trial based on (C, E) CD82 or (D, F) CD53 expression.
Figure 3: CD82 overexpression promotes AML chemoresistance.
Caspase activity was measured in KG1a cell lines differentially expressing CD82, which were treated with vehicle (A, B), 1.7μM daunorubicin or (C) 0.5μM cytarabine over 72 hours. (C) Caspase activity was measured in Kasumi-1 cells transfected with mCherry vector control or mCherry-CD82 and treated with vehicle or 1.7μM daunorubicin over 72 hours. (D) Control and CD82OE KG1a cells were treated with vehicle or 1.7μM daunorubicin for 24 hr and then fixed and stained for apoptosis marker, Annexin V (green), CD82 (red) and DAPI (blue) and imaged by confocal microscopy Scale bars: 10 μm. Means
± SD (n = 3 in A - E). 2-Way ANOVA statistical significance in B - D is in comparison to the matching time-point of vehicle treated control cells. *p < 0.05, **p < 0.01 and ****p < 0.0001.
Figure 4: PKCα is required for CD82-mediated chemoresistance

(A) Principal component analysis and condensed heatmap for control and CD82OE cells treated with vehicle or Daunorubicin. Total and phospho-PKCα levels were analyzed by Western blot in control and CD82OE cells: KG1a (B, C) and Kasumi-1 (D, E). (F) Caspase activity measured in KG1a cells with PKCα inhibitor, Go 6976 (2.5 nM), alone or in combination with 1.7μM daunorubicin. (G) Western blot analysis for PKCα 24hr after shRNA transfection of KG1a cells. (H) Caspase activity measured in PKCα shRNA or scrambled shRNA treated control or CD82OE KG1a cells, which were treated with 1.7μM daunorubicin. Means ± SD (n = 2 in A, 3 in B - H) *p < 0.05, **p < 0.01 and ***p < 0.001.
Figure 5: Chemotherapy activates β1 integrin downstream of PKCα.

(A) Pathway analysis for genes upregulated in CD82OE cells upon treatment with daunorubicin. (B) Total and (C) active β1 integrin levels were analyzed by flow cytometry in control and CD82OE KG1a cells 24 hours after treatment with vehicle, daunorubicin (1.7μM), PKCα inhibitor, Go 6976 (2.5 nM), or Go 6976 (2.5 nM) and daunorubicin (1.7μM). (D) Active β1 integrin levels were measured by flow cytometry following PKCα shRNA or scrambled shRNA treatment of control and CD82OE KG1a cells, that were treated for 24 hr with vehicle or daunorubicin (1.7μM). β1 integrin activation was quantified.
by flow cytometry in (E) control or CD82OE Kasumi-1 cells and (F) primary AML patient samples following vehicle or daunorubicin (1.7μM) treatment for 24 hr. (G) Western blot analysis of total and phospho-Talin-1 levels analyzed in control and CD82OE KG1a cells 24 hours after treatment with vehicle or daunorubicin (1.7μM). (H) Caspase activity was measured in control and CD82OE KG1a cells following treatment with vehicle, daunorubicin (1.7μM), α4β1 integrin inhibitor, Bio 5192 (20 nM) or Bio 5192 (20 nM) in combination with daunorubicin (1.7μM) over 72 hours. Means ± SD (n = 2 in A, 3 in B – E, 2 in F, 3 in G and H) *p < 0.05, **p < 0.01, ***p < 0.001 and ****p < 0.0001.
Figure 6: CD82 expression alters β1 integrin membrane clustering.
Representative dSTORM images of control and CD82OE KG1a cells labeled for β1 integrin after treatment with (A) vehicle or (B) 1.7 μM daunorubicin. Blue boxes represent 5×5μm ROIs used for clustering analysis; scale bar: 5μm. Enlarged regions show β1 integrin clustering determined by DBSCAN analysis; parameters used were ε = 50nm and n = 20 points; scale bar = 200nm. Cumulative distribution plots of (C) cluster area and (D) cluster-to-cluster distance were derived from DBSCAN clustering data; statistical significance determined using the Kolmogorov-Smirnov test. (E) Average number of clusters per ROI.
and (F) average number of cluster localizations per 0.01μm² were determined using DBSCAN. For all analyses, n ≥ 9 cells per condition. Error bars, SEM; ***p < 0.001 and ****p < 0.0001.
Figure 7: Chemotherapy activates p38 MAPK downstream of PKCα and β1 integrin signaling. 

(A) Western blot analysis of total and phospho-p38 levels in control and CD82OE KG1a cells 24 hours after treatment with vehicle, daunorubicin (1.7μM), Go 6976 (2.5 nM), or Go 6976 (2.5 nM) in combination with daunorubicin (1.7μM). Densitometry analysis of total (B) and phospho-p38 (C) Western blots from (A). (D) Western blot analysis of total and phospho-p38 levels in control and CD82OE KG1a cells 24 hours after treatment with vehicle, daunorubicin (1.7μM), Bio 5192 (20 nM), or Bio5192 (20 nM) in combination with daunorubicin (1.7μM). Densitometry analysis of total (E) and phospho-p38 (F) Western
blots from (D). (G) Caspase activity was measured in control and CD82OE KG1a cells following treatment with vehicle, daunorubicin (1.7μM), p38 MAPK inhibitor, BMS 582949 (25 nM) or BMS 582949 (25 nM) in combination with daunorubicin (1.7μM) over 72 hours. (H) Schematic model of CD82 mediated chemoresistance (created using BioRender). Chemotherapy treatment (1) stimulates increased CD82 surface expression and PKCα activation (2), which promotes PKCα localization to the plasma membrane (3). Active PKCα stimulates the activation of β1 integrin via talin-1 mediated inside out signaling (4) and increased CD82 expression promotes β1 integrin clustering (5). p38 MAPK is activated downstream of PKCα and β1 integrin activation promoting increased AML cell survival (6). (n = 3 in A – F) *p < 0.05 and **p < 0.01 and ****p < 0.0001.