Assessment of epigenetic mechanisms and DNA double-strand break repair using laser micro-irradiation technique developed for hematological cells

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A B S T R A C T

Background: Certain tumors rely heavily on their DNA repair capability to survive the DNA damage induced by chemotherapeutic agents. Therefore, it is important to monitor the dynamics of DNA repair in patient samples during the course of their treatment, in order to determine whether a particular drug regimen perturbs the DNA repair networks in cancer cells and provides therapeutic benefits. Quantitative measurement of proteins and/or their posttranslational modification(s) at DNA double strand breaks (DSBs) induced by laser microirradiation provides an applicable diagnostic approach to examine DNA repair and its dynamics. However, its use is restricted to adherent cell lines and not employed in suspension tumor cells that include the many hematological malignancies.

Methods: Here, we report the development of an assay to laser micro-irradiate and quantitatively measure DNA repair transactions at DSB sites in normal mononuclear cells and a variety of suspension leukemia and lymphoma cells including primary patient samples.

Findings: We show that global changes in the H3K27me3-ac switch modulated by inhibitors of Class I HDACs, EZH2 methyltransferase and (or) H3K27me3 demethylases do not reflect the dynamic changes in H3K27me3 that occur at double-strand break sites during DNA repair.

Interpretation: Results from our mechanistic studies and proof-of-principle data with patient samples together show the effectiveness of using the modified micro-laser-based assay to examine DNA repair directly in suspension cancer cells, and has important clinical implications by serving as a valuable tool to assess drug efficacies in hematological cancer cells that grow in suspension.

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1. Introduction

Many new components of the DNA damage response were identified in the past two decades, which include proteins that function as DNA damage sensors or transducers, or those that act in the actual DNA repair. Because the majority of these factors act primarily at the sites of DNA breaks, they can be detected using immunofluorescence (IF) as coalesced structures called foci after exposure to DNA damaging agents. However, this standard IF approach fails to detect proteins that bind transiently to DNA break sites, and it does not permit the quantitative measurement of the spreading of DNA damage signal and the temporal examination of the assembly/disassembly of repair proteins at break sites. These limitations of the standard IF approach were overcome by the development of localized irradiation technique using micro-laser-fitted microscopes and its combination with immunofluorescence. Moreover, laser micro-irradiation has enabled the induction of breaks at discrete regions within the nucleus and has allowed the assessment of DNA repair dynamics in real time. Using the laser micro-irradiation technique, novel roles for several chromatin modifiers and remodelers at the sites of DNA damage were elucidated [1–3]. While this versatile approach has contributed significantly to the DNA repair field, its use has been restricted to adherent cells, particularly HeLa and U2OS cell lines [3–5], and it was not optimized for use in liquid tumors or suspension cells that include hematological malignancies, such as, leukemia and lymphoma.

Misregulated DNA repair contributes to the pathogenesis of and/or treatment resistance in many hematological cancers [6,7]. An underlying aberrant or altered epigenetic mechanism, such as histone post-translational modifications, contributes to the poor prognosis in many of these cancers. For instance, changes in histone H3 lysine-27...
Research in context

Evidence before this study

Investigations into DNA repair dynamics at DNA break sites using microlaser-induced DNA damage has provided valuable insights into various facets of the DNA repair process. However, the use of this powerful technique was restricted to solid tumors or adherent cells, and not employed in liquid tumors or suspension cells that include the many hematological cancers. We have previously used the microlaser technique to demonstrate the effects of histone deacetylase (HDAC) inhibitors on DNA repair in leukemia and lymphoma cells.

Added value of this study

We present here a detailed methodology to study DNA repair at microlaser-induced double strand breaks in a variety of suspension hematological cells that include normal blood cells and liquid tumors. We show that the microlaser technique can be used as an assay to examine the dynamics of DNA repair in hematological cancer cells. Using the optimized methodology, we provide insights into the mechanisms that regulate H3K27me3 epigenetic mark at double-strand break sites during DNA repair in hematological cancer cells. Overall, this study is of value to the HDACs, DNA repair, epigenetics and chromatin fields, and it also has an important application in the field of cancer therapeutics—by serving as a diagnostic tool to evaluate treatment efficacy in either preclinical or clinical setting.

Implications of all the available evidence

Using the refined microlaser technique and small molecule inhibitors, we demonstrate the mechanistic interplay between epigenetic enzymes—HDAC1,2, EZH2 H3K27 methyltransferase and H3K27me3 demethylases—during DNA repair in suspension cancer cells. Therefore, findings from our study show that the microlaser technique can be used to examine epigenetic and other regulatory mechanisms that occur during DNA repair, directly in hematological cells. Thus, obviating the need to extrapolate results obtained from adherent cancer cells to liquid tumors. Moreover, the methodology described here provides a diagnostic test in clinics, to evaluate the efficacies of drugs, especially those targeting aberrant DNA repair, and guide treatment plans with mono- or combination therapies for hematological cancers.

trimethylation (H3K27me3) are associated with poor prognosis in lymphoid malignancies [8–10]. Overexpression of EZH2 H3K27 methyltransferase occurs in many lymphoid malignancies including Burkitt lymphoma, follicular lymphoma, diffuse large B-cell lymphoma (DLBCL), mantle cell lymphoma and multiple myeloma, and EZH2 gain-of-function mutation is observed in germinal center-derived DLBCL and follicular lymphomas [11,12]. UTx, the demethylase for H3K27, is mutated in multiple hematological malignancies [13] and its inhibition is proposed as a potential epigenetic therapy for T-cell acute lymphoblastic leukemia [14]. EZH2 and H3K27me3 are linked to DNA double-strand break repair [15,16]. The H3K27 residue undergoes mutually exclusive acetylation (ac) and methylation modifications. We developed a method to laser micro-irradiate suspension cells, and using this modified approach, we previously reported a novel functional relationship between epigenetic enzymes, histone deacetylase 1 and 2 (HDAC1,2) and the EZH2 H3K27 methyltransferase, regulating the H3K27me3-ac switch during DNA repair in suspension EZH2 gain-of-function mutant DLBCL cells [3]. Translating the findings from the micro-laser-based assay, we reported that histone deacetylase (HDAC) inhibitors either in isolation or in combination with standard chemotherapies impair DNA repair, reset the altered H3K27me3-ac epigenetic switch and provide therapeutic benefits in the EZH2 gain-of-function (GOF) mutant DLBCL cells [3]. Therefore, based on this published study and given the importance of DNA repair in hematological malignancies and their drug response [6,7,17], it has become essential to develop a methodology to track and measure DNA repair dynamics in suspension cancer cells, which in turn could greatly enhance our ability to treat these cancers in an effective manner.

Here, we now present a detailed methodology to micro-irradiate a variety of suspension cancer cells including those obtained from primary patients with hematological cancers, and to detect DNA repair proteins or protein post-translational modifications at the sites of laser-induced DNA damage directly in suspension cells. We further highlight the use of the modified micro-laser technique for mechanistic studies by demonstrating the effects of small molecule inhibitors of HDACs, EZH2 or UTx on the deposition of H3K27me3 mark at DNA break sites in suspension cells. Overall, the micro-laser technique described here provides a powerful system to further our fundamental understanding of DNA repair mechanisms in suspension cancer cells, while advancing the clinical applicability of the DNA repair-based assays as biomarkers for cancer treatment.

2. Materials used in the study

2.1. Cell culture

Human lymphoma cells were maintained in culture medium in an exponentially growing state in a 37 °C, 5% CO2 incubator. Culture medium was prepared by adding 20% heat inactivated fetal bovine serum (FBS), 1% Penicillin-Streptomycin, and 1% L-Glutamine to RPMI 1640. Primary lymphoma patient cells and normal peripheral blood mononuclear cells were maintained in Mononuclear Cell Medium purchased from PromoCell. The patient sample culture was carried out in accordance with the University of Utah IRB guidelines. The samples are de-identified samples and hence we do not have access to the patient history. SupB15 leukemia cells were cultured in IMDM supplemented with 20% fetal bovine serum and penicillin-streptomycin.

2.2. Reagents used

Karpas-422 DLBCL cell line (Sigma, 06101702-1VL), SupB15 leukemia cell line (ATCC, CRL-1929), RPMI 1640 Medium (Caisson, RPL09-500ML), Fetal Bovine Serum (HyClone, SH30910.03), L-Glutamine (Corning, 25-005-CI), NaHCO3 (Sigma, S4019), NaOH Pellets (Mallinkrodt Analytical, 7708-10), Cell-Tak Cell and Tissue Adhesive (Corning, 354240), HCl (Fisher, A144-212), NaCl (Fisher, S671-3), KCl (Fisher, P217-500), Na2HPO4 (Fisher, S374-1), KH2PO4 (Fisher, P285-500), Hoechst 33342 Solution in Water (Thermo, H3570), Formalin 1:10 (Fisher, 23-245-685), Triton X-100 (Sigma, X100-1L), Normal Goat Serum (Sigma, G9023-10ML), Mouse α-yH2AX (Millipore, 05-636), Rabbit α-yH2AX (Cell Signaling, 2577), Rabbit α H3K27me3 Cell Signaling, 9733), Mouse α-pS1981-ATM (Rockland, 200–301–400S), Rabbit α-Cleaved Caspase-3 (Cell Signaling, 9661), Goat α Mouse Alexa 546 (Thermo-Life Technologies, A11030), Goat α Rabbit Alexa 488 (Thermo-Life Technologies, A11034), Goat α Mouse Alexa 488 (Thermo-Life Technologies, A11029), Goat α Rabbit Alexa 546 (Thermo-Life Technologies, A11035), Fluoromount-G (Southern Biotech, 0100–01). Sodium Bicarbonate Solution: Dissolve 0.42 g of NaHCO3 in 50 mL of sterile H2O to make a 0.1 M solution and add HCl to bring solution to pH 8. Cell Tak Solution: To prepare 100 μl of solution, 1.25 μl 1 M NaOH and 2.5 μl Cell-Tak was added to 96.25 μl 0.1 M NaHCO3 to obtain final concentrations of 12.5 mM NaOH and 5 μg Cell-Tak in 0.1 M NaHCO3.
2.3. Preparation of chromatin

Chromatin fractions from DLBCL and Pre-B-ALL cells were made as described previously [24]. Briefly, 3-4 × 10^6 cells were pelleted and resuspended in buffer A containing a mixture of 10 mM HEPES (pH 7.9), 10 mM KCl, 1.5 mM MgCl2, 0.34 M sucrose, 10% glycerol, 1 mM dithiothreitol and protease inhibitor cocktail. Triton-X-100 was added to the samples at a final concentration of 0.1% and incubated on ice for 8 min. Nuclei were collected by centrifugation at 8000 rpm for 5 min at 4 °C and lysed on ice for 30 min in buffer B (3 mM EDTA, 0.2 mM EGTA, 1 mM dithiothreitol and protease inhibitor cocktail). The chromatin fraction was separated from the soluble fraction by centrifugation at 10,000 rpm at 4 °C for 5 min. The supernatant was discarded and the chromatin pellet was resuspended in RIPA buffer with protease inhibitor and sonicated prior to immunoblotting.

2.4. Transient attachment of suspension cells to chamber slides

The Corning™ Cell-Tak reagent is a solution derived from the ‘polyphenolic proteins’ secreted by the marine mussel that enable it to anchor itself onto solid structures [42]. The Cell-Tak reagent acts as a cell adhesive, and therefore can be used to attach the suspension leukemia or lymphoma cancer cells onto the chamber dish. The Cell-Tak reagent was diluted in a solution of NaHCO3 and NaOH outlined above. We coated the chamber dish wells with this solution (100 μl/cm² well of an 8-well chamber) and left the chamber undisturbed in the tissue culture hood at room temperature for 30 min. Following this incubation, the solution was removed and wells were washed with sterile water prior to adding the cells. During the last 10 min of this 30-min coating period, cells that were either untreated or pre-treated with drugs were centrifuged (4000 rpm, 5 min) and re-suspended in 300 μl fresh culture media before adding them to the Cell-Tak-treated wells. Cells were then allowed to settle for 15–20 min in a 5% CO2 incubator at 37 °C. Following the attachment of cells to the dish and prior to micro-irradiation, Hoechst 33342 solution was added to a final concentration of 55 μM to stain DNA to identify the nucleus. We found that the timing between plating of cells and laser micro-irradiation is critical for the successful implementation of this protocol. Imaging of live cells with or without laser micro-irradiation showed that cells get washed off 4–8 h after their initial attachment to the Cell-Tak-treated chamber (Fig. 1A). Therefore, it is important to perform micro-irradiation immediately following a 15–20 min incubation of cells in the Cell-Tak coated culture dishes.

2.5. Laser micro-irradiation

For micro-irradiation, we used a 405 nm solid-state laser, and a 60× oil-immersion Plan Apo,1.4 numerical aperture (NA) objective with a pinhole of 1.8 airy units (AU) on a Nikon A1R (or Nikon A1) confocal microscope that was controlled using the Nikon Instruments Software (NIS-Elements). During irradiation, the microscope was programmed to utilize the laser to irradiate specific regions in the well. The areas targeted for laser irradiation are termed as regions of interest (ROIs) in the NIS-Elements software (Fig. 1B). Because B-cells (and their nuclei) are smaller in size compared to the adherent cells (such as HeLa or U2OS), we chose to use 24 ROIs for irradiation as opposed to 12 ROIs used for HeLa or U2OS cells, in order to irradiate the maximum number of B-cells (Fig. 1B–C).

Utilizing the NIS-Elements software, we created target areas using the ROI manager by selecting the “rectangle ROI” option and drawing rectangles across the field of view. We then duplicated the rectangle until there were 24 ROIs spanning the region (Fig. 1B), with each ROI measuring 208 × 1.7 μm². When creating these ROIs, it was important to ensure that they were labeled as “stimulation ROIs” in order for the laser to recognize them as areas to be irradiated. We set the 405 nm laser to irradiate the 24 ROIs spanning the field under view. Each well was irradiated over a period of five minutes and this time period included moving to each field of view within the same well. We focused on a corner of the well and then worked our way across the edge to irradiate 2–3 fields of view (Fig. 1C). We found that the ~2 min 5× irradiation cycle in addition to focusing on the next field of view allowed for 2–3 fields to be irradiated in the allotted 5 min. Following the 5 min of micro-irradiation, cells in each well were allowed to recover for an additional 10 min before fixing them with 3.7% formalin for 10 min.

2.6. Immunofluorescence

Immunofluorescence following micro irradiation was performed according the protocol described previously except that 0.1% instead of 0.5% Triton-X-100 was used [43]. The images were overlaid using the NIS elements software.

2.7. Statistical analyses

At least 100 cells with γH2AX lines per experiment were used to determine if H3K27me3 co-localizes with γH2AX and each experiment had at least three biological replicates. Averages and standard errors from three or more biological replicates were determined. All the p-values were calculated using the student’s two tailed t-test.

3. Results

3.1. Detection of DSB tracks in suspension cancer cells

Having optimized the conditions to immobilize suspension cells onto chamber slides using the Cell-Tak reagent (see methods section for details), we then determined the correct laser settings to induce double strand breaks. We used a wide range of laser power and found that the damage lines or tracks are most distinct when a 100% power of irradiation was used. We further validated the suitability of this high-power setting by micro-irradiating immobilized B-cells and performing time-lapse live cell imaging. A number of regions were subjected to micro-irradiation and each region was imaged every half hour for 8 h. We also performed multiple cycles of irradiation to identify the optimal dose of DNA damage for suspension cells. One cycle includes one round of irradiation of all the regions of interest (ROIs), and five cycles of 24 ROIs refers to zapping each of the 24 micro-lasered lines five times for a total of 120 irradiations over a particular field of view. No significant changes in cell morphology were observed in the irradiated regions under these conditions (Fig. 1D). We also performed immunostaining of laser irradiated cells to detect the cleaved form of caspase-3 (an apoptosis marker), which showed that cell viability was not compromised when laser irradiation was performed for 5 cycles, but a statistically significant number of cells undergoing active apoptosis were observed when irradiation was done for ≥10 cycles (Fig. 1D).

γH2AX or H2AX serine-139 phosphorylation is the gold standard for detecting DNA damage [18]. We manipulated the scan speed (that is, speed at which one field of view is imaged or irradiated) and pixel dwell (which is inversely related to the scan speed; a higher scan speed refers to a lower laser dwell time), in order to further examine if changing the irradiation intensity affected the formation of γH2AX lines. Our results showed that the settings detailed above can effectively cause laser-induced damage lines at both scan speeds and pixel dwell time (Fig. 2A). We also irradiated the same field of view with increasing number of cycles (2–8 cycles) and found that increasing the cycle number magnifies the spreading of the damage signal, as evidenced by the broadening of the γH2AX tracks with increasing cycle numbers (Fig. 2B). An increased number of cycles may be optimal when using an older laser, as overall laser power can decrease over time with usage of the microscope. We measured the time for γH2AX tracks to be formed or seen after laser irradiation. We find that it takes 5–10 min after laser irradiation to detect the γH2AX-labeled tracks (Fig. 2B). In our standard γH2AX immunofluorescence assays, we see
that it can take $>24$ h for the disappearance of radiation-induced $\gamma$H2AX foci (Supplementary Fig. 1). This poses a technical impediment to measure dephosphorylation of $\gamma$H2AX in our micro-laser assays, as the suspension cells start to detach from the assay plate after $4–8$ h following Cell-Tak-mediated adherence (Fig. 1A).

The DNA damage tracks were observed in both lymphoma (Karpas-422) as well as leukemia (SupB15) cells (Fig. 3A), demonstrating the feasibility and effectiveness of the technique across different cell types. In addition, the settings that were used in SupB15 and Karpas-422 cells also created $\gamma$H2AX damage and H3K27me3 stripes in adherent U2OS cells, except that we used 12 ROIs instead of 24 ROIs and did not use the Cell-Tak reagent for U2OS cells (Fig. 3B). Therefore, comparison of repair dynamics in suspension as well as adherent cells can also be made using our newly developed technology (Fig. 3B).

3.2. Detection of DSB proteins at micro-laser damage sites in DLBCL lines as well as in primary DLBCL patient samples

To detect proteins at laser-induced DNA breaks following immunofluorescence, imaging was done in various Z-stacks to determine...
whether a protein of interest co-localizes with γH2AX containing damage lines in different planes. Cells were imaged by Z-stacks utilizing a 60× Plan Apo objective on a Nikon A1R (or Nikon A1) confocal microscope. The software was set to perform imaging at increments of 0.5 μm to result in an average of 5–15 steps per Z-stack. In order to visualize the nucleus, the protein-of-interest and γH2AX, three channels were selected for imaging: 405 nm (blue), 488 nm (green), and 561 nm (red), respectively. Using the selected stack, the number of γH2AX lines within a focal plane were counted. Only lines for the protein-of-interest that co-localized with the γH2AX line were counted as a true DNA damage signal. Approximately, 100–300 cells with γH2AX lines per treatment were counted and the experiment was repeated at least three times in order to obtain statistically significant data. To confirm that our laser settings induced DNA double strand breaks (DSBs), we

Fig. 2. Optimization of the irradiation cycles for B-cells. A. Representative images of micro-irradiation at different scan speeds, illustrating the use of different laser intensities in Karpas-422 (EZH2 GOF DLBCL) cells. Localization of γH2AX immunofluorescence to the damage lines is shown in this fig. B. SUDHL4 cells were irradiated for increasing numbers of cycles and allowed to recover for the indicated time periods: 0–5 min, cells were fixed between 0 and 5 min post start of irradiation. That is, when irradiating for 8 cycles it takes over 3 min to irradiate all 24 lines for 8 times each and therefore the closest to no recovery possible is approximately 3–4 min post start of irradiation. Cells were then stained with anti-γH2AX antibody to evaluate the appearance and intensity of laser-induced damage.
examined DNA repair proteins that localize to DSB sites and not to single-strand DNA breaks, namely, components of the MRN complex (Mre11–Rad50–Nbs1) [19]. Our results showed an efficient localization of Mre11, Rad50 and Nbs1 to the laser-induced break sites (Fig. 4A), verifying that the micro-laser technique that we have optimized for blood-related or hematological cancers creates double-strand DNA breaks.

ATM is a key regulator of DNA damage response [20]. During DNA repair, ATM is phosphorylated at Serine-1981 in response to DSB induction, and ATM phosphorylation dissociates the inactive dimers to active monomers [21,22]. We then examined whether our optimized protocol can detect post-translational modifications of DNA repair factors as effectively as histone modifications. Given the link we established between HDAC1,2 activity and DSB repair [3,23–25], we examined whether HDAC1,2 inhibition affects phospho-ATM levels at DSB sites in EZH2 gain-of-function mutant DLBCL cells. Control DMSO-treated cells or the HDAC1,2 inhibitor (RCY957) treated cells were co-stained with antibodies that detect H3K27me3 or phospho-ATM along with γH2AX. Our data showed that HDAC1,2 inhibition decreases H3K27me3 at DSB sites in agreement with our previous results [3], and that HDAC1,2 inhibition does not affect phospho-ATM at DSB sites (Fig. 4B-C). These findings together indicate that HDAC1,2 inhibition impairs EZH2-mediated but not ATM-dependent DNA repair signaling. Collectively, our results (as shown in Fig. 4A-C) demonstrate that the micro-laser technique we optimized can successfully detect proteins and their post-translational modifications at DSBs in suspension cells.

We used the settings detailed above to successfully irradiate and detect γH2AX tracks in primary mantle cell lymphoma (Fig. 5A), primary Burkitt lymphoma mononuclear cells (Fig. 5B), and in normal human peripheral blood mononuclear cells (PBMCs) (Fig. 5C). Importantly, we found that HDAC1,2 inhibition reduced H3K27me3 in primary mantle cell lymphoma (Fig. 5D), similar to that seen in EZH2 gain-of-function mutant DLBCL cells (Fig. 4B) [3]. Therefore, regulation of H3K27me3 by HDAC1,2 appears to be a general phenomenon seen in diverse cell lines and also occurs in primary patient samples. More importantly, these results demonstrate that our micro-laser technique can be applied to monitor DNA repair dynamics directly in normal PBMCs and in primary patient suspension cancer cells.

3.3. Use of the optimized micro-laser technique to examine H3K27me3 during DNA repair in suspension DLBCL cells following small molecule inhibition of epigenetic modifiers

Next, we employed our modified micro-laser approach to determine the changes in the DNA repair–associated H3K27me3 at DSBs following treatment of suspension diffuse large B-cell lymphoma cells with small molecule inhibitors of enzymes that regulate this epigenetic mark. We chose two different DLBCL cell lines for the study: SUDHL4, a germinal
center-derived diffuse large B-cell lymphoma (GC-DLBCL) line with a gain-of-function mutation in EZH2 [3] and SUDHL2, an activated B-cell-derived diffuse large B-cell lymphoma (ABC-DLBCL) line with wild-type EZH2 and expressing a truncated mutant form of the p300 histone acetyltransferase (HAT) that catalyzes histone H3K27 acetylation [26]. We treated these lymphoma lines with small molecule inhibitors of three different epigenetic enzymes: 1) DZNep that degrades the PRC2 complex that contains the EZH2 H3K27 methyltransferase [27]; 2) GSKJ4, an inhibitor of JMJD3/UTX H3K27 demethylases [28][29,30]; and 3) RCY957, an HDAC1,2 selective inhibitor that could indirectly impact H3K27me3 via regulating H3K27 acetylation [3].

We first examined the effects of these small molecule inhibitor treatments on bulk H3K27me3 and H3K27ac levels using immunoblotting. A dose dependent increase in bulk H3K27me3 levels was observed upon

Fig. 4. Localization of DSB repair proteins and histone H3K27me3 mark to γH2AX tracks in DLBCL cells. A. Karpas-422 (EZH2 GOF DLBCL) cells were micro-irradiated as described in the main text and stained with anti-γH2AX along with anti-Mre11, Rad50 or NBS1. B. Karpas-422 DLBCL cells were micro-irradiated as described in the main text and stained with anti-γH2AX and H3K27me3. A reduction in H3K27me3 is seen following treatment of cells with HDAC1,2 inhibitor (RCY957). C. Karpas-422 DLBCL cells were micro-irradiated and stained with anti-γH2AX and phospho-ATM. No change in phospho-ATM localization was observed following treatment of Karpas-422 cells with HDAC1,2-selective inhibitor (RCY957).
treatment of SUDHL2 or SUDHL4 cells with GSKJ4 (Fig. 6A–B), confirming the inhibition of H3K27 demethylases by this compound. We previously showed that treatment of SUDHL4 cells with HDAC1,2 inhibitor RCY957 or EZH2 inhibitor DZNep does not affect global H3K27me3 levels [3]. A robust increase in H3K27ac was obtained in both SUDHL4 and SUDHL2 cells following treatment with RCY957, demonstrating its ability to inhibit the deacetylase activities of HDAC1,2 (Fig. 6C–D, compare lanes 1–2). We find that SUDHL2 cells possess H3K27ac even in the absence of a functional p300 acetyltransferase, suggesting that the homologous CBP acetyltransferase might be catalyzing H3K27ac (Fig. 6C). A subtle increase in H3K27ac was observed following treatment with DZNep (Fig. 6C–D, compare lanes 1–3), which could be ascribed to an increased access to the substrate H3K27 residue by acetyltransferases as a result of the reported DZNep-mediated disruption of the EZH2-containing PRC2 complex [27]. A decrease in bulk H3K27ac was observed following treatment with GSKJ4 (Fig. 6C–D, compare lanes 1–4), which correlates well with the increase in bulk H3K27me3 levels upon inhibiting the H3K27 demethylases.
Fig. 6. The effect of HDAC, EZH2 and UTX/JMJD3 inhibitors on H3K27ac and H3K27me3 levels in DLBCL cells. A-B. SUDHL2 or SUDHL4 cells were treated with increasing concentrations of GSKJ4 (1–5 μM) for 12 h. Immunoblotting were performed with whole cell lysates obtained from control and treated cells. Blots were probed with anti-H3K27 trimethyl (me3) or anti-H3 antibody. C-D. SUDHL2 or SUDHL4 cells were treated for 12 h with single drugs, 2 μM RCY957, 0.5 μM DZNep or 5 μM GSKJ4 or treated with a single drug for 12 h, washed with PBS and treated with the second drug for an additional 12 h. Following indicated treatments, the chromatin fractions were prepared and immunoblotting was performed. Blots were probed with either anti-H3K27 acetyl (ac) or anti-H3 antibody. SE or LE, short or long exposure, respectively. E-F. SUDHL2 or SUDHL4 cells were treated for 24 h with single drugs RCY957, GSKJ4 or DZNep. Dual treatments were performed for 12 h with the first drug followed by washing of cells and treatment with the second drug for an additional 12 h. Following micro-irradiation, cells were stained with anti-γH2AX and H3K27me3. The overlay of γH2AX and H3K27me3 signals with the hoechst stained nucleus is shown in this figure. E. The p values calculated using the two tailed t-test are as follows: SUDHL2 cell line: DMSO vs 957: 0.0098 (#1), DMSO vs GSKJ4: 0.5243 (#2), DMSO vs 957 then GSKJ4: 0.164 (#3), DMSO vs DZNep: 0.107 (#4), DMSO vs 957 then DZNep: 0.0016 (#5) and DMSO vs DZNep then RCY957: 0.003 (#6). F. SUDHL4 cell line: DMSO vs 957: 0.0148 (#1), DMSO vs GSKJ4: 0.842 (#2), DMSO vs 957 then GSKJ4: 0.156 (#3), DMSO vs DZNep: 0.0303 (#4) and DMSO vs 957 then DZNep: 0.0212 (#5) and DMSO vs DZNep then RCY957: 0.05 (#6). The images were overlaid using the NIS elements software.
preexisting factor or protein post-translational modification. Micro-laser assays in general, wherein increased accumulation of a substrate by acetyltransferases and inhibition of HDAC1,2 activities. Pre-treatment of SUDHL2 or SUDHL4 cells with the H3K27 demethylase inhibitor GSKJ4 prior to HDAC1,2 inhibition prevented the accumulation of H3K27ac (Fig. 6C–D, lane 7), suggesting an acquisition of a ‘saturated’ methylated state at H3K27 globally that effectively blocks acetylation. Addition of GSKJ4 to cells pretreated with RCY957 also prevented the accumulation of H3K27ac (Fig. 6C–D, lane 8), showing that the effects of RCY957 are reversible. Taken together, these results show the effectiveness of the small molecules in inhibiting the enzymes involved in establishing the H3K27me3-ac epigenetic switch in DLBCL cells.

We then tested the effects of these small molecule inhibitors on cell viability and the cell cycle progression of SUDHL2 and SUDHL4 lines after different treatment times. The HDAC1,2 inhibitor (RCY957) or EZH2 inhibitor (DZNep) treatment showed no effect on cell viability or the cell cycle phases after 24 h treatment (Supplemental Fig. 2), but increased the sub-G1 or dead cell population after 72 h treatment (Supplemental Fig. 3), which agrees well with our previous study [3]. The H3K27 demethylase inhibitor GSKJ4 alone or when added before or after RCY957 caused death in SUDHL2 and SUDHL4 cells after 12 h treatment (Supplemental Fig. 2), and resulted in a robust cell death after 72 h treatment (Supplemental Fig. 3). Surprisingly, SUDHL4 cells expressing the hyperactive mutant EZH2 and containing elevated levels of H3K27me3 were more sensitive to GSKJ4 treatment than SUDHL2 cells expressing wild-type EZH2 (Supplemental Figs. 2 and 3). This result reveals for the first time that increasing the already high H3K27me3 levels by inhibiting H3K27 demethylation in the EZH2 gain-of-function mutant SUDHL4 cells can also be detrimental. Collectively, results from these cell cycle assays suggest that altering the levels of H3K27me3 and H3K27ac compromises the viability of SUDHL2 and SUDHL4 DLBCL cells.

We next used our optimized micro-laser assay to examine the effects of these small molecule inhibitors on H3K27me3 at DSB sites in the suspension DLBCL cells. For this assay, we chose treatment time points where the inhibitors showed no significant induction of cell death. Therefore, we used the 24 h treatment time-point for HDAC1,2 inhibitor (RCY957) or EZH2 inhibitor (DZNep) and the 12 h time-point for the H3K27 demethylase inhibitor (GSKJ4), (Supplemental Figs. 2 and 4). While treatment with GSKJ4 increased global H3K27me3 levels (Fig. 6A–B), an increase in H3K27me3 was not evident at the micro-laser-induced DSBs (Fig. 6E) and the difference was statistically insignificant between DMSO and GSKJ4 treatments in SUDHL2 (p = .5243) or SUDHL4 cells (p = .842). This finding indicates to a limitation of the micro-laser assays in general, wherein increased accumulation of a preexisting factor or protein post-translational modification at DSBs are not easily measurable. Interestingly, DZNep treatment caused a very modest reduction in H3K27me3 at DSB sites in SUDHL2 cells (p = .107), but showed greater reduction in SUDHL4 cells (p = .0303) (Fig. 6E–F), alluding to a likely higher sensitivity of the EZH2 hyperactive mutant-containing PRC2 complex in the latter cell line to the DZNep-mediated disassembly. While HDAC1,2 inhibition by RCY957 robustly increases global H3K27ac (Fig. 6C–D), it does not reduce global H3K27me3 [3]. However, H3K27me3 deposition at DSBs is severely reduced following RCY957 treatment in both SUDHL2 (p = .0098) and SUDHL4 (p = .0148) cells (Fig. 6E–F), suggesting that increased H3K27ac as a result of HDAC1,2 inhibition blocks the active deposition of EZH2-catalyzed H3K27me3 at DSBs during DNA repair. We were unable to reliably assess H3K27ac at DSB sites as it exhibited a pan nuclear staining pattern in immunofluorescence (data not shown). Nevertheless, our results show that the micro-laser assay can be effectively used to detect and measure factors that are actively recruited to or deposited at DSBs during DNA repair, such as, the MRN complex and phospho-ATM (Fig. 4A–C) in addition to H3K27me3. Moreover, these results also demonstrate that the micro-laser assay can provide valuable insights into the mode-of-action of small molecule inhibitors at DSBs during DNA repair.

We also used our optimized micro-laser assay to examine the effects of sequential inhibitor treatments on H3K27me3 at DSBs. Pre-treatment with RCY957 did not significantly impact the effects of the H3K27 demethylase inhibitor GSKJ4 on H3K27me3 at DSBs in either SUDHL2 (p = .164) or SUDHL4 (p = .156) cell lines (Fig. 6E–F). Because of its high cytotoxicity (Supplemental Figs. 2 and 3), we were unable to examine the effects of pretreatment with GSKJ4. However, pretreatment of SUDHL2 or SUDHL4 cells with RCY957 before DZNep or vice versa, both caused a statistically significant decrease in H3K27me3 when compared to DZNep treatment alone (p < .05) (Fig. 6E–F). Based on this observation, we tested the effects of RCY957 and DZNep combination treatments on cell viability. Our results showed a robust cell death induced by the combination or sequential treatments when compared to the single agent treatments (Supplemental Fig. 3) [3]. This result suggests that the HDAC1,2 and EZH2 inhibitors can possibly serve as a mechanism-based combination therapeutic strategy for EZH2 gain-of-function DLBCL. Importantly, these results strongly support the use of micro-laser assay to guide treatment plans in the clinic.

4. Discussion

Here we report the development and application of the micro-laser DNA repair assay that can now be used for liquid tumor cells. We report extensive optimizations for this assay that include determining the adherence strategy, the cell density, the laser power and microscope settings for successful micro-irradiation and imaging of suspension lymphoma and leukemia cells (Figs. 1–3). We have confirmed that the optimized settings and conditions do not compromise cell viability (Fig. 1D).

We recently reported that inhibition of Class I family histone deacetylases HDAC1 and HDAC2 (HDAC1,2) reduces the amount of DNA break-induced silencing in cis (DISC) during DNA repair, a phenomenon that normally affects several kilobases flanking DSBs [25]. DISC is regulated by two histone modifications belonging to the EZH2 repair pathway: H3K27me3 installed by EZH2, and the PBAF nucleosome remodeler complex controlled H2AK119 monoubiquitination (H2AK119ub1), another transcriptional repression mark that acts downstream of EZH2 during DNA repair [1,16,25]. H3K27 acetylation (ac) acts as a positive epigenetic signal during transcription [31]. Acetylation blocks methylation at H3K27 residue, and thus creating an epigenetic switch between these two signaling states. Consistent with this model, using the micro-laser technique, we have found that HDAC1,2 inhibition decreases H3K27me3 at DSBs during DNA repair and increases global levels of H3K27ac in DLBCL cells [3] (Fig. 6). Thus, using our optimized micro-laser technique, we have made a novel link between HDAC1,2 and EZH2 in DNA repair for the first time in DLBCL cells [3]. Overall, our optimized microlaser technique enables determining the recruitment of factors including chromatin modifiers or remodelers to DSB sites, and uncovering their functions in DNA repair, directly in liquid tumor cells.

DLBCL is the most common disease entity of B-cell non-Hodgkin lymphoma (NHL) in adults, and has one of the highest mortality rates [32]. DLBCL subtypes, GC and ABC, originate from germinal center B-cells or activated B-cells, respectively [33]. While >50% of DLBCL patients are cured with the available multi-agent chemo, radio- or immune-therapies, ~30–40% of patients develop a relapsed or refractory disease [34] and many respond poorly to a second line therapy using high-dose chemotherapy and autologous stem cell transplantation [35–37]. Therefore, there is a compelling need for new and effective therapeutic strategies and prognostic biomarkers. Sequencing of lymph node biopsies from DLBCL patients has identified recurrent...
somatic mutations in the polycomb group proto-oncogene EZH2 [38], occurring in ~22% of GC-DLBCL subtype. These are gain-of-function (GOF) mutations within the epigenetic modifier EZH2's catalytic SET domain resulting in a hyperactive enzyme producing high levels of H3K27 trimethylation (K27me3) [8,39]. EZH2 GOF GC-DLBCL cells with aberrantly elevated H3K27me3 are chemoresistant to DNA damaging agents such as doxorubicin compared to GC-DLBCL with wild-type EZH2 [40].

Small molecule inhibitors of EZH2 have reached clinical trials for these refractory EZH2 GOF B-cell lymphomas. In addition to EZH2 inhibitors, we have shown a novel strategy of targeting the EZH2 gain-of-function mutant DLBCL using HDAC1,2 selective inhibitors [3]. We reported that HDAC1,2 inhibitor decreases the survival advantages promoted by the EZH2 GOF mutation in DLBCL cells. In addition to EZH2, H3K27 demethylase JMJD3 or KDM6B is overexpressed in germinal center-derived DLBCL and this high expression of JMJD3 is also correlated with poor prognosis [41]. Here, we have now shown that GSKJ4, an inhibitor of H3K27me3 demethylase, causes rapid and robust apoptosis in GC-DLBCL cells (Supplementary Figs. 2 and 3). Interestingly, the EZH2 gain-of-function mutant DLBCL cells with high H3K27me3 are more sensitive to GSKJ4 treatment than DLBCL cells with wild-type EZH2 (Supplemental Figs. 2 and 3). This finding along with those obtained using EZH2 and HDAC1,2 inhibitors together allude to a model where maintaining the H3K27me3-H3K27ac balance is critical for DNA repair capabilities and survival of certain subset of cancer cells. The use of laser-based repair assay optimized here for hematological liquid cancers fills a major void that currently exist between assessing global histone modifications and chromatin dynamics that occur globally and those that occur specifically at DSB sites in liquid tumors. Because the suspension cells remain attached to the Cell-Tak-coated culture dish for a limited time (4–8 h), one limitation of the assay is the inability to measure γH2AX dephosphorylation and attenuation of DNA damage signal, which take several hours to be completed (Supplemental Fig. 1).

The era of using the same treatment plan for all DLBCLs is fading away as personalized medicine approaches gain ground. As illustrated by our data, the technique presented here could lend itself to important clinical applications as we demonstrate the applicability of this technique in primary lymphoma patient sample cultures. We performed micro-irradiation in two different primary patient lymphoma samples and were able to detect γH2AX and/or H3K27me3 at the DSB laser stripes (Fig. 5A–B). We could also observe a decrease in H3K27me3 at the DSB laser tracks following HDAC1,2 inhibitor treatment in a primary DLBCL patient sample (Fig. 5D). Thus, our data demonstrate an important and direct applicability of our optimized micro-laser technique to predict the response of a blood cancer patient to chemotherapy and guide the manipulation of the drug dosage or regimen for subsequent treatments using bone marrow aspirates. The optimized micro-irradiation followed by immunofluorescence described here can be used to assess the DNA repair efficiency in patients recruited to clinical trials, as the standard immunofluorescence technique to examine epitigenic histone marks or DNA damage markers would not inform us about the rapid DNA repair events that occur at DSB sites. In summary, the method described here is not only useful to decipher the mechanisms of action of novel drugs, but can also provide a valuable diagnostic tool that could be used to devise mechanism-based therapeutic strategies.

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Conflict of interest

The authors have no competing financial interests.

Author contributions

DJ and GSSB optimized the laser conditions for hematological cells. DJ, GSSB and SB designed experiments and critically evaluated the data. DJ, GSSB and KC performed the experiments described in this study. DJ and SB wrote the manuscript. SB obtained funding for this work.

Appendix A. Supplementary data

Supplementary data to this article can be found online at https://doi.org/10.1016/j.ebiom.2019.03.083.

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