IGH/MYC Translocation Associates with BRCA2 Deficiency and Synthetic Lethality to PARP1 Inhibitors

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Abstract

Burkitt lymphoma/leukemia cells carry t(8;14)(q24;q32) chromosomal translocation encoding IGH/MYC, which results in the constitutive expression of the MYC oncogene. Here, it is demonstrated that untreated and cytarabine (AraC)-treated IGH/MYC–positive Burkitt lymphoma cells accumulate a high number of potentially lethal DNA double-strand breaks (DSB) and display low levels of the BRCA2 tumor suppressor protein, which is a key element of homologous recombination (HR)-mediated DSB repair. BRCA2 deficiency in IGH/MYC–positive cells was associated with diminished HR activity and hypersensitivity to PARP1 inhibitors (olaparib, talazoparib) used alone or in combination with other chemotherapies.

Implications: This study postulates that IGH/MYC–induced BRCA2 deficiency may predispose Burkitt lymphoma cells to synthetic lethality triggered by PARP1 inhibitors.

Visual Overview: http://mcr.aacrjournals.org/content/molcancerres/15/8/967/F1.large.jpg. Mol Cancer Res; 15(8); 967–72. ©2017 AACR.

Introduction

Burkitt lymphoma/leukemia is a highly aggressive mature B-cell neoplasm characterized by chromosomal rearrangements of the c-myc oncogene resulting in the overexpression of MYC transcription factor (1). The most common translocation is the t(8;14)(q24;q32) (85% of all cases), which involves MYC and IGH loci to generate IGH/MYC. Deregulation of MYC, a potent proto-oncogene and transcriptional regulator, contributes to lymphomagenesis through alterations in cell-cycle regulation, cell differentiation, apoptosis, adhesion, and metabolism.

Burkitt lymphoma treatment consists of high-intensity chemotherapy protocols that include cyclophosphamide, cytarabine (AraC), and doxorubicin. Current therapies have achieved a very favorable outcome resulting in complete remission in 75% to 90% of Burkitt lymphoma patients and a survival rate of 70% to 80%. However, the current treatment for Burkitt lymphoma is suboptimal in elderly patients or patients with advanced stage disease, in the setting of HIV infections, as well as in the setting of relapsed disease. Therefore, new therapeutic strategies are necessary to improve the outcomes in Burkitt lymphoma diseases in the poor prognosis patients.

It has been reported that overexpression of MYC caused accumulation of potentially lethal DNA double-strand breaks (DSB; ref. 2). DSBs are usually repaired by BRCA1/2–mediated homologous recombination (HR) and DNA-PK–mediated non-homologous end joining (NHEJ; D-NHEJ), whereas PARP1-dependent NHEJ serves as a backup (B-NHEJ) pathway (3). In addition, PARP1 may prevent accumulation of potentially lethal DSBs, either by stimulation of base excision repair and single-strand break repair and/or by promoting stalled replication fork restart (4, 5).

Here, we show that IGH/MYC–positive cells display down-regulation of BRCA2 tumor suppressor protein and are hypersensitive to synthetic lethality triggered by PARP1 inhibitors, either used alone, or in combination with a cytotoxic agent.
Materials and Methods

Cells

Epstein–Barr virus (EBV)–immortalized lymphoblastoid cell lines (LCL) 1–3 from healthy donors (6), and Burkitt lymphoma–derived EBV-positive B-cell lines Mutu (generated in Dr. Paul Lieberman's laboratory, Wistar Institute, Philadelphia, PA) and Raji (ATCC-CCL-86), and EBV-negative DG75 cell line (ATCC-CRL-2625) were cultured in RPMI supplemented with 10% FBS and 5% antibiotic/antimycotic solution. All cell lines were tested for the presence or absence of EBV by Western blot analysis for expression of viral antigens and DNA quantitative PCR for viral genome. Primary Burkitt lymphoma bone marrow samples were obtained from the University of Texas MD Anderson Cancer Center [Houston, TX; karyotype: 47, XX, t(8;14)(q24;q32), der(14), t(1;14)(q11;p12)] and Perelman Stem Cell and Xenograft Core at the University of Pennsylvania [Philadelphia, PA; karyotype: 46,XX, t(8;14)(q24;q32)] and were cultured in StemSpan SFEM growth medium (Stemcell Technologies) supplemented with 15% FBS, 5% antibiotic/antimycotic solution, 20 ng/mL IL3, 100 ng/mL FLT3, and 10% antibiotic/antimycotic solution, 20 ng/mL IL3, 100 ng/mL FLT3, 50 ng/mL SCF, and 10 ng/mL IL7 (all from Cell Sciences, Inc.). These studies were approved by the appropriate Institutional Review Boards and met all requirements of the Declaration of Helsinki.

Flow cytometry

The cells were fixed and permeabilized using PerFix EXPOSE Kit (Beckman Coulter), followed by staining with specific antibodies: mouse monoclonal IgG2b anti-BRCA1 (#MAB22101, R&D Systems), mouse monoclonal IgG1 anti-BRCA2 (#MAB2476, R&D Systems), and rabbit polyclonal anti-PALB2 (#A301-246, Millipore) conjugated with fluorochromes (Alexa Fluor-405, Alexa Fluor-488 or allophycocyanin) using Zenon Labeling Kit (Life Technologies). Isotype control antibodies (Abcam) were used as controls. Flow cytometry analysis was performed using the FACSAria. Western blot analysis

Cells were lysed at 95°C in buffer containing 50 mmol/L Tris-HCl pH 6.8, 10% glycerol, and 2% SDS and analyzed by Western blot analysis using the antibodies detecting BRCA1 (#MAB22101, R&D Systems), BRCA2 (#MAB2476, R&D Systems), RAD51 (#sc6862, Santa Cruz Biotechnology), Ku70 (#A302-623A, Millipore), Ku80 (#MA5-15873, Thermo Fisher Scientific), PARP1 (#sc7150, Santa Cruz Biotechnology), and RAD52 (#sc8350, Santa Cruz Biotechnology). Secondary goat anti-mouse and donkey anti-rabbit antibodies conjugated with IRDye 800CW and IRDye 680RD were from Li-Cor Biosciences. Fluorescent signal from membranes was collected and analyzed using Li-Cor Biosciences equipment and software.

HR activity

HR activity was detected as described before with modifications (7). Five million cells were nucleofected with 5 μg of I-SceI-linearized HR reporter plasmid and 2.5 μg of pDsRed plasmid using Nucleofector (Lonza; program U-008, Human CD34 Cell Nucleofector Kit). HR event restores functional GFP expression. After 72 hours, the percentage of GFP+/Red+ cells in Red+ cells was analyzed by flow cytometry to assess HR activity.

Nucleofection of DG75 cells

Five million cells in 100 μl nucleofection solution containing 1 μg of pmaxGFP plasmid and/or 3 μg of phCMV1_2XMBP-BRCA2 plasmid (kindly provided by Dr. Stephen Kowalczykowski, Department of Microbiology and Molecular Genetics and Department of Molecular and Cellular Biology, University of California, Davis, CA; ref. 8) were transfected using the SF Cell Line 4D-Nucleofector Kit (Lonza). GFP+ cells were sorted 72 hours later.

In vitro treatment

A triplicate of 1 × 104 cells and 2 × 104 primary cells were plated in 100 μl growth medium in a 96-well plate and treated with olaparib, talazoparib, and/or AraC (all from Selleckchem).

γ-H2AX nuclear foci

γ-H2AX nuclear foci were detected as described before with modifications (10). Briefly, cells were treated with 2.5 μmol/L olaparib for 24 hours, followed by cyt centrifugation, fixation, permeabilization, and blocking using the ImageIT Fluorescence Microscopy Kit (Life Technologies). Foci were detected using a rabbit mAb specific for γ-H2AX (Millipore), and a DyLight596-conjugated goat anti-rabbit secondary antibody (Life Technologies) with 100 nmol/L DAPI used as a nuclear counter stain. The images were acquired with a Nikon fluorescence microscope. Postacquisition image processing was performed using Photoshop CS6 software, with identical settings for resolution, thresholds, and scaling applied to each image within the set.

Fixable viability dye/γ-H2AX assay

Cells were untreated or treated with 10 nmol/L talazoparib, followed by staining with Fixable Viability Dye eFluor 780 (eBioscience), fixation in formaldehyde, and permeabilization with 100% methanol. Cells were stained with Alexa Fluor 647 anti-H2AX (pS139; BD Biosciences) and analyzed by flow cytometry with BD FACSCanto II (BD Biosciences).

Figure 1. IGH/MYC-positive Burkitt lymphoma cell lines display downregulation of BRCA2 protein and sensitivity to PARPi inhibitors. A, Left, representative Western blots of the expression of indicated proteins in total cell lysates obtained from Burkitt lymphoma cell lines (DG75, Mutu, and Raji) and LCLs (clones #1, #2, #3); middle, quantification of normalized BRCA2 levels to PARPi. Bars, mean percentage volume intensity ± SD. Right, immunofluorescence analysis of the expression levels of BRCA2. Overlay of representative histograms are shown. Mean values of BRCA2 immunofluorescence signal ± SD are presented in the top right corner on each graph for three control (in red) and three lymphoma (in green) cell lines. B, HR activity in LCL#1 and Mutu cells was measured by restoration of GFP expression from HR reporter cassette. Results, mean ± SD from triplicates/sample; *, P < 0.001. C, Left, cells were left untreated or treated with 2.5 μmol/L olaparib for 48 hours followed by detection of γ-H2AX fluorescence intensity; right, representative γ-H2AX foci are shown. D, Cells were untreated (0) or treated with 5 μmol/L olaparib for 24 and 48 hours followed by staining with anti-γ-H2AX antibody and cell viability dye. Representative diagrams illustrate accumulation of γ-H2AX (right quadrangles) and dead cells (upper quadrangles). E, Burkitt lymphoma cell lines DG75, Raji and Mutu, and LCL clones 1–3 were treated with the indicated concentrations of olaparib or talazoparib, and cells were counted in Trypan blue after 72 hours. F, DG75 GFP+ and DG75 BRCA2”GFP+” cells were treated with olaparib and counted in Trypan blue after 72 hours.
Primary leukemia xenograft
NSG female mice (The Jackson Laboratory) were injected intravenously with \(1 \times 10^6\) Burkitt lymphoma patient cells. One week later, mice were randomized into four groups: control (drug vehicles), AraC (50 mg/kg i.v. days 1–5; ref. 10), talazoparib (0.33 mg/kg/day by oral gavage for 7 days; ref. 11), and AraC + talazoparib. Human leukemia cells were analyzed by flow cytometry using anti-human CD45 antibody as described before (12). Median survival time was determined. The experiments were executed in compliance with institutional guidelines and regulations.

Statistical analyses
Data are presented as mean ± SD from three independent experiments and were compared using the unpaired two-tailed Student t test; \(P\) values less than 0.05 were considered significant. Median survival time of the mice ± SE was calculated by Kaplan–Meier log-rank survival analysis. Synergistic activity was determined by two-sample t test, two-way ANOVA, or log-rank test.

Results
IGH/MYC-positive cells display low levels of BRCA2 protein, reduced HR activity, and sensitivity to PARP1 inhibitors
Western blot analysis and flow cytometry detected low levels of BRCA2 protein in three Burkitt lymphoma–derived cell lines carrying IGH/MYC in comparison with three EBV-transformed lymphoblastoid cell lines (Fig. 1A). Expression levels of the other key proteins in HR (BRCA1, PALB2, RAD51, RAD52) and also those in NHEJ (Ku70, Ku80, PARP1) were not altered. BRCA2 is a key mediator in HR, and its downregulation impairs DSB repair (13). In concordance, low expression of BRCA2 protein in MuT cells was associated with reduced HR activity (Fig. 1B). Impaired HR was accompanied by accumulation of higher numbers of DSBs marked by γ-H2AX nuclear foci in BRCA2-deficient IGH/MYC-positive cell lines in comparison with BRCA2-proficient LCL cells; this effect was even more pronounced after treatment with a PARP1 inhibitor olaparib (Fig. 1C). Upregulation of γ-H2AX in IGH/MYC-positive cell line was associated with increased cell death detected by the staining with a cell viability dye (Fig. 1D). As BRCA2 deficiency is associated with synthetic lethality triggered by PARP1 inhibitors (14), we tested the sensitivity of IGH/MYC-positive cell lines to two structurally different PARP1 inhibitors, olaparib and talazoparib. Although olaparib (Lynparza) has been recently FDA approved to treat advanced ovarian cancer associated with defective BRCA genes, talazoparib is a highly potent PARP1 inhibitor in a phase III study for the treatment of breast cancer with BRCA1 or BRCA2 mutations.

Burkitt lymphoma cell lines were much more sensitive to AraC, talazoparib, and the combination of these drugs. When used individually, AraC and talazoparib reduced the number of hCD45+ Burkitt lymphoma cells in peripheral blood, but the combination of AraC + talazoparib exerted synergistic effect (Fig. 2B). Untreated mice succumbed to the leukemia/lymphoma in 29.1 ± 0.7 days (Fig. 2C), whereas those treated with AraC or talazoparib developed deadly disease after 49.5 ± 0.9 and 42.4 ± 0.2 days, respectively (\(P < 0.01\) when compared with untreated animals). Of note, combination of AraC and talazoparib exerted synergistic therapeutic effect as the Burkitt lymphoma–bearing mice treated with the combination survived 75.5 ± 4.2 days (\(P < 0.01\) in comparison with individual drug treatments). All mice succumbed to Burkitt lymphoma–like disease as confirmed by flow cytometry detecting abundant presence of hCD45+ Burkitt lymphoma cells in peripheral blood.

Discussion
Cancer-specific defects in DNA repair pathways create the opportunity to employ synthetic lethality, already applied against cancer cells harboring deleterious mutations in BRCA1 and BRCA2 by using PARP1 inhibitors (14). Here, we show that, in comparison with nontransformed cells, Burkitt lymphoma cells carrying IGH/MYC translocation accumulate high numbers of DSBs, display low levels of BRCA2 tumor suppressor protein accompanied by inhibition of HR activity, and are hypersensitive to PARP1 inhibitors used either alone or in combination with AraC. In addition, we detected that IGH/MYC-positive cells were also sensitive to RAD52 inhibitor (Supplementary Fig. S3), known to induce synthetic lethality in BRCA2-deficient cells (15).

Synergistic anti–Burkitt lymphoma activity of AraC combined with PARP1i is probably due to the fact that AraC causes DNA single-strand breaks (16), which, if not repaired by PARP1, may lead to accumulation of lethal DSBs in BRCA2-deficient cells. This hypothesis is supported by the observation that AraC combined with olaparib resulted in selective accumulation of DSBs in Burkitt lymphoma cell line, which was associated with enhanced cell death.

The mechanism responsible for low levels of BRCA2 protein in IGH/MYC-positive cells does not appear to depend on differences in proliferation rates, because 26% ± 2% of EBV-transformed cells and 31% ± 4% of IGH/MYC-positive cells were in S-G2-M cell cycle phase. However, the presence of high levels of BRCA2 mRNA in Burkitt lymphoma cells suggests posttranscriptional regulation (Supplementary Fig. S4).
levels of c-MYC were associated with downregulation of BRCA2 protein (Supplementary Fig. S5; ref. 17). Moreover, it has been suggested that overexpression of MYC upregulates miR-1245, which directly suppresses BRCA2 3'-UTR and translation (17). In concordance, miR-1245 was upregulated in Burkitt lymphoma cells implicating its role in downregulation of BRCA2 (Supplementary Fig. S6).

The biological relevance of low levels of BRCA2 tumor suppressor protein in IGH/MYC-positive cells remains to be determined. High copy number of MYC has been detected in prostate cancer from BRCA2 germline mutation carriers (18), suggesting that tumor cells displaying deregulated MYC or the treatment of other malignancies associated with deregulation of MYC, including diffuse large B-cell lymphoma and ALK-positive large B-cell lymphoma.

Disclosure of Potential Conflicts of Interest
No potential conflicts of interest were disclosed.

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Grant Support
This work was supported by NIH/NCI 1R01 CA186238 (to T. Skorski).

Received December 16, 2016; revised April 3, 2017; accepted May 19, 2017; published OnlineFirst June 20, 2017.
References

Molecular Cancer Research

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doi:10.1158/1541-7786.MCR-16-0468

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http://mcr.aacrjournals.org/content/suppl/2017/06/02/1541-7786.MCR-16-0468.DC1

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