

ORIGINAL ARTICLE

EBV-miR-BHRF1-2 targets *PRDM1/Blimp1*: potential role in EBV lymphomagenesisJ Ma¹, K Nie¹, D Redmond², Y Liu¹, O Elemento², DM Knowles¹ and W Tam¹

PRDM1/Blimp1, a master regulator of B-cell terminal differentiation, has been identified as a tumor suppressor gene in aggressive lymphomas, including diffuse large B-cell lymphoma (DLBCL). It has been shown in DLBCL and Hodgkin lymphoma that *PRDM1* is downregulated by cellular microRNAs. In this study, we identify the Epstein–Barr virus (EBV) microRNA (miRNA), EBV-miR-BHRF1-2, as a viral miRNA regulator of *PRDM1*. EBV-miR-BHRF1-2 repressed luciferase reporter activity by specific interaction with the seed region within the *PRDM1* 3' untranslated region. EBV-miR-BHRF1-2 inhibition upregulated *PRDM1* protein expression in lymphoblastoid cell lines (LCL), supporting a role of miR-BHRF1-2 in *PRDM1* downregulation *in vivo*. Discordance of *PRDM1* messenger RNA and protein expressions is associated with high EBV-miR-BHRF1-2 levels in LCLs and primary post-transplant EBV-positive DLBCL. Enforced expression of *PRDM1*-induced apoptosis and cell cycle arrest in LCL cells. Inhibition of EBV-miR-BHRF1-2 negatively regulates cell cycle and decreases expression of *SCARNA20*, a small nucleolar RNA that is also downregulated by *PRDM1* overexpression. The interaction between EBV-miR-BHRF1-2 and *PRDM1* may be one of the mechanisms by which EBV-miR-BHRF1-2 promotes EBV lymphomagenesis. Our results support the potential of EBV-miR-BHRF1-2 as a therapeutic target in EBV-associated lymphoma.

Leukemia (2016) 30, 594–604; doi:10.1038/leu.2015.285

INTRODUCTION

PRDM1/Blimp1 is a DNA-binding, positive regulatory domain-containing transcription repressor with a critical role in the terminal differentiation of B cells as well as in the homeostatic maintenance of T cells.¹ Its function in regulating differentiation, activation and homeostasis also extends to other cell and tissue types. *PRDM1* is a tumor suppressor gene in B- and T-cell as well as natural killer cell lymphomas.² In these lymphomas, *PRDM1* has been shown to be inactivated by nonsense and missense point mutations, allelic deletion, transcription repression by translocated *BCL6* and promoter hypermethylation.^{3–8} *PRDM1* has also been previously proposed as a target for microRNA (miRNA) downregulation in classical Hodgkin lymphoma, diffuse large B-cell lymphoma (DLBCL), and extranodal NK/T-cell lymphoma.^{9–11} Downregulation of *PRDM1* by miRNA in these tumors likely contributes to the pathogenesis of these tumors.

Epstein–Barr virus (EBV) miRNAs were first identified by sequencing small RNA libraries generated from EBV-positive cell lines infected by B95.8 strain of EBV.^{12,13} To date, ~25 precursors and 44 mature miRNA have been identified in EBV. Three of them (EBV BHRF1-1, -2 and -3) are derived from the *BHRF1* cluster, and the remaining are encoded by the *BART* clusters.¹² Among the *BHRF1*-derived miRNAs, EBV-miR-BHRF1-1 is found in the promoter region, whereas EBV-miR-BHRF1-2 and EBV-miR-BHRF1-3 are located in the 3' untranslated region (UTR) of the *BHRF1* cluster.¹⁴ In addition, 8 *BART* precursors (EBV mir-BART1, -BART3-6 and -BART15-17) are located at the cluster I of *BART*, and 13 other miRNAs (EBV mir-BART7-14, and -BART18-22) are located at the cluster II of *BART*.¹⁵ *BHRF1* miRNA is highly expressed in stage III

latency but barely detectable in either stage I or II latency during viral life.¹³ In contrast, *BART* miRNAs are expressed in all EBV-positive cell lines including Burkitt lymphoma (BL), lymphoblastoid cell lines (LCLs) and nasopharyngeal cancer.¹⁶ There is emerging evidence that EBV miRNAs regulate expression of cellular messenger RNAs (mRNAs) with important effects in biological processes such as cell proliferation and survival.^{17–21} The interaction between EBV miRNAs and cellular mRNAs may be rather pervasive, as demonstrated by the multiple cellular targets identified globally by an immunoprecipitation of Argonaute protein-containing RNA-induced silencing complexes followed by microarray analysis (RIP-Chip) technique to look for interaction between EBV miRNAs and 3'UTR of the cellular target genes within the RNA-induced silencing complexes complex.²² These results suggest capability of EBV viral miRNAs to regulate diverse cellular pathways including p53, B-cell signaling, cell proliferation and apoptosis.¹³

Interestingly, one of the members of the *BHRF1* miRNAs, EBV-miR-BHRF1-2, was found by this technique to bind to the 3'UTR of *PRDM1*, suggesting that *PRDM1* may be a target of this EBV miRNA.²² In our studies, we provide functional evidence that *PRDM1* is indeed a cellular target for EBV-miR-BHRF1-2. We demonstrate that the inhibition of *PRDM1* expression by EBV-miR-BHRF1-2 is likely to confer growth advantage to EBV-infected B cells by dampening *PRDM1*-mediated functions. These findings are highly relevant to the pathobiology of EBV-associated lymphoproliferative disorder and provide one of the mechanisms by which EBV can promote B-cell lymphoma development through its miRNA functions.

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Received 9 June 2015; revised 4 September 2015; accepted 28 September 2015; accepted article preview online 4 November 2015; advance online publication, 18 December 2015

MATERIALS AND METHODS

Patient formalin-fixed and paraffin-embedded tissues, cell block and cell lines

Seven EBV-positive post-transplant lymphoproliferative disorders (PTLD) formalin-fixed and paraffin-embedded patient samples were obtained from the Department of Pathology and Laboratory Medicine according to the protocols approved by the Institutional Review Board of Weill Medical College of Cornell University. All PTLD cases show > 80% of tumor content, and were reviewed and classified according to the World Health Organization 2008 classification. LCLs ARH77, CCL156, CCL159 and TIB-190 were purchased from ATCC (Manassas, VA, USA), and JY25 was previously described.²³ These LCL cell lines were derived from infecting peripheral blood B-lymphocytes with EBV. All LCLs were cultured in RPMI1640 medium supplemented with 100 ng/ml penicillin/streptomycin and 20% fetal bovine serum. BL cell line MUTU-1 was purchased from ATCC, and cultured in RPMI1640 supplemented with 100 ng/ml penicillin/streptomycin and 10% fetal bovine serum. Multiple myeloma cell line U266 was also purchased from ATCC and cultured in RPMI1640 supplemented with 100 ng/ml penicillin/streptomycin and 20% fetal bovine serum. All cell lines were maintained in the humidified 37 °C/5% CO₂ incubator. Formalin-fixed, paraffin-embedded cell blocks were made according to a previously described methods.²⁴

Antibodies

See Supplementary Materials and Methods.

miRNA inhibitor and siRNA

See Supplementary Materials and Methods.

Plasmids construction

See Supplementary Materials and Methods.

Transfection

For the PRDM1 overexpression experiments, 0.5×10^6 CCL159 and JY25 cells were transfected either with 1 µg pMSCV-PRDM1 plasmid or pMSCV-PIG empty vector. For the EBV-miR-BHRF1-2 inhibition study, 0.5×10^6 CCL156 and CCL159 cells were transfected either with 1 µM EBV-miR-BHRF1-2 inhibitor or *mirvana* miRNA inhibitor negative control (Life Technologies Inc., Grand Island, NY, USA), with or without 1 µM PRDM1 small interfering RNA (siRNA). For the PRDM1 and SCARNA20 co-expression study, 0.5×10^6 of CCL156 or CCL159 cells were transfected with 1 µg of pMSCV-PRDM1 plasmid, and/or 1 µg of pcDNA3.1-SCARNA20. pMSCV-PIG and pcDNA3.1 empty vectors were included as vehicle controls. All transfections were performed in two independent experiments with an optimized transfection program #16 (for plasmid) and #15 (for siRNA and miRNA inhibitor) using Neon transfection system (Life Technologies Inc.) per manufacturer's instructions. Representative transfection efficiencies were illustrated in Supplementary Figure. 1.

Western blotting and immunohistochemistry

See Supplementary Materials and Methods.

miRNA quantitation

See Supplementary Materials and Methods.

Firefly luciferase assay

See Supplementary Materials and Methods.

Reverse transcriptase, quantitative PCR and human gene expression assay probes

Reverse transcription and quantitative PCR were carried out as previously described.⁹ See Supplementary Methods for gene expression probes details.

Cell cycle analysis and apoptosis assay

See supplementary Materials and Methods.

Whole-transcriptome sequencing (RNA-seq)

See supplementary Materials and Methods.

RNA-seq data processing

The transcriptome sequencing reads were mapped to the human reference (hg19) via TopHat and the RefSeq (March 2014) transcript levels (units of FPKM) were quantified using Cufflinks, and differential expression was tested in CuffDiff according to a standard protocol (<http://www.nature.com/nprot/journal/v7/n3/full/nprot.2012.016.html>).²⁵ Heatmaps of select transcripts were then plotted with a red-black-green scale using the R package heatmap2 (<https://cran.r-project.org/web/packages/gplots/index.html>) from the RNA-seq expression data.

Total and miRNA extraction from formalin-fixed paraffin-embedded samples

See supplementary Materials and Methods.

Statistical analysis

Unpaired Student *t*-test was used to compare the difference between two groups, and a one way-analysis of variance test was performed on assays involving at least three groups for comparison. Correlation between the EBV-miR-BHRF1-2 and PRDM1 mRNA expression in EBV-positive PTLD samples were performed using linear regression analysis.

RESULTS

EBV-miR-BHRF1-2 targets the 3'UTR of PRDM1 mRNA

Argonaute-based RIP-CHIP assay has identified with high confidence a putative miRNA interaction site for EBV-miR-BHRF1-2 in a short RNA fragment corresponding to 1578–1601 nt of the 3'UTR of PRDM1.²² This short region harbors a 7mer site located from 1581 to 1588 nt of PRDM1 3'UTR that matches the seed

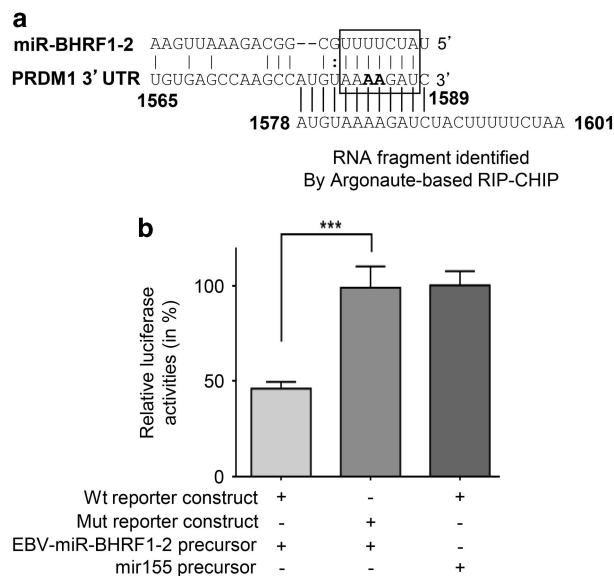


Figure 1. EBV-miR-BHRF1-2 targets PRDM1 3'UTR. **(a)** Schematic representation of EBV-miR-BHRF1-2 putative binding site located from 1565 to 1589 on the 3'UTR of PRDM1. The highly conserved 7-bp seed pair is highlighted by open box. The short RNA fragment identified by Argonaute-based RIP-CHIP assay²² was also depicted below. **(b)** Either 20 nm reporter luciferase plasmids containing wild-type (Wt) sequence or mutant (harboring point mutations at the two positions in the EBV-miR-BHRF1-2 binding site complementary to bases 5 and 6 of EBV-miR-BHRF1-2 (highlighted in bold) were co-transfected with the indicated miRNA precursors into 293T cells. Luciferase activities were measured 24 h post transfection, and normalized against firefly luciferase activities. Error bars represent the s.e.m. from technical replicates. ****P* < 0.005.

region (bases 2–8) of EBV-miR-BHRF1-2^(ref. 22) (Figure 1a). As a first step to validate that *PRDM1* is indeed a functional target of EBV-miR-BHRF1-2 mediated through this miR binding site, we used luciferase reporters containing either the wild-type *PRDM1* 3' UTR or the *PRDM1* 3'-UTR harboring a point mutation in the critical seed region of the EBV-miR-BHRF1-2 binding site (Figure 1a). Wild-type or mutant reporter plasmid was then co-transfected with 20 nM EBV-miR-BHRF1-2 precursor molecules into 293T cells, and their luciferase activities were measured at 24 h. EBV-miR-BHRF1-2 repressed wild-type *PRDM1* luciferase reporter activity by $46.2 \pm 3.5\%$ (Figure 1b, $P < 0.005$) but had no effect on the luciferase activity of the mutant *PRDM1* plasmid. MiR-155, for which no putative binding sites were predicted within the *PRDM1* 3'UTR, did not alter the *PRDM1* luciferase activity (Figure 1b). These results indicate that EBV-miR-BHRF1-2 can repress expression of *PRDM1* by direct and specific interaction with the *PRDM1* 3'UTR.

EBV-miR-BHRF1-2 downregulates endogenous *PRDM1* protein expression in EBV-immortalized lymphoblastoid cells

Many EBV-positive B-cell lymphomas are postulated to originate from EBV-infected B cells with latency III program of EBV gene expression. Thus, EBV-immortalized LCLs, which are of latency III type, serve as a good model to study EBV lymphomagenesis.²⁶ To further investigate that *PRDM1* is an *in vivo* target for EBV-miR-BHRF1-2, EBV-immortalized LCLs were transfected either with EBV-miR-BHRF1-2 inhibitor or miRNA inhibitor negative control. Although CCL159 and CCL156 cells transfected with EBV-miR-BHRF1-2 showed no significant changes in *PRDM1* mRNA levels compared with controls (Figure 2a), *PRDM1* protein expression was induced by 40% on EBV-miR-BHRF1-2 inhibitor transfection in

CCL159 and CCL156 cells at 48 h post transfection (Figures 2b and c). These results demonstrate that *PRDM1* is a target for translational repression by EBV-miR-BHRF1-2 in LCL cells.

Discordance of *PRDM1* mRNA and protein expressions in LCLs is associated with high EBV-miR-BHRF1-2 levels

If EBV-miR-BHRF1-2 downregulates *PRDM1* expression in LCL cells predominantly by a translational repression mechanism, one should expect to see an association of high EBV-miR-BHRF1-2 expression with low *PRDM1* protein expression in LCLs. We analyzed the total *PRDM1* mRNA expression by quantitative transcription PCR in five LCLs cell lines. CCL156 expressed the highest level of *PRDM1* mRNA. The *PRDM1* transcript levels in ARH77 and TIB-190 were also higher than that of the myeloma cell line U266. CCL159 and JY25 showed relatively lower levels of *PRDM1* mRNA (Figure 3a). *PRDM1* protein expression in these cell lines was also determined by immunohistochemistry performed on cell blocks and western blotting. All LCL cell lines examined showed much lower levels of *PRDM1* protein compared with U266 (Figures 3b–d). Quantification of protein/mRNA ratio of each cell line indicated that the LCL cells have a lower protein/mRNA ratio compared with U266, suggestive of lower translation efficiency in LCLs (Figure 3e). Interestingly, all five LCL cell lines had markedly higher levels of EBV-miR-BHRF1-2 relative to U266, a multiple myeloma cell line, which lacks EBV and showed undetectable level of EBV-miR-BHRF1-2 (Figure 3f). These results are compatible with an inhibitory effect of EBV-miR-BHRF1-2 on *PRDM1* expression in LCL cells. MUTU-1, a BL cell line of type I EBV latency, showed undetectable level of EBV-miR-BHRF1-2.

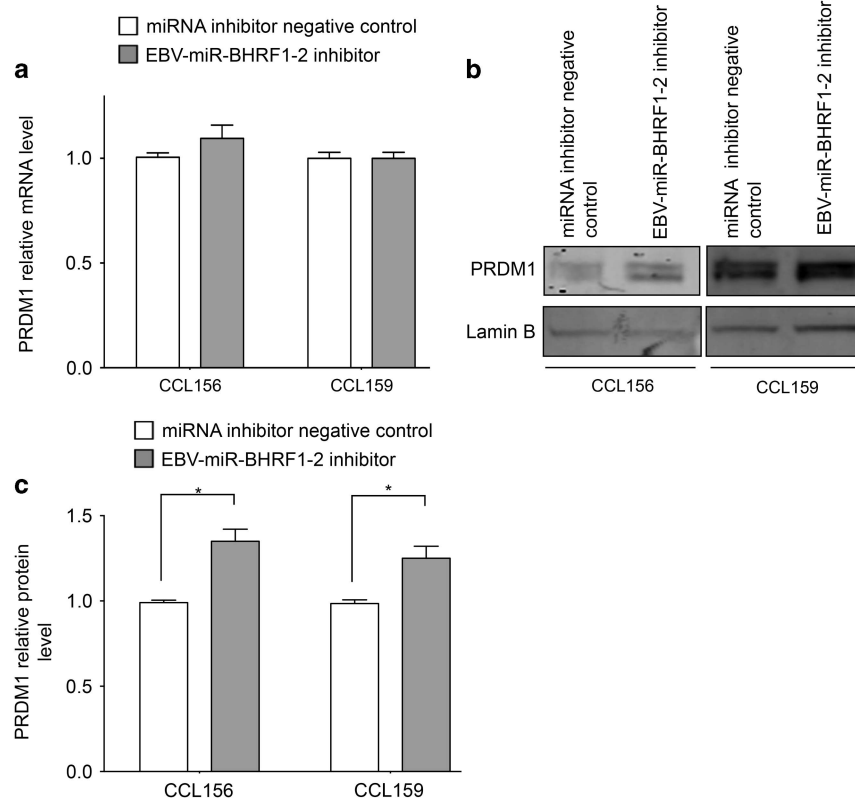


Figure 2. EBV-miR-BHRF1-2 downregulates *PRDM1* protein levels in LCL cells. (a) CCL159 or CCL156 cells were transfected either with $1 \mu\text{M}$ of EBV-miR-BHRF1-2 inhibitor, or with $1 \mu\text{M}$ *mirvana* miRNA inhibitor negative control. *PRDM1* mRNA level was measured 48 h post transfection, and normalized to negative control. (b) Whole cell lysates were prepared from the above indicated transfectants at 48 h. *PRDM1* protein expression was determined by western blotting. Lamin B was included as a loading control. Note: The doublet appears to be artifactual and was not seen in the immunoblots in Figures 3c and 6b. (c) Normalized *PRDM1* protein was expressed as a percentage relative to the negative control, and graph was plotted using the prism 6 software (La Jolla, CA, USA). Error bars represent the s.e.m. from technical replicates. * $P < 0.05$.

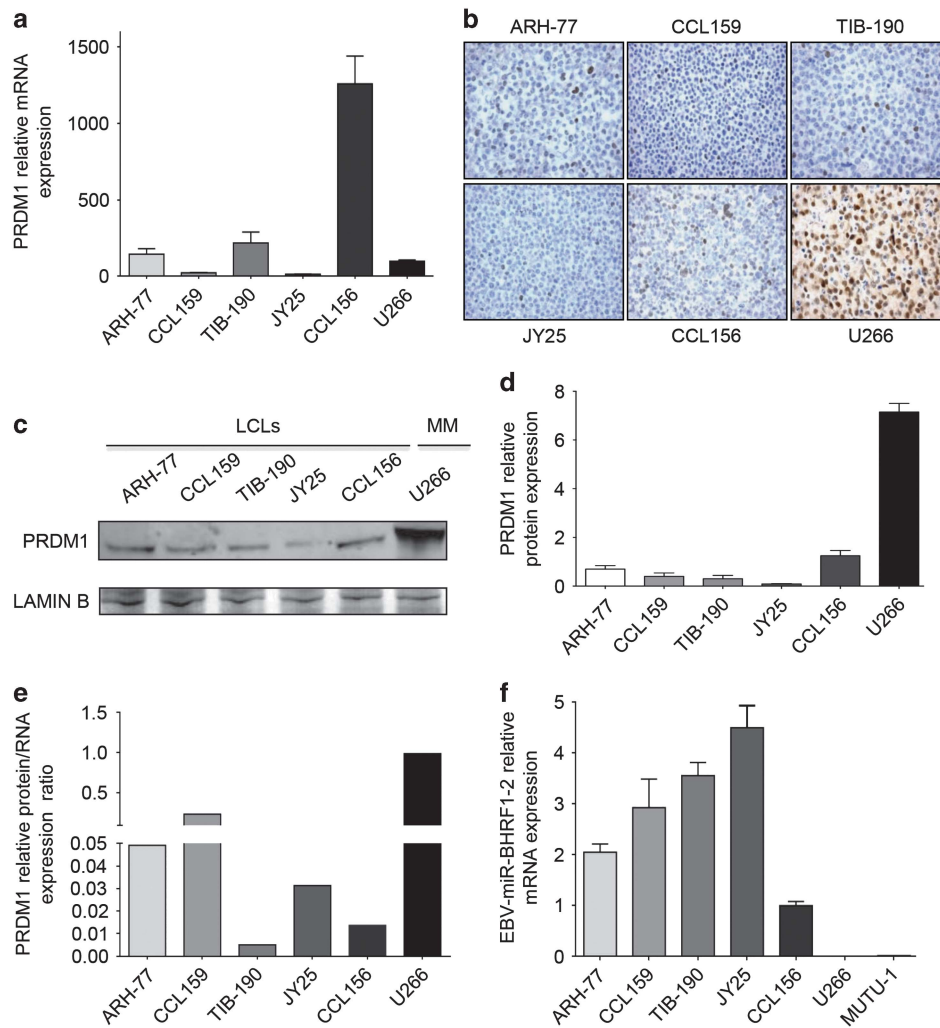


Figure 3. High levels of EBV-miR-BHRF1-2 are associated with discordance of *PRDM1* mRNA and protein expressions in LCLs. **(a)** qPCR analysis of *PRDM1* expression in five LCLs. The relative *PRDM1* mRNA levels of each cell line are shown (U266 as 100). **(b)** Immunoperoxidase staining of *PRDM1* protein on formalin-fixed, paraffin-embedded cell blocks. **(c)** Immunoblotting analysis of whole cell lysates extracted from the indicated cell lines using the antibody against *PRDM1*. Nuclear protein Lamin B was used as loading control. **(d)** Immunoblotting band intensities were measured using the Image J software (National Institutes of Health, Bethesda, MD, USA), and bar graph was plotted using GraphPad prism 6 (National Institutes of Health) to demonstrate relative *PRDM1* expressions among cell lines. **(e)** Quantification of protein/mRNA ratio in five LCLs cell lines and a multiple myeloma cell line U266. **(f)** Real-time PCR analysis of EBV-miR-BHRF1-2 expression in five LCLs cell lines, one multiple myeloma cell line (U266), and one Burkitt lymphoma cell line (MUTU-1). The relative mRNA levels are shown (The value in CCL156 set as 1). Error bars represent the s.e.m. from two independent experiments. qPCR, quantitative PCR.

High expression of EBV-miR-BHRF1-2 correlates with low *PRDM1* expression in EBV-positive lymphomas

To further investigate whether there is also a negative correlation between EBV-miR-BHRF1-2 levels and *PRDM1* expression in primary EBV-positive lymphoma cases, EBV-miR-BHRF1-2 and *PRDM1* expressions were assessed in seven cases of EBV-positive PTLD (Supplementary Table 1), all of which harbored tumor content of > 80%. Cases 1 A, 3 and 5 showed relatively high levels of miR-BHRF1-2 and *PRDM1* mRNA (Figure 4a). In contrast, the levels of EBV-miR-BHRF1-2 and *PRDM1* mRNA were detected at very low levels in the rest of the cases. Overall, the expression of EBV-miR-BHRF1-2 is positively associated with *PRDM1* mRNA expression in these patients (Figure 4b, $P < 0.0001$), suggesting both are activated during EBV type III latency. This positive association between EBV-miR-BHRF1-2 and *PRDM1* mRNA expressions is well demonstrated in patient 1, whose samples 1A and 1B represent the initial diagnostic and relapse samples, respectively.

Interestingly, while sample 1A had high EBV-miR-BHRF1-2 and *PRDM1* mRNA, very low EBV-miR-BHRF1-2 and *PRDM1* mRNA were detected in sample 1B. The mechanism underlying these alterations between the diagnostic and relapse samples is not known. Two of those three cases exhibiting increased levels of *PRDM1* mRNA have discordantly low levels of *PRDM1* protein as demonstrated by immunohistochemistry. An example was shown in Figure 4c. In the third case (Pt. 3), higher levels of *PRDM1* were seen overall compared with other cases (Figure 4d). However, on closer inspection, it appears that there are variations in *PRDM1* expression between areas within the tumor (Figure 4d). We performed EBNA2/*PRDM1* double immunostaining in this case (Figure 4e), and observed a negative correlation between EBNA2 and *PRDM1* expressions (Figure 4d). As EBNA2 is associated with type III latency²⁷ and EBV-miR-BHRF1-2 is expressed during type III latency,¹³ this finding indicates a negative correlation between EBV-miR-BHRF1-2 and *PRDM1* expression between different tumor cell subpopulations.

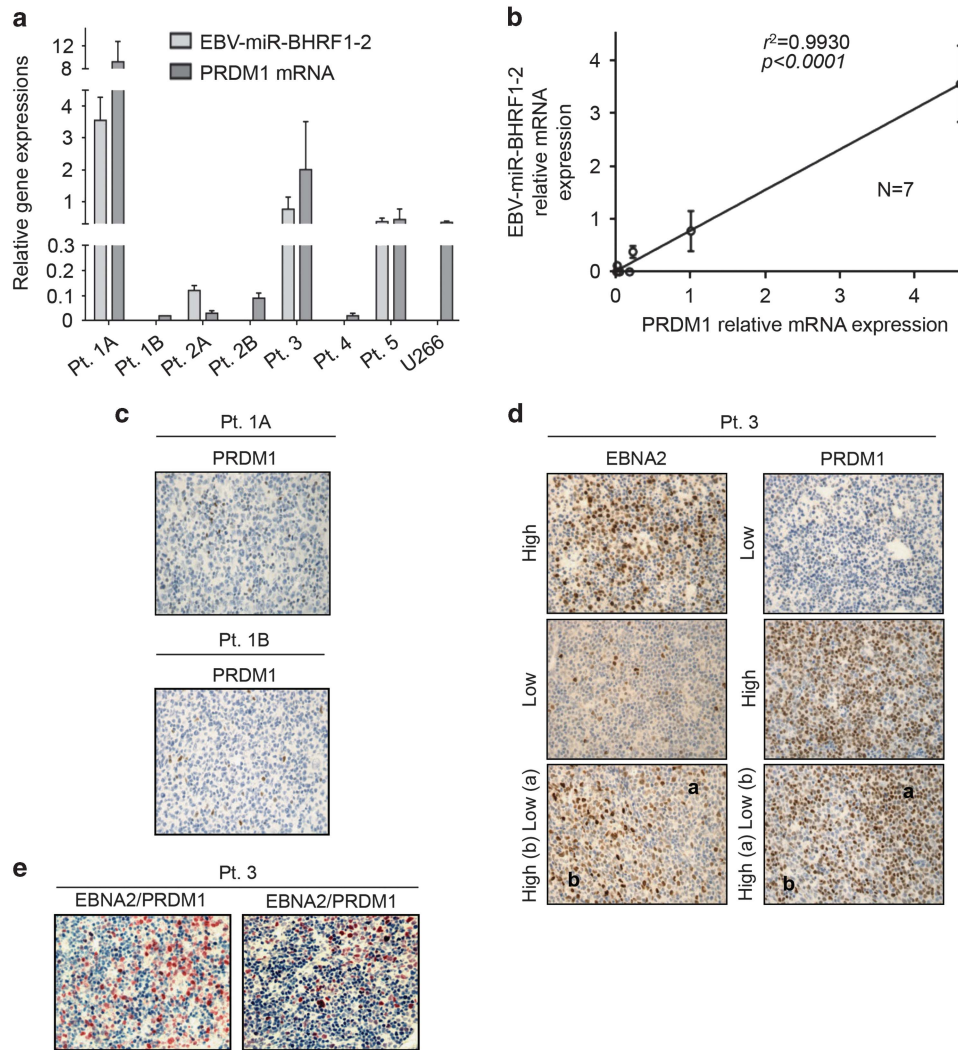


Figure 4. EBV-miR-BHRF1-2 and PRDM1 expressions in EBV⁺ PTLD patients. Total RNA and miRNA was extracted from seven formalin-fixed and paraffin-embedded patient samples and 1 cell block (U266). EBV-miR-BHRF1-2 and PRDM1 mRNA expressions were determined by qRT-PCR, and PRDM1 protein expression was evaluated using immunohistochemistry. (a) EBV-miR-BHRF1-2 and PRDM1 mRNA expressions in individual cases. U266 cell block was included as a positive control. (b) Linear regression of EBV-miR-BHRF1-2 and PRDM1 mRNA expression in all cases ($N=7$). (c) Immunostaining of PRDM1 in Pt. 1A, which has high PRDM1 mRNA expression, and in Pt. 1B, which has low PRDM1 mRNA expression. Note the low PRDM1 protein expressions in both cases. (d) Immunostaining of PRDM1 and EBNA2 in Pt. 3. Note the negative correlation between PRDM1 and EBNA2 expressions in different areas. **a**, EBNA2 low, PRDM1 high; **b**, EBNA2 high, PRDM1 low. (e) PRDM1 (blue)/EBNA2 (red) double staining in Pt. 3. The vast majority of the cells show either blue (PRDM1 positive) or red (EBNA2 positive) nuclear staining, indicative of the negative relationship between PRDM1 and EBNA2 expressions. Purple staining, indicative of PRDM1 and EBNA2 co-expressions, constitutes < 10% of the total nuclei. qRT-PCR, quantitative reverse transcription PCR.

Overexpression of PRDM1 induced S phase reduction and cell death in CCL159 and JY25 cells

To determine the potential effects of EBV-miR-BHRF1-2-mediated inhibition of PRDM1 expression and how this inhibition might contribute to lymphomagenesis, we first investigated the biological consequences of increased PRDM1 on LCL cells. To accomplish this, we constructed a pMSCV-PRDM1 plasmid to overexpress PRDM1 in LCL cells (Figures 5a and b). A significant cell apoptosis was induced following overexpression of PRDM1 in both CCL159 and JY25 cells ($P < 0.01$; Figure 5c). The percentage of S phase was reduced from 43.4% (vector control) to 27.6% ($P < 0.05$, Figures 5d and e) in CCL159 cells and from 39.5% (vector control) to 27.9% ($P < 0.05$, Figures 5d and f) in JY25 cells following PRDM1 overexpression at 48 h. A concomitant increase in G₀/G₁ phase was also detected in CCL159 cells ($P < 0.05$, Figures 5d and e). Collectively, these data indicate that increased PRDM1 expression has a negative effect on LCL cell survival and

cell proliferation. Thus, counteracting an increase in PRDM1 by EBV-miR-BHRF1-2 can potentially confer advantage to lymphomagenesis by a positive effect in cell survival and proliferation.

EBV-miR-BHRF1-2 inhibitor blocks G₁ to S phase transition, which can be relieved by knockdown of endogenous PRDM1 in LCLs cells

We then explored the biological effects of inhibiting EBV-miR-BHRF1-2 on LCLs and whether any of these effects may be mediated through increase of PRDM1 levels. We either overexpressed this inhibitor alone or in combination with PRDM1 siRNA in LCLs (Figures 6a and b). Quantitative reverse transcription PCR, immunoblotting and cell cycle assays were conducted at 48 h. Although PRDM1 mRNA level remained unchanged on EBV-miR-BHRF1-2 inhibitor (plus scramble siRNA control) transfection compared with negative control (miRNA inhibitor negative control plus scramble siRNA control), western blotting revealed

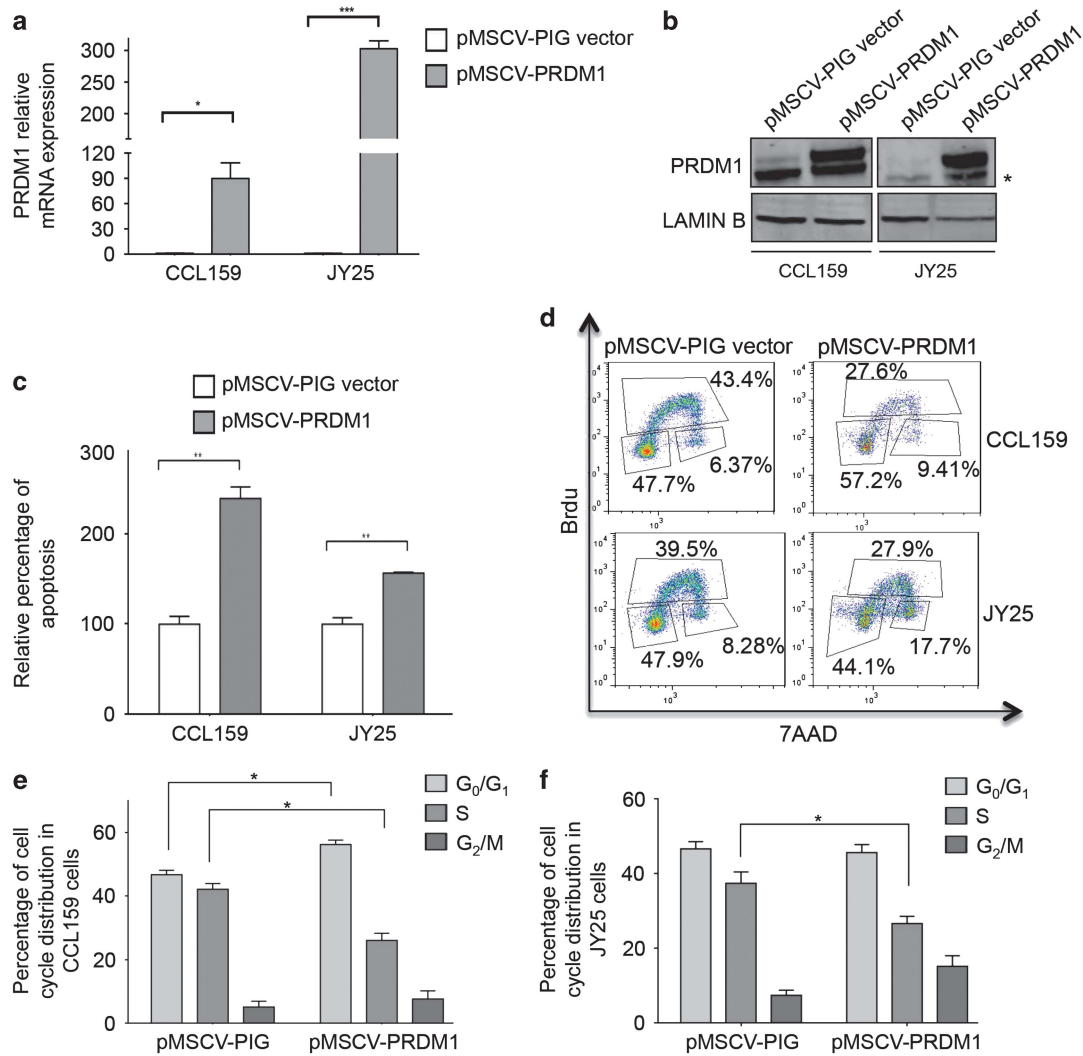


Figure 5. Overexpression of PRDM1 induces cell cycle arrest and apoptosis in LCL cells. CCL159 and JY25 cells were transfected either with pMSCV-PIG vector or pMSCV-PRDM1 plasmid. Total RNA or protein was collected at 48 h time point, *PRDM1* expression was determined by real-time PCR (a) or western blotting (b). Lamin B was included as a loading control. The lower band seen (indicated by asterisk) in (b) is most likely a non-specific band, as previously reported.⁴⁸ (c) Cells collected at 48 h were stained with Annexin V and 7AAD using apoptosis detection kit (BD Biosciences, San Jose, CA, USA) and subjected to flow cytometry analysis. (d) Cell cycle analysis of GFP positive cells at indicated time point. The percentage of each cell cycle phase distribution in CCL159 (e) and JY25 (f) cells was calculated by FlowJo software (Ashland, OR, USA) and graph was generated using the prism 6 software. Error bars represent s.e.m. from technical replicates. * $P < 0.05$; ** $P < 0.01$; *** $P < 0.005$. GFP, green fluorescent protein.

induction of *PRDM1* protein level as previously demonstrated in Figure 2b. This is associated with a decrease in cell cycle progression based on BrdU incorporation assay. As shown in Figures 6c and d, the S phase was reduced from 22.2% in CCL159 cells co-transfected with miRNA inhibitor negative control and siRNA scramble control, to 18.1% on EBV-miR-BHRF1-2 inhibitor plus scramble siRNA transfection ($P < 0.01$). To test if this inhibitory effect on cell cycle can be attributed to the increase in PRDM1, we transfected EBV-miR-BHRF1-2 inhibitor along with *PRDM1* siRNA. Transfection of *PRDM1* siRNA (plus miRNA negative control) reduced *PRDM1* mRNA and protein levels (Figures 6a and b), which resulted in a significant increase in S phase from 22.2% to 32.6% (Figure 6c, $P < 0.05$). This result is in line with our results on *PRDM1* overexpression (Figure 4) in LCL cells, further supporting a critical role of *PRDM1* in cell cycle control in LCL cells. Co-transfection of *PRDM1* siRNA essentially abrogated the PRDM1 induction seen in transfection with EBV-miR-BHRF1-2 inhibitors. More importantly, instead of a decrease in S phase as seen in EBV-miR-BHRF1-2 inhibitor transfectants, the percentage

of cells in S phase was now increased from 22.2% to 32.7%, similar to the effect of *PRDM1* siRNA alone (Figures 6c and d, $P < 0.001$). Taken together, these results suggest that EBV-miR-BHRF1-2 may promote cell cycle progression through its inhibitory effect on *PRDM1* expression.

Inhibition of EBV-miR-BHRF1-2 decreases *SCARNA20* expression in LCLs through downregulation of *PRDM1*

To explore how a modest increase in *PRDM1* on EBV-miR-BHRF1-2 inhibition may modulate *PRDM1* function, we compared the changes in gene expression on *PRDM1* overexpression and on EBV-miR-BHRF1-2 inhibition and determined if they shared any genes in common. LCLs cells were transiently transfected with either pMSCV-PRDM1 plasmid or EBV-miR-BHRF1-2 inhibitor. Total RNA was extracted from treated cells at indicated time points, and gene expression profiles were determined using transcriptome sequencing (RNA-seq). Cufflinks software (<http://cole-trapnell-lab.github.io/cufflinks/>) based on the relative abundance of these

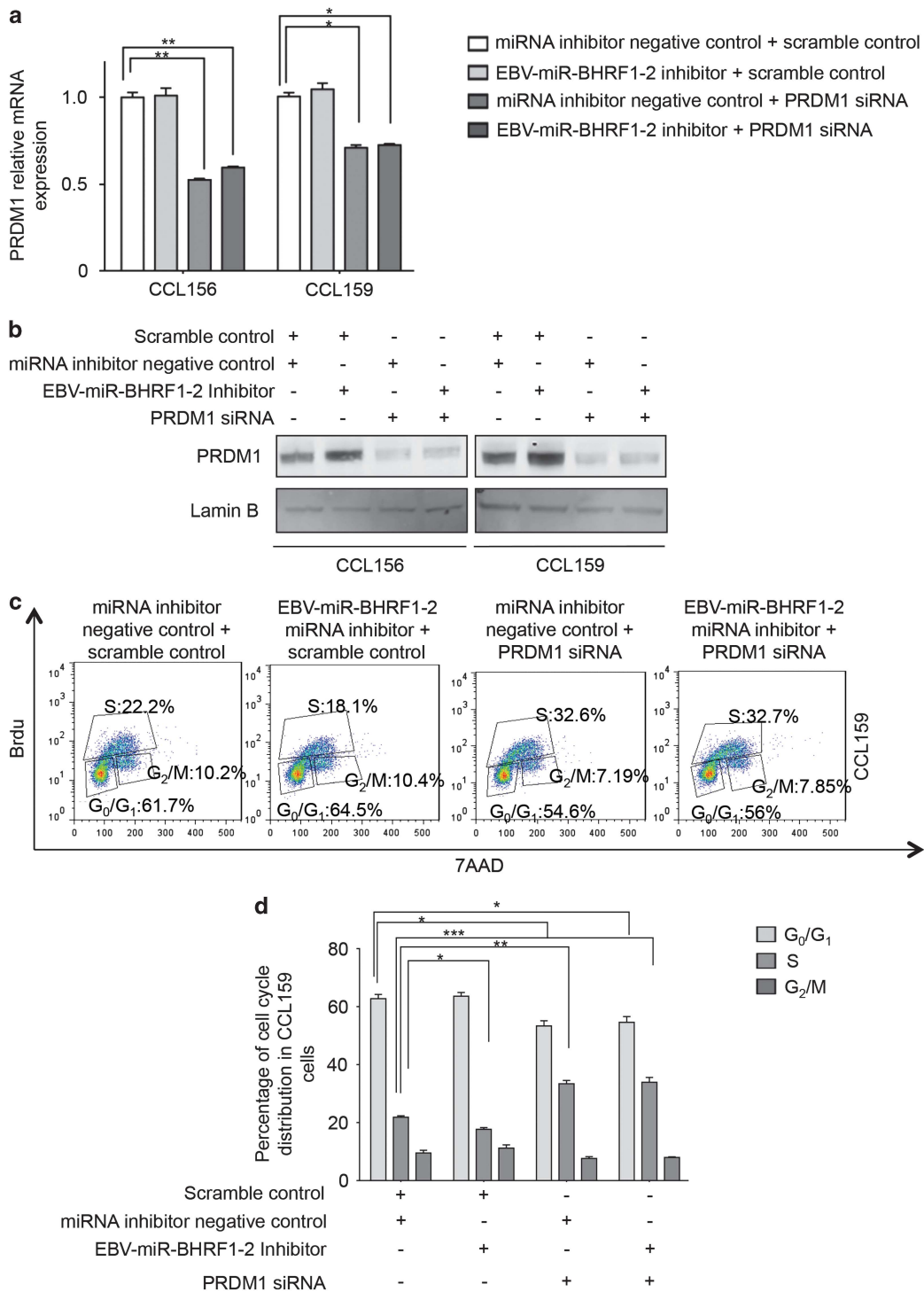


Figure 6. EBV-miR-BHRF1-2 inhibitor induces S phase arrest, which can be relieved by PRDM1 knockdown. CCL159 and CCL156 cells were co-transfected with one of the four miRNA inhibitor/siRNA combinations: miRNA inhibitor negative control plus siRNA scramble control; EBV-miR-BHRF1-2 inhibitor plus scramble control; miRNA inhibitor negative control plus PRDM1 siRNA; EBV-miR-BHRF1-2 inhibitor plus PRDM1 siRNA. Total RNA or whole cell lysates were collected at 48 h, and PRDM1 expression was determined by qRT-PCR (a) or immunoblotting (b) in LCLs transfected with the indicated miRNA/siRNA. Lamin B was included as a loading control. (c) BrdU incorporation analysis of the distribution of cell cycle in CCL159 cells transfected with the indicated transfectants. (d) Quantification of cell cycle phase distribution. Error bars represent the s.e.m. from technical replicates. **P* < 0.05; ***P* < 0.01; ****P* < 0.005. qRT-PCR, quantitative reverse transcription PCR.

transcripts analyzed the change of expression in each gene. Figure 7a demonstrated a small subset (six genes) of down-regulated genes by EBV-miR-BHRF1-2 inhibition overlapping with those by PRDM1 overexpression. Furthermore, we attempted to validate this set of six common genes by quantitative PCR. One of

these genes, SCARNA20 (ACA66) (*P* < 0.05) was consistently downregulated in PRDM1 overexpressing and EBV-miR-BHRF1-2 inhibitor-transfected CCL159 (Figure 7b) or CCL156 cells (Figure 7c). To demonstrate that the SCARNA20 downregulation can be a result of PRDM1 induction in LCL cells treated with

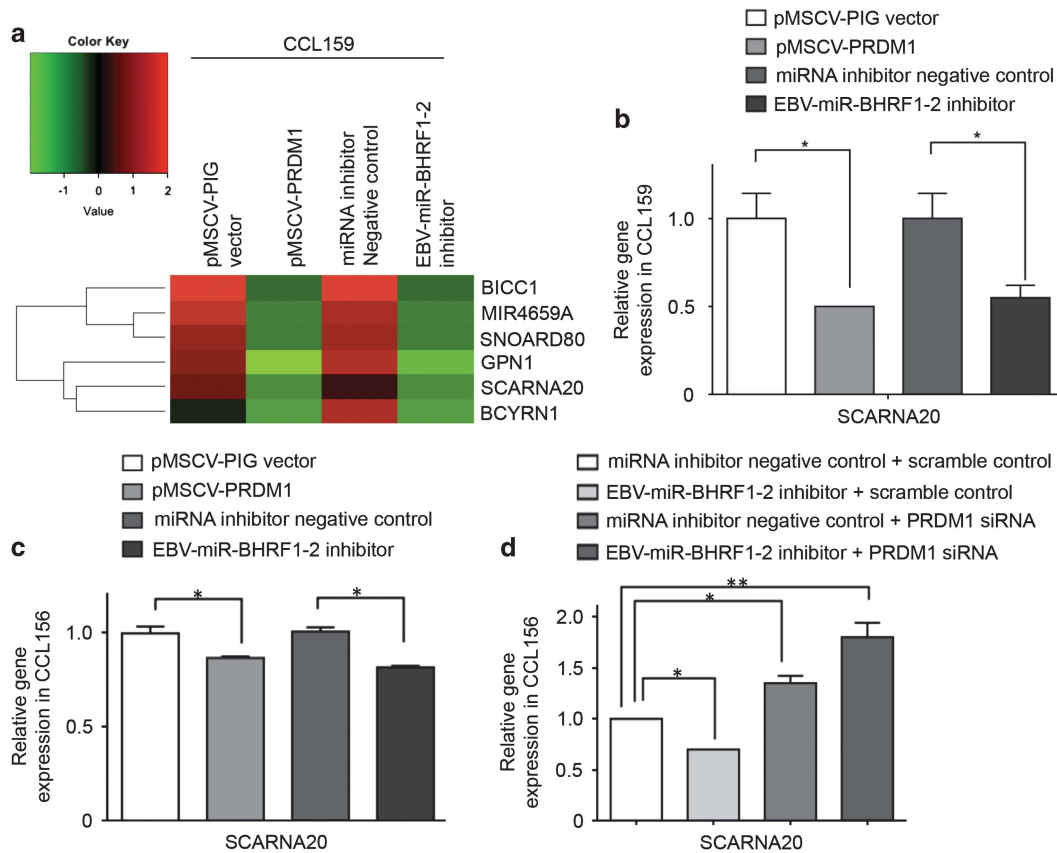


Figure 7. EBV-miR-BHRF1-2 inhibition represses *SCARNA20* via *PRDM1* downregulation. CCL159 and CCL156 cells were co-transfected with pMSCV-PIG vector, pMSCV-PRDM1 plasmid, miRNA inhibitor negative control or EBV-miR-BHRF1-2 inhibitor. Total RNA was extracted at 48 h time point, and gene expression profile was determined by RNA-seq analysis. (a) Each column indicates the representative data from two independent transfectants of CCL159. Each row represents a significantly repressed gene (green) on transfection compared with controls. A gradient color bar showed the fold decrease of gene expression as a ratio of expression in PRDM1-transfected or EBV-miR-BHRF1-2-inhibitor-transfected cells (green) versus control cells (red). qRT-PCR validation of *SCARNA20* in CCL159 cells (b) or in CCL156 cells (c) CLL156 cells were co-transfected with one of the four miRNA inhibitor/siRNA combinations: miRNA inhibitor negative control plus siRNA scramble control; EBV-miR-BHRF1-2 inhibitor plus siRNA scramble control; miRNA negative control plus PRDM1 siRNA; EBV-miR-BHRF1-2 inhibitor plus PRDM1 siRNA. Total RNA was extracted at 48 h, and *SCARNA20* expression relative to vehicle control was determined by qPCR analysis (d). Error bars represent the s.e.m. from technical replicates. * $P < 0.05$; ** $P < 0.01$. qRT-PCR, quantitative reverse transcription PCR.

EBV-miR-BHRF1-2 inhibitors, we transfected LCL cells with miR-EBV-BHRF1-2 inhibitor, *PRDM1* siRNA, or both in CCL156 cells. *SCARNA20* was reduced on overexpression of EBV-miR-BHRF1-2 inhibitor, and was induced by knocking down of *PRDM1* (Figure 7d). Abrogation of the EBV-miR-BHRF1-2 inhibitor-mediated *PRDM1* induction through simultaneous addition of *PRDM1* siRNA abolished the decrease in *SCARNA20* expression (Figure 7d). These data suggest that the negative modulatory effect of EBV-miR-BHRF1-2 on *PRDM1* expression, even though it may be possibly small, is sufficient to alter its function, leading to increased expression of a small subset of its target genes, for example *SCARNA20*.

SCARNA20 partially counteracts *PRDM1*-induced apoptosis and cell cycle arrest in LCLs

To determine whether *SCARNA20* has a role in mediating the function of *PRDM1*, we generated a *SCARNA20*-expressing plasmid (pcDNA3.1-*SCARNA20*) and transfected it along with pMSCV-PRDM1 in CCL156 and CCL159 cells. Overexpression of pcDNA3.1-*SCARNA20* significantly increased *SCARNA20* levels (Figures 8a and b). Elevated *SCARNA20* expression was capable of partially inhibiting PRDM1-mediated apoptosis and cell cycle arrest (Figures 8c–f). Cell apoptosis decreased by 15% in CCL156 cells,

and by 13% in CCL159 cells ($P < 0.05$). There was also a modest but significant increase (~10–20% increase relative to PRDM1 overexpression alone) in fraction of cells in S phase for both cell lines on *PRDM1* and *SCARNA20* co-expression. These findings provide supportive evidence for a role of *SCARNA20* in mediating *PRDM1* functions.

DISCUSSION

Our current studies are relevant to the pathogenesis of EBV-associated lymphoproliferative disorders such as PTLD, EBV-positive DLBCL and EBV-positive BL. EBV⁺ PTLD and DLBCL frequently have latency III type of EBV infection,²⁸ in which all latent genes, including LMP1, LMP2A, LMP2B, EBNA2 and EBNA3–6 are all expressed. EBV LMP1 and EBNA2 can activate NF- κ B.^{29–32} In addition, the majority of EBV-positive PTLD and DLBCL are of non-germinal center B-cell origin.^{33,34} Constitutively, activated NF- κ B pathway is a characteristic feature of non-GCB type of DLBCL.³⁵ As this pathway has the capacity to active proliferation but also induce *PRDM1* expression and, hence terminal differentiation, downregulation of *PRDM1* is critical to lymphoma development. High levels of EBV-miR-BHRF1-2 expressions in EBV-positive PTLD and DLBCL with latency III pattern will therefore be advantageous to the growth of lymphoma cells by

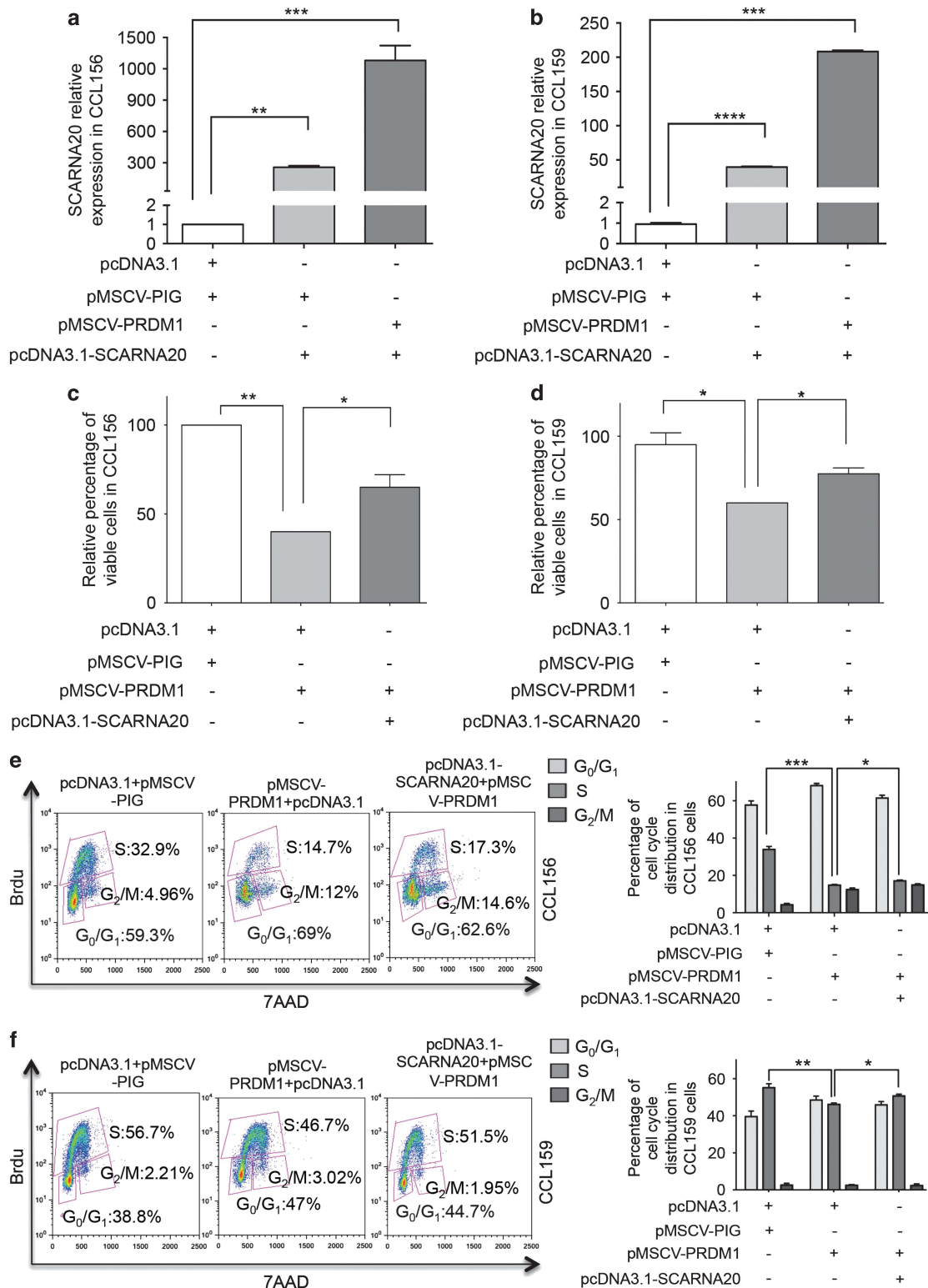


Figure 8. SCARNA20 partially inhibits PRDM1-induced apoptosis and cell cycle arrest in LCL cells. CCL156 and CCL159 cells were co-transfected with one of the following plasmid combinations: pcDNA3.1 plus pMSCV-PIG; pcDNA3.1-SCARNA20 plus pMSCV-PIG; pMSCV-PRDM1 plus pcDNA3.1; pcDNA3.1-SCARNA20 plus pMSCV-PRDM1. Total RNA, live cells or fixed cells was collected at 48 h. (a and b) SCARNA20 expression was determined in CCL156 and CCL159 cells transfected with the indicated plasmid. Apoptosis detection assay of Annexin V and 7AAD doubled stained CCL156 (c) and CCL159 (d) cells. BrdU incorporation analysis and quantification of the distribution of cell cycle in CCL156 (e) and CCL159 cells (f) transfected with the indicated plasmids. Error bars represent the s.e.m. from technical replicates. * $P < 0.05$; ** $P < 0.01$; *** $P < 0.005$; **** $P < 0.0001$.

limiting *PRDM1* expression that would otherwise be upregulated by activated NF- κ B. Interestingly, our study of a small cohort of monomorphic EBV-positive PTLD appears to show a positive correlation between *PRDM1* mRNA and EBV-miR-BHRF1-2 levels, further supporting a potential function of EBV-miR-BHRF1-2 to counteract the possible rise of *PRDM1* protein level associated with *PRDM1* mRNA induction in latency III infection. EBV-positive BL characteristically exhibits EBV latency type I, which does not have detectable EBV-miR-BHRF1-2 expression. However, it has been proposed that the precursor of EBV-positive BL is derived initially from an EBV-infected B-cell with type III latency akin to the LBL cell lines.³⁶ Thus, though BHRF1-2 is not expressed at high levels in EBV⁺ BL in the final stage, it is likely to be highly expressed and beneficial to the initial stage of EBV⁺ BL pathogenesis. Recently, it has been shown that LMP1 can downregulate *PRDM1a* transcription in GCB cells.³⁷ Consistent with this, LCLs derived from GCB cells, in contrast to the LCLs currently used in our study, expressed low levels of *PRDM1* transcripts. It is possible that the effect of LMP1 on *PRDM1* mRNA expression may vary depending on the stage of differentiation of the B cells. It will be intriguing to determine if EBV-miR-BHRF1-2 levels remain high in LCLs derived from GCB cells.

Several previous studies have described *PRDM1* as a functionally relevant target of miRNAs.^{9–11} We demonstrate for the first time that *PRDM1* is a potential miRNA target for an EBV miRNA with likely pathogenic consequences, and adds to the expanding list of cellular genes targeted and regulated by viral miRNAs. To our knowledge, this is the first validated cellular target of EBV miR-BHRF1-2. In this study, we demonstrated that even a modest increase in *PRDM1* on treatment of LCL cells with miR-BHRF1-2 inhibitor is sufficient to alter cell cycle, but it is conceivable that the negative regulatory effect of EBV-miR-BHRF1-2 on *PRDM1* expression and the consequential biological effect on cell cycle *in vivo* is higher than what we demonstrated here. We might likely have underestimated the actual extent of downregulation of *PRDM1* by EBV miR-BHRF1-2 because of the < 100% transfection and knockdown efficiency of the miR inhibitors. However, it has been proposed that miRNAs function to fine-tune gene expression³⁸ and the change in target gene expression may not be large on changes in miRNA levels. Thus, even had a complete knockdown of EBV-miR-BHRF1-2 function been achieved, it is unlikely it could have resulted in an increase in *PRDM1* expression the same extent as a directly enforced expression. More likely, the increase in *PRDM1* expression, if totally uninhibited by EBV-miR-BHRF1-2, will be expectedly somewhere in between what has been achieved using the conventional EBV-miR-BHRF1-2 inhibitors and plasmid-driven overexpression.

To further demonstrate that even a small EBV-miR-BHRF1-2 mediated change in *PRDM1* expression levels can be functionally significant, we investigate whether the modest increase in *PRDM1* on EBV-miR-BHRF1-2 inhibition can be sufficient to alter expressions of *PRDM1* target genes in LCL cells. To this end, we explored whether there is any overlap in genes repressed by *PRDM1* overexpression and EBV-miR-BHRF1-2 inhibition. Notably, *PRDM1* target genes identified in our study are distinctly different from those identified previously,³⁹ which may be attributed to the different cell lines used for investigation. The set of genes shared by both *PRDM1* overexpression and EBV-miR-BHRF1-2 inhibition potentially represent a subset of *PRDM1* target genes, whose expressions are most sensitive to repression by *PRDM1* as its level is elevated from baseline by miR-BHRF1-2 inhibition. Indeed, target genes of a transcription factor may have graded dose response to levels of transcription factor.⁴⁰ Our findings suggest that even a modest increase in *PRDM1* as a result of miR-BHRF1-2 inhibition is sufficient to effect a change in expression in a subset of *PRDM1* target genes. We identified several such genes and among them were able to validate *SCARNA20* (*ACA66*). We also provided preliminary evidence that *SCARNA20* mediates, at least

partially, *PRDM1* functions. *SCARNA20* is a small nucleolar RNA (snoRNA) localized in the Cajal's bodies.^{41,42} It belongs to the H/ACA class of snoRNA, which functions to guide site-specific pseudo-uridylation of small nuclear RNA U12, a component of the minor spliceosome.⁴¹ The minor spliceosome regulates the splicing of the U12-type intron, which represents a small minority of introns with low splicing efficiency but is present in genes regulating diverse cellular processes.⁴³ The role of U12-type intron splicing in human cancer has started to emerge. Recently, a subset of myelodysplastic syndrome was found to exhibit aberrant splicing of U12-type intron.⁴⁴ Thus, it is conceivable that EBV-miR-BHRF1-2 contributes to lymphomagenesis by downregulating *PRDM1* expression to a sufficiently low level at which *SCARNA20* expression is relatively uninhibited by *PRDM1*, resulting in aberrant U12 RNA modification and splicing of U12-type introns. SnoRNAs are generally encoded in the intron of protein coding genes and derived from processing of their precursor molecules.⁴⁵ Dysregulation of snoRNAs and their precursors in cancer development has been increasingly recognized.⁴⁶ Further experiments are necessary to determine whether *PRDM1* downregulates *SCARNA20* expression directly or through altering the level of its host gene.

In summary, we presented evidence that *PRDM1* is a target of EBV-miR-BHRF1-2 with likely pathogenic consequences. Previously, we have demonstrated promoter hypermethylation of *PRDM1* as a potential pathogenic event in EBV-positive BL.⁶ Thus, *PRDM1* appears to be an important gene target for inactivation or suppression in EBV-associated lymphomas. The mechanisms employed may depend on the latency type. EBV-miR-BHRF1-2 has also other potential targets^{22,47} and, therefore its effect on EBV-infected cells can be pleiotropic. The interaction between EBV-miR-BHRF1-2 and *PRDM1* may represent one of the many ways by which EBV-miR-BHRF1-2 promotes EBV lymphomagenesis. Our results support the potential of EBV-miR-BHRF1-2 as a therapeutic target in EBV-associated lymphomas.

CONFLICT OF INTEREST

The authors declare no conflict of interest.

ACKNOWLEDGEMENTS

This project is funded by the Department of Pathology and Laboratory Medicine of Weill Cornell Medical College. We would like to thank Felicia Chaviano for sample coordination.

AUTHOR CONTRIBUTIONS

JM designed and performed the experiments, analysed the data and wrote the manuscript; KN and LY performed experiments; DR and OE analysed the RNA-seq data; DMK contributed clinical cases, useful discussions and suggestions; WT conceptualized and directed the project, designed the experiments and wrote the manuscript.

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