Inhibition of Epstein-Barr Virus (EBV) Reactivation by Short Interfering RNAs Targeting p38 Mitogen-Activated Protein Kinase or c-myc in EBV-Positive Epithelial Cells

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Latent Epstein-Barr virus (EBV) is reactivated by 12-O-tetradecanoylphorbol-13-acetate (TPA) in EBV-infected cells. In this study, we found that TPA up-regulated phosphorylation of p38, a mitogen-activated protein kinase, and activated c-myc mRNA in EBV-positive epithelial GT38 cells. The EBV immediate-early gene BZLF1 mRNA and its product ZEBRA protein were induced following TPA treatment. Protein kinase C inhibitors, 1-(5-isoquinolinesulphonyl)-2, 5-dimethylpyraperazine (H7) and staurosporine, inhibited the induction of p38 phosphorylation and the activation of c-Myc by TPA. The p38 inhibitor SB203580 blocked both p38 phosphorylation and ZEBRA expression by TPA. Pretreatment of GT38 cells with the nitric oxide (NO) donor S-nitroso-N-acetylpenicillamine inhibited p38 phosphorylation and c-Myc activation by TPA, suggesting that NO may inhibit EBV reactivation via both p38 and c-Myc. By using short interfering RNA (siRNA) targeting either p38 or c-myc, we found that p38 or c-myc siRNA specifically inhibited expression of the respective gene and also suppressed the induction of ZEBRA and EBV early antigen. The interferon (IFN)-responsive gene expression tests ruled out the possibility that the antiviral effect of siRNA is dependent on IFN. Our present study demonstrates for the first time that either p38 or c-myc siRNA can efficiently inhibit TPA-induced EBV reactivation in GT38 cells, indicating that p38- and/or c-myc-associated signaling pathways may play critical roles in the disruption of EBV latency by TPA.

Epstein-Barr virus (EBV) is a human herpesvirus that infects B cells and epithelial cells to establish latent infection. EBV is associated with a variety of cancers, including Burkitt’s lymphoma (BL) (41), nasopharyngeal carcinoma (57), and gastric carcinoma (44). Upon primary infection, EBV infects epithelial cells, where it undergoes lytic replication, and B cells, where it usually remains latent (32, 41). Expression of the EBV immediate-early proteins BZLF1 (ZEBRA and Zta) or BRLF1 (Rta) is sufficient to convert EBV infection from its latent to lytic form (7, 50). Latency can be disrupted by certain BRLF1 (Rta) is sufficient to convert EBV infection from its immediate-early proteins BZLF1 (ZEBRA and Zta) or

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The c-myc proto-oncogene encodes a nuclear transcription factor that contains a basic helix-loop-helix leucine zipper domain and binds to the cis element CACGTG when dimerized with another nuclear factor, Max. Through regulating expression of various target genes, c-myc is actively involved in the control of cellular proliferation, differentiation, and apoptosis (39). c-myc is a key downstream target of the EBV latency-associated gene EBNA2 in infected B lymphocytes (3), and induction of c-myc expression by latency-associated genes likely plays a crucial role in promoting cell cycle progression (21, 40). Although latency-associated EBV genes can induce c-myc expression, these genes are not expressed in EBV-associated tumor in immunocompetent individuals (21). Instead, c-myc is translocated in BL (36) or overexpressed through other mechanisms in nasopharyngeal carcinoma (33). It is well known that TPA reactivates latent EBV in infected cells (13, 56) and activates c-myc in resting lymphocytes (23), T cells (35), chronic lymphocytic leukemia cells (28), and BL cells (8). However, the relationship between c-myc activation and EBV reactivation is not well understood. Therefore, study of the involvement of c-myc in EBV reactivation would be valuable to a further exploration of the mechanisms underlying EBV latency.

RNA interference (RNAi) is a highly conserved mechanism found in almost all eukaryotes and is believed to serve as an
antiviral defense mechanism. The molecular details are becoming clear due to combined genetic and biochemical approaches (51, 48). On entry into the cells, the double-stranded RNA (dsRNA) is cleaved by an RNase III-like enzyme, Dicer, into short interfering RNAs (siRNAs) (4, 17, 25, 26, 55). The siRNAs are incorporated into a multisubunit protein complex, the RNA-induced silencing complex, which directs the siRNA to the appropriate mRNA. This complex, when activated, can specifically silence or downregulate gene expression. RNAi has been used to study gene function in multiple model organisms, including plants (52), flies (24), Caenorhabditis elegans (11), and mice (53). However, in most mammalian cells, dsRNAs longer than 30 nucleotides activate an interferon (IFN) response, leading to nonspecific degradation of RNA transcripts and a general shutdown of host cell protein translation (47).

This nonspecific effect can be circumvented by the use of synthetic siRNA that are 21 nucleotides long with short 3′ overhangs (9). The synthetic siRNA has been shown to induce homology-dependent degradation of cognate mRNA and has been used to knock down expression of endogenous and heterologous genes in mammalian cell lines (5, 16, 19, 29, 38). RNAi interferes with the replication of a number of animal viruses including human immunodeficiency virus (19, 31, 38), hepatitis C virus (22), and gammaherpesviruses (20). However, the effect of RNAi on EBV has not been reported.

The aim of this study was to determine whether p38 and c-myc play a role in TPA-induced EBV reactivation. Here, we demonstrated that TPA-induced EBV reactivation in GT38 cells is dependent on PKC-mediated phosphorylation of p38 and c-myc activation and that the suppression of p38 phosphorylation by the specific inhibitor inhibited ZEBRA induction. Furthermore, we found that the RNAi efficiently inhibited TPA-induced ZEBRA expression and EBV early antigen (EA) through interference with either p38 or c-myc expression, suggesting that p38 and c-myc play key roles in the reactivation of EBV. siRNA targeting to either p38 or c-myc is sufficient to efficiently interfere with EBV reactivation by TPA in EBV-infected GT38 cells.

**MATERIALS AND METHODS**

**Cell line and reagents.** Cells of the GT38 cell line are EBV-positive epithelial cells derived from human gastric tissue (49). The cells were cultured in RPMI 1640 medium, supplemented with 10% fetal bovine serum, penicillin (100 U/ml), and streptomycin (100 μg/ml). Cells were incubated in an atmosphere of 95% air and 5% CO2. For EBV reactivation, cells were treated with 20 ng of TPA (Sigma Chemical Co., St. Louis, Mo.) per ml 1-(3-Isosquolinesulphonyl)-2,5-dimethylpyrroleperazine (HT) and staurosporine were purchased from Sigma. SB203580 and 5-nitroso-N-acetylpenicillamine (SNAP) were obtained from Calbiochem (Bio- science, Inc., La Jolla, Calif.) and dissolved in dimethyl sulfoxide just before being added to the culture medium.

**Preparation of RNA and Northern blot analysis.** Total RNA was prepared from cells by using an ISOGEN kit (Nippongene Inc., Tokyo, Japan) according to the manufacturer’s protocol as described previously (13, 14). The RNA was dissolved in diethyl pyrocarbonate-treated water and stored at −20°C. For all RNA samples, the ratio of the optical density at 260 nm to the optical density at 280 nm was >1.50.

Northern blot analysis was carried out as described previously (13, 14). Total RNA samples were denatured in 50% formamide and 2.2 M formaldehyde at 57°C for 15 min. Aliquots containing 20 μg of total RNA were loaded into 1% agarose gel containing 2.2 M formaldehyde and were electrophoresed with running buffer containing 50 mM MOPS (morpholinepropanesulfonic acid) acetate (pH 7.0), 10 mM sodium acetate, and 1 mM EDTA. The RNA was then transferred onto Hybond N+ membranes (Amerham, Buckinghamshire, United Kingdom) by capillary transfer and UV autocross-linked. Membranes were treated with prehybridization buffer for 6 to 24 h at 42°C. The solution was replaced with fresh hybridized solution containing the [α-32P]dCTP-labeled probe of c-myc (6) at a final concentration of 106 cpm/ml, and hybridization continued for 24 h at 42°C. The blots were washed three times at 65°C in 1× SSC (1× SSC is 0.18M NaCl, 0.01M NaHPO4), and 1 mM EDTA. The RNA was then exposed to Kodak X-AR film (Eastman Kodak Co., Rochester, N.Y.) at −80°C for 1 to 3 days. Northern blot membranes were stripped by boiling in 5 mM EDTA–0.1% SDS. They were then rehybridized with a specific probe for glyceroldehyde-3-phosphate dehydrogenase (GAPDH) (13) to control for variations in the amount of RNA per lane. Each experiment of at least three separate experiments performed at different times, unless otherwise specified.

**RT-PCR analysis.** Reverse transcription (RT)-PCR was performed essentially as previously described (13). The sequences of sense and antisense primers used for amplification of specific human 2′,5′-oligoasemadentanyl synthetase 1 (2,5 OAS; 40 kDa) cDNA were 5′-TGGCTGATTACCCCATGCTT-3′ (sense) and 5′-TGGACAGGGAGTGTGAAAT-3′ (antisense) to amplify a 265-bp product. The PCR primers for 2,5 OAS2 (71 kDa) were 5′-TTAAATGATAATCCCCAGGCC C-3′ (sense) and 5′-AAAGATTCTGCTGCGTCA-3′ (antisense) to amplify a 424-bp product; β-tubulin primers were 5′-TGATCTCTAGAACCCTGGGACC AT-3′ (sense) and 5′-ACCATTGITTACGTCGAACTGC-3′ (antisense). The amplified PCR products were analyzed after electrophoreses in a 1.2% agarose gel to compare relative induction levels of 2,5 OAS.

**RNA interference assays.** The 19-nucleotide p38 SMARTpool siRNA was purchased from Dharmacon Research, Inc. (Lafayette, Colo.). With SMART- pool, four individual siRNAs targeting the same p38 gene were combined into one pool by using a proprietary pooling algorithm of five parameters. The individual p38 siRNA sequences were as follows: 5′-GCAAGAAGACUAAUU ACAGU-3′, 5′-GAAUGCGGGUUAUCUUAAAC-3′, 5′-GAAUCUGGCGAAGUAUU-3′, and 5′-CAAGGCUCUGAGGAUACAA-3′. The scrambled p38 siRNA containing nonspecifically pooled duplex was obtained from Dharmacon Research, Inc. Silencer c-myc siRNA kits were obtained from Ambion Inc. (Austin, Tex.). The c-myc siRNA sequence is derived from the 3′ untranslated region of the human c-myc mRNA sequence. The scrambled c-myc negative control siRNAs, which have no significant homology to any known gene sequences from mouse, rat, or human, were obtained from Ambion Inc. According to the protocol of the manufacturer, the siRNA duplexes were resuspended with siRNA universal dilution buffer at a concentration of 20 μM. Cells were seeded in 12-well plates at a density of 5 × 105 to 7 × 105 cells well, incubated for 24 h, and then transfected with 100 nM p38 or c-myc siRNAs by using 2 μl of Lipofectamine 2000 transfection reagent (Invitrogen, Carlsbad, Calif.) according to the manufacturer’s optimized protocol; the transfected cells were then incubated for 48 to 72 h at 37°C.

**Western blot analysis.** Western blot analysis was performed on cell lysates as described previously (13). Protein samples (20 μg) were separated in SDS–10% polyacrylamide gel and electrophoretically transferred onto polyvinylidene difluoride membranes (Millipore Co., Bedford, Mass.), by using the semidry transfer blot system. The membrane was blocked with 5% skim milk in Tris-buffered saline containing 0.1% Tween 20 and subsequently incubated with primary antibody. Immunoblottings were performed by using the following antibodies: a 1:100 dilution of anti-c-Myc and a 1:200 dilution of anti-β-actin (Santa Cruz Biotechnology, Inc., Calif.); a 1:1,000 dilution of anti-phospho-c-Myc, anti-phospho-p38, and anti-p38 (New England, Biolabs Inc., Beverly, Mass.); and a 1:100 dilution of anti-ZEBRA monoclonal antibodies (developed in our laboratory).

After washing, the membrane was incubated with peroxidase-conjugated goat anti-rabbit or anti-mouse immunoglobulin G antibody (Amersham Biosciences Co.,) and the specific bands were detected by the Amersham ECL enhanced chemiluminescence technique system.

**Immunofluorescence analysis.** Approximately 2 × 105 cells were seeded in Lab-Tek eight-well chamber slides (Nalge Nunc International, Naperville, Ill.). After 24 h incubation, the cells were transfected with 100 nM concentrations of the siRNAs according to the procedure described above. At 2 days posttransfection, the cells were treated with or without TPA (20 ng/ml) for 72 h, washed twice with phosphate-buffered saline (PBS), and then fixed with acetone for 10 min at room temperature. Thereafter, cells were incubated with anti-EA antibodies, which were derived from a patient’s serum, at a final dilution of 1:40 for 45 min at 37°C. After washing with PBS, fluorescein isothiocyanate-conjugated anti-human antibodies (Organon Teknika Co., Durham, N.C.) were added and incubated for another 45 min at 37°C. The slides were washed and mounted with 1:glycerol-PBS and finally were examined under a fluorescence microscope.

**Colorimetric (MITT) assay for cell proliferation.** Cell growth was assayed by incorporation of MTT 3-[4,5-dimethylthiazol-2-yl]-2,5-diphenyl tetrazoliu bro
mide dye (Chemicon International, Inc., Temecula, Calif.). Cells were seeded on 96-well plates (Corning Inc., Corning, N.Y.) at a density of 5 × 10² cells/100 μl per well. At the indicated times, 10 μl of MTT solution (5 mg/ml) was added to each well, and the plates were incubated for 4 h at 37°C. One hundred microliters of isopropanol with 0.04 N HCl was added to each well and mixed by repeated pipetting. Absorbance was measured at a test wavelength of 570 nm and a reference wavelength of 655 nm with a microplate reader (model 550; Bio-Rad, Richmond, Calif.).

Statistical analysis. The results for Northern and Western blotting analyses were quantified according to a method described previously (13). Data are expressed as means ± standard deviations. The data shown were mean values of at least three different experiments. A Student's t test was used to compare the results. A P value of less than 0.01 is considered statistically significant.

RESULTS

Effects of TPA on the expression and/or activation of BZLF1 or BRLF1, c-myc, and p38 in GT38 cells. We previously reported that the expression of EBV immediate-early genes BZLF1 and BRLF1 in GT38 cells was induced by TPA (13). To further investigate cellular genes that are associated with EBV reactivation by TPA, we focused on p38 and c-myc expression. GT38 cells were incubated with 20 ng of TPA per ml from 0 to 24 h. BZLF1 and BRLF1 mRNAs were induced at 0.5 h and reached a maximum level at 8 h. The c-myc mRNA level began to increase at 0.5 h, reached maximum at 8 h, and then declined at 12 h after TPA stimulation (Fig. 1A). To determine whether activation of p38 or/and c-myc is associated with EBV reactivation, we used Western blot analysis to examine p38 phosphorylation and c-myc activation in GT38 cells treated with TPA (Fig. 1B). Phosphorylation of p38 reached a maximum at 0.5 h, decreased to a lower level at 1 h, and returned to the basal level at 4 to 24 h. TPA treatment did not alter the expression levels of total p38. The expression of c-Myc was increased at 1 h, reached a plateau at 2 to 8 h, and then decreased at 24 h after TPA treatment. ZEBRA was detected at 8 and 24 h. In the parallel samples without TPA treatment, the phosphorylation of p38, the activation of c-Myc, and ZEBRA expression remained essentially unchanged at different time points (data not shown). These results indicated that TPA is able to induce the concomitant activation of p38 and c-myc during EBV reactivation in GT38 cells.

Effects of p38 and PKC inhibitors on p38 phosphorylation and expression of c-myc and ZEBRA. TPA is a potent activator of PKC (37). It was previously reported that TPA induced
EBV reactivation via the PKC pathway in GT38 cells (13). To examine whether the activation of the c-myc gene is associated with TPA-induced PKC activation, GT38 cells were pretreated with potent PKC inhibitors, H7 and staurosporine, and the p38 inhibitor SB203580 for 1 h and then treated with TPA for 4 h (Fig. 2A). Consequently, the induction of c-myc mRNA by TPA was greatly inhibited by H7 pretreatment. Likewise, staurosporine also inhibited c-myc mRNA expression, although the inhibitory effect was weaker than that of H7. In contrast, the p38-specific inhibitor SB203580 did not affect the induction of c-myc mRNA by TPA (data not shown). To examine whether TPA induces EBV reactivation via the p38 and/or c-myc pathways as regulated by PKC, we performed immunoblotting analyses (Fig. 2B). H7 and staurosporine greatly inhibited the phosphorylation of p38 and activation of c-Myc and ZEBRA by TPA. SB203580 blocked the phosphorylation of p38 and ZEBRA expression by TPA but did not affect the expression of c-Myc. The protein levels of total p38 were not affected by SB203580, H7, or staurosporine. To examine whether p38 regulates the phosphorylation of c-Myc, we performed Western blot analyses using anti-phospho-c-Myc antibodies. The phosphorylation pattern of c-Myc was similar to that of c-Myc activation. Both H7 and staurosporine inhibited the phosphorylation of c-Myc by TPA, but SB203580 had no effect on the phosphorylation of c-Myc. Taken together, these results indicated that PKC is an upstream effector of p38 and c-myc activation, which are essential for ZEBRA expression by TPA, and that the phosphorylation of p38 and c-myc may be in independent pathways regarding their roles in TPA-induced EBV reactivation.

Effects of SNAP, an NO donor, on p38 phosphorylation and c-Myc activation by TPA in GT38 cells. It was previously re-
siRNA targeting to either p38 or c-myc inhibits EBV reactivation in GT38 cells. To further study the critical roles of p38 and c-Myc in EBV reactivation, we analyzed the loss of function of p38 and c-myc on EBV reactivation with RNAi methods. We transfected GT38 cells with p38 siRNAs by using Lipofectamine 2000 and then treated the cells with TPA (Fig. 4A). The phosphorylation of p38 was markedly downregulated by p38 siRNAs at 48 and 72 h after transfection. Expression of total p38 was also downregulated by the siRNA. TPA-mediated induction of ZEBRA expression was also inhibited by p38 siRNAs. Similarly, GT38 cells were transfected with c-myc siRNA, and thereafter the expression of c-Myc was examined by Western blot analysis (Fig. 4B). As a result, activation of c-Myc by TPA was inhibited by c-myc siRNA at 48 and 72 h posttransfection. ZEBRA induction by TPA was also inhibited by c-myc siRNA. To investigate the effect of siRNA targeted to p38 on c-myc expression and siRNA targeted to c-myc on p38 phosphorylation, GT38 cells were transfected with p38 or c-myc siRNA for 48 or 72 h and then treated with TPA. The phosphorylation of p38 was not affected by c-myc siRNA transfection, and p38 siRNA also had no effect on c-Myc activation (data not shown). These results demonstrated that siRNAs targeted to p38 and c-myc can efficiently inhibit TPA-induced ZEBRA expression in GT38 cells and that both the p38- and c-Myc-associated pathways are crucial for TPA-mediated EBV reactivation.

p38 and c-myc siRNAs inhibit EA induction in GT38 cells. To further examine whether p38 or c-myc siRNA could inhibit EA expression in GT38 cells, the cells were transfected with p38 or c-myc siRNA and then treated with TPA at 48 h posttransfection. The expression of EA was analyzed by the immunofluorescence method (Fig. 5). EA was detected in 2.9% of untreated GT38 cells. After TPA treatment for 72 h, the percentage of EA-positive cells was increased to 9.4% (threefold increase; P = 0.001). However, the percentage of EA-positive cells was decreased to 4.1 and 4.3% (2.3-fold [P = 0.003] and 2.2-fold [P = 0.002], respectively) in cells transfected with p38 and c-myc siRNAs, respectively, whereas no significant reduction of EA-positive cells was observed in the scrambled siRNA-transfected cells (P = 0.468). These results indicated that both p38 and c-myc siRNAs can block TPA-mediated induction of EA in GT38 cells and strongly suggested that siRNAs targeted to p38 and c-myc are able to inhibit EBV reactivation.

siRNA transfection neither activates IFN-induced 2,5 OAS induction nor induces cell-growth inhibition. To rule out the possibility that the siRNA-mediated inhibition of EBV reactivation in GT38 cells could be mediated by dsRNA-induced activation of the IFN pathway, we analyzed the expression levels of 2,5 OAS1 and 2,5 OAS2 mRNA, known as the IFN-induced gene (22). The relative amounts of 2,5 OAS1 and 2,5 OAS2 transcripts were determined by RT-PCR. IFN-α induced the expression of 2,5 OAS1 and 2,5 OAS2 mRNAs in GT38 cells (Fig. 6A). However, 2,5 OAS1 mRNA was not detected in any group of cells that were untransfected or transfected with siRNA targeted to p38 or c-myc or with an irrelevant scrambled siRNA. Although low-level expression of 2,5 OAS2 mRNA was detected in untransfected GT38 cells and in GT38 cells transfected with p38, c-myc, or scrambled siRNAs, no difference was observed between untransfected cells and

![Experimental figure](http://jvi.asm.org/)

**FIG. 3.** TPA-induced phosphorylation of p38 and activation of c-Myc were inhibited by the NO donor SNAP. GT38 cells were pretreated with SNAP at indicated concentrations for 1 h prior to TPA (20 ng/ml) treatment. Cell lysates were prepared for phosphorylation of p38 and total p38 (t-p38) and of c-Myc at 0.5 and 4 h, respectively, after TPA treatment. Cell lysates were separated by SDS-PAGE and blotted onto membranes. Phosphorylated p38, p38, and c-Myc were detected by Western blotting with anti-phospho-p38, anti-p38, anti-c-Myc or anti-β-actin antibodies. The p38 and β-actin were used as internal standards. The relative densitometric units of phosphorylation of p38 and activation of c-Myc were determined to p38 and β-actin signals, respectively. The data represent three independent experiments and the standard deviations are shown.

Support that NO downregulates the induction of EBV reactivation by TPA (13). To determine the effects of NO on p38 and/or c-myc activation, GT38 cells were treated with TPA in the absence or presence of the NO donor SNAP, which generates NO in an aqueous solution, resulting in the formation of peroxynitrite. The phosphorylation of p38 and expression of c-Myc were examined by immunoblotting analysis (Fig. 3). TPA-mediated phosphorylation of p38 and c-Myc activation were inhibited by SNAP in a dose-dependent manner; SNAP (0.1 mM) only has a slight inhibition effect, and the maximum inhibition was observed at a concentration of 1 mM SNAP. TPA-mediated induction of BZLF1 mRNA and ZEBRA protein were also inhibited by SNAP (reference 13 and data not shown). These results suggested that NO may inhibit TPA-induced EBV reactivation via the inhibition of the p38 and c-myc pathways.
cells transfected with p38, c-myc, or scrambled siRNAs. We concluded that IFN induction is not involved in the decrease in the phosphorylation of p38 or the expression of c-myc and ZEBRA by the siRNAs.

To confirm whether the RNAi-mediated inhibition of EBV reactivation was not due to siRNA transfection-induced cytotoxicity, cell growth was monitored by an MTT assay (Fig. 6B). TPA treatment did not affect cell growth, and the growth pattern of siRNA-transfected cells was quite similar to that of cells not treated with TPA or treated only with TPA. These results indicated that the suppression of EBV reactivation in the siRNA-transfected cells was not due to the cytotoxicity of p38 and c-myc siRNA in the cells.

**DISCUSSION**

It has been reported that EBV latency can be disrupted by TPA in the EBV-positive epithelial cell lines GT38 and GT39 and that the effect of TPA was mediated by the PKC- and MAPK-associated pathways (13). In this study, we focused on

![FIG. 4](image-url)  
**FIG. 4.** siRNAs targeting p38 and c-myc specifically inhibit the expression of respective genes and ZEBRA expression. (A) GT38 cells were transfected with p38 siRNA (0.1 μM) or scrambled siRNA (scr) and incubated for 48 and 72 h or for 72 h and then treated with TPA for 30 min for phosphorylated p38 and total p38 (t-p38) and 48 h for ZEBRA. The cell extracts were analyzed for phosphorylation of p38, total p38, ZEBRA, and β-actin by Western blotting. (B) GT38 cells were transfected with siRNA (0.1 μM) targeted to c-myc for 48 or 72 h or with scrambled (scr) siRNA for 72 h and then treated with TPA (20 ng/ml) for 4 or 48 h for the expression analysis of c-Myc and ZEBRA, respectively. Cell lysates were harvested from siRNA-transfected or untransfected GT38 cells at indicated times posttransfection. C-Myc and ZEBRA expression were analyzed by Western blotting by using anti-c-Myc, anti-ZEBRA, and anti-β-actin antibodies. The signals of phosphor-p38, t-p38, and ZEBRA (panel A) and of c-Myc and ZEBRA (panel B) were quantified by a phosphorimage system. The relative protein level was measured as ratio of β-actin protein. The data represent three independent experiments, and the standard deviations are shown.

![FIG. 5](image-url)  
**FIG. 5.** p38 or c-myc siRNA blocks EA induction by TPA. GT38 cells were transfected with siRNA (0.1 μM) targeting p38 and c-myc or with scrambled (scr) siRNA. At 48 h posttransfection, cells were treated with TPA (20 ng/ml) for 72 h. EA-positive cells were determined by indirect immunofluorescence. The data shown represent three independent experiments. Bars represent standard deviations. (*) P < 0.01 versus the TPA-positive and siRNA-negative cells.)
the roles of p38 MAPK and c-Myc that were suggested to be the downstream signaling molecules of PKC in the TPA-induced EBV reactivation in the GT38 cell line and found that TPA induced phosphorylation of p38 and c-myc activation. We analyzed whether PKC activates p38 and c-myc during EBV reactivation. To address this question, we treated GT38 cells with the PKC inhibitors H7 and staurosporine. The phosphorylation of p38 and c-myc activation by TPA were abolished by the inhibitors. Furthermore, GT38 cells were treated with p38 inhibitor SB203580; consequently, both phosphorylation of p38 and expression of ZEBRA were simultaneously inhibited. These findings indicate that p38 and c-myc are downstream effectors of PKC on EBV reactivation and that those pathways may play important roles in regulating the stringency of viral latency.

Previous reports have shown that TPA induces BZLF1 expression by inhibiting inducible nitric oxide synthase expression in GT38 cells (13, 14). TPA-mediated induction of BZLF1 expression was abolished by PKC inhibitors, and TPA down-regulated inducible nitric oxide synthase gene expression was reversed by PKC inhibitors (13). In order to extend the observation, in the present study we further examined the effects of the NO donor SNAP in the TPA-mediated EBV reactivation in GT38 cells and found that the phosphorylation of p38 and c-myc activation by TPA were inhibited by SNAP in a dose-dependent manner. This finding suggested that NO first down-regulates the phosphorylation of p38 and c-Myc and then leads to the inhibition of BZLF1 expression.

c-myc can be activated by TPA in many kinds of cells including BL cells (8, 23, 28, 35). The normal c-myc allele in BL cells is activated by TPA. Based on these observations, we studied the effects of TPA on c-Myc activation and EBV reactivation and found that c-Myc was activated by TPA in GT38 cells and that c-myc activation is closely associated with BZLF1 and ZEBRA expression. These results suggested that activation of c-Myc may be important for efficient lytic EBV infection. The results that PKC inhibitors inhibited the activation of c-Myc suggested that PKC, as an upstream kinase of EBV reactiva-

![Fig. 6. Effects of siRNAs targeted to p38 and c-myc on the 2,5 OAS induction and cell growth.](http://jvi.asm.org/)

(A) GT38 cells were transfected with p38, c-myc, or scrambled (scn) siRNA. At 48 h posttransfection, cells were treated with TPA (20 ng/ml) for 48 h. As the positive control of 2,5 OAS, GT38 cells were treated with IFN (300 U/ml) for 6 h. The expression levels of 2,5 OAS1 and 2,5 OAS2 were determined by RT-PCR. β-tubulin mRNA was amplified from the same cDNA as an internal standard. (B) GT38 cells were plated in 96-well plates and incubated for 24 h and then transfected with siRNA duplexes targeted to c-myc or p38 or scrambled (scn) siRNA. Cell growth was analyzed by using MTT assays.
tion, may directly and/or indirectly activate c-Myc and finally lead to EBV reactivation. SB203580 did not affect the activation and phosphorylation of c-Myc. This indicates that the phosphorylation of p38 and c-myc activation are in independent pathways. To our knowledge, this is the first report that provides direct evidence linking c-Myc activation to EBV reactivation.

Previous reports demonstrated that Zta downregulated c-Myc protein expression (42) and that the c-myc transfectants expressed low levels of BZLF1 protein and of its mRNA (10). These observations seem to contradict our present findings. However, in the report by Fais et al. (10) it was also shown that TPA could reinduce the expression of EBV lytic antigens. It indicated that c-myc transfectants did not completely lose the capacity to initiate the lytic cycle. Although the molecular mechanisms underlying this are not yet clearly understood, these observations reiterated the correlations existing between c-myc expression and regulation of the EBV lytic cycle.

We have demonstrated that siRNAs targeted to p38 and c-myc blocked TPA-mediated EBV reactivation in transfected GT38 cells. The p38 and c-myc siRNAs specifically inhibited expression of their respective proteins, whereas the scrambled siRNA, which has no significant homology with the targeting gene sequences, affected neither p38 nor c-Myc protein levels. ZEBRA and EA inductions were significantly inhibited by p38 and c-myc siRNAs but not by an irrelevant scrambled siRNA. siRNA designed against p38 had no effect on c-Myc expression; siRNA targeted against c-myc also did not affect the phosphorylation of p38 (data not shown). These results further confirmed that the TPA-induced phosphorylation of p38 and activation of c-myc belong to independent transduction pathways.

The percentages of EA-positive cells were also significantly reduced by p38 or c-myc siRNAs. To rule out the possibility that the inhibition may be due to the cytotoxicity of siRNA transfection, we showed by using an MTT assay that both p38 and c-myc siRNAs inhibited EA induction in a cytotoxicity-independent manner. This finding indicated that the inhibiting effects of siRNAs are not the result of cytotoxicity.

Recently, it has been reported that transfection of siRNAs probably cause IFN-mediated activation of the Jak-STAT pathway, and this effect is mediated by the dsRNA-dependent protein kinase PKR and 2,5 OAS, which are activated by siRNAs and are required for upregulation of IFN-β in response to siRNA (45). To further clarify this question, transcripts of the IFN-inducible 2,5 OAS gene were examined by RT-PCR in siRNA-transfected GT38 cells. Consequently, no induction of 2,5 OAS gene was found in cells transfected with p38 or c-myc siRNAs. These results ruled out the possibility of an IFN response in the downregulating effects of p38, c-myc, and ZEBRA by siRNA transfection. Thus, our results showed that RNAi could be used to study the ability of host genes to regulate EBV reactivation in this system. Recently, many viruses have been shown to be susceptible to inhibition by RNAi, suggesting that RNAi may play an adaptive antiviral role in controlling these viral infections (2, 19, 34, 38). In addition, some viruses have evolved mechanisms to evade RNAi. For example, flock house virus induces RNAi in Drosophila cells, yet the flock house virus B2 protein suppresses this antiviral effect (31), reflecting an attempt to escape from this endogenous antiviral host response.

In conclusion, our study demonstrates that p38 and c-myc play critical roles in TPA-induced EBV reactivation in EBV-positive epithelial GT38 cells and that RNAi provides an antiviral response in EBV-infected cells. Using the efficient RNAi method, we can identify essential genes involved in EBV reactivation and suppress EBV reactivation and replication in infected cells. RNAi may potentially represent a powerful therapy for EBV-associated disorders such as BL, nasopharyngeal carcinoma, gastric carcinoma, etc.

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