Identification of a Boundary Domain Adjacent to the Potent Human Cytomegalovirus Enhancer That Represses Transcription of the Divergent UL127 Promoter†

ANA ANGULO,1 DAVID KERRY,1 HUANG HUANG,2 EVA-MARIA BORST,2 ALISON RAZINSKY,1 JUN WU,1,‡ URS HOBOM,2 MARTIN MESSERLE,2 AND PETER GHAZAL1,*

Department of Immunology and Molecular Biology, Division of Virology, The Scripps Research Institute, La Jolla, California 92037,1 and Max von Pettenkofer Institute, Munich, Germany2

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Transcriptional repression within a complex modular promoter may play a key role in determining the action of enhancer elements. In human cytomegalovirus, the major immediate-early promoter (MIEP) locus contains a highly potent and complex modular enhancer. Evidence is presented suggesting that sequences of the MIEP between nucleotide positions −556 and −673 function to prevent transcription activation by enhancer elements from the UL127 open reading frame divergent promoter. Transient transfection assays of reporter plasmids revealed repressor sequences located between nucleotides −556 and −638. The ability of these sequences to confer repression in the context of an infection was shown using recombinant viruses generated from a bacterial artificial chromosome containing an infectious human cytomegalovirus genome. In addition to repressor sequences between −556 and −638, infection experiments using recombinant virus mutants indicated that sequences between −638 and −673 also contribute to repression of the UL127 promoter. On the basis of in vitro transcription and transient transfection assays, we further show that interposed viral repressor sequences completely inhibit enhancer-mediated activation of not only the homologous but also heterologous promoters. These and other experiments suggest that repression involves an interaction of host-encoded regulatory factors with defined promoter sequences that have the property of proximally interfering with upstream enhancer elements in a chromatin-independent manner. Altogether, our findings establish the presence of a boundary domain that efficiently blocks enhancer-promoter interactions, thus explaining how the enhancer can work to selectively activate the MIEP.

The major immediate-early promoter (MIEP) enhancer region of cytomegalovirus (CMV) is required for optimal infection (3) and plays an important role in determining the cell type tropism and state of activation in vivo (4, 5, 24, 28). The MIEP is highly complex and contains one of the most potent transcriptional enhancers known to date (7; reviewed in reference 20). Previous studies have demonstrated the importance of a wide variety of positive cis-acting enhancer modules in promoting high levels of transcription from the human CMV (HCMV) MIEP. In addition, the viral IE2 (IE86 and L40) proteins act as autorepressors of the HCMV MIEP by binding the cis repression sequence (14, 30, 34, 39, 41, 42). The location of this sequence, between the initiation site of transcription and the TATA box sequence, enables IE2 proteins to competitively block the recruitment of RNA polymerase II at the MIEP (31, 42). Repression is also predicted to be essential for establishing stringent regulation of the MIEP enhancer, which is likely to be an important feature in determining temporal and cell-type-specific patterns of viral gene expression as well as preventing promiscuous promoter interactions.

In the species-specific mouse CMV (MCMV), a similarly complex but distinct enhancer/promoter region is involved in controlling expression from its major immediate-early (MIE) genes (15). A divergent promoter upstream from its MIEP drives the expression of the MCMV ie2 gene (Fig. 1) (37). This gene is dispensable for growth in tissue culture and in vivo and comes under control of the nearby enhancer, thereby displaying IE kinetics of expression (9, 35). The observation that the MCMV enhancer plays a dual role in regulating ie1 and ie2 expression is not unexpected since by definition enhancers function equally well in either orientation. Similar to the organization of MCMV MIE locus, the genome sequence from HCMV (11) identified a divergent open reading frame (ORF) (UL127) in close proximity to the enhancer (Fig. 1) and which overlaps the modulator domain of the human MIEP (Fig. 2A). While the HCMV UL127 ORF is dispensable for replication in tissue culture, it does not appear to have any significant homology with the mouse viral ie2 gene (36, 37). In contrast to the MCMV ie2 gene, expression of UL127-encoding transcript is not detected upon infection of human foreskin fibroblasts (HFF cells) by HCMV (10). The UL127 promoter region contains an excellent TATA box (10 out of 11 nucleotides [nt] match with the MIEP TATA box [Fig. 1]) and binds CTF-1/ NF-1 at the expected CAAT box position (18). Thus, the observation for inactivity of this divergent promoter is unexpected due to the strong potency of the CMV enhancer and raises the question as to why the enhancer is unable to influence expression of the UL127 promoter site. It is conceivable that repressors stringently regulate the UL127 promoter. To investigate this issue, we have examined the requirement of cis-acting sequences in the promoter-proximal region of UL127 for limiting enhancer action.

Here, we present evidence that sequences located between −556 and −673 of the MIEP serve to completely block en-
hancer interactions with the UL127 promoter site. This region therefore plays an important role in defining the boundary of action of the MIEP enhancer in HCMV.

MATERIALS AND METHODS

Cells. The cell lines U373-MG, HeLa, NT-2/D1, and MRC-5, as well as the HFF cells, were grown in Dulbecco's modified Eagle medium supplemented with 2 mM glutamine, 100 U of penicillin per ml, 100 \( \mu \)g of gentamicin per ml, and 10% (vol/vol) fetal bovine serum.

Plasmid constructions and transfections. Plasmid pSnaB-Bam(HI)CAT was constructed by inserting a BamHI/SpeI fragment from pMIEP(HI)CAT (17) into BamHI/Spel-digested pSnaB-BamCAT. Plasmid pSnaB-BamCAT contains regulatory sequences of the HCMV MIE gene from nt 2243 to 2741 linked to the chloramphenicol acetyltransferase (CAT) gene. To construct pGACC(2531) and pGACC(2673), recombinant plasmid pGACC was generated by inserting annealed oligonucleotides G53 (5'-GATCCATTGGTTATATAGCATACTA-GTAAGCTTCTACGTAC-3') and G52 (5'-AGCTGTACGTAGAAGCTTACTAGTTATGCTATATAACCAATG-3'), containing SnaBI, HinIII, and SpeI sites, into BamHI/HinIII-digested pMIEP(266/112)CAT (42). A HinIII/SnaBI fragment from D1 and pR55 (kindly provided by B. Fleckenstein) was cloned into pGACC digested with HinIII/SnaBI to create pGACC(2531) and pGACC(2598), respectively. Plasmids pGACC(2531) and pGACC(2673) were digested with SnaBI and BamHI and ligated into Smal/BglII-digested pGL3-Basic (Promega, Madison, Wis.) to create pGL3(−531) and pGL3(−673), respectively. To construct pE(−66/−112)CAT and pEus(−66/−112)CAT, a 190-bp BamHI/HindIII fragment from pMIEP(−66/−112) (44) that contains HCMV MIEP sequences from −66 to −112 was inserted into BamHI/HindIII-digested pGACC(−531) and pGACC(−598), respectively. Plasmids pGL3(−634), pGL3(−581), pGL3(−556), and pGL3(−531) were constructed using the PCR primers promExt-1 (5'-TGTACGTAGATGTACTGCCAAGT-3'), promExt-2 (5'-TGAAGCTTGGTCATTAGTTCATAGCCAT-3'), promExt-3 (5'-TGAAAGCTTATGTTATAATAGTAATCAATTAC-3'), promExt-4 (5'-TGAAAGCTTATGTTATAATAGTAATCAATTAC-3'), and promExt-5 (5'-TGAAGCTTATGTTATAATAGTAATCAATTAC-3'). All PCRs were performed with pMIEP(−1145/−112)CAT (19) and primer promExt-1. Primers promExt-2, promExt-3, promExt-4, and promExt-5 were used to generate pGL3(−634), pGL3(−581), pGL3(−556), and pGL3(−531), respectively. Primer promExt-1 contains a 5' SnaBI linker, while primers promExt-2, promExt-3, promExt-4, and promExt-5 contain a 5' HindIII linker. The corresponding PCR products were digested with SnaBI and HindIII and inserted into SnaBI/HindIII digested pGL3(−531). The SnaBI/BamHI fragments excised from the recombinant plasmids generated were then inserted into Smal/BglII-digested pGL3-Basic to generate plasmids pGL3(−634), pGL3(−581), pGL3(−556), and pGL3(−531). Plasmids pGL3(Δ−531/−634) and pGL3(Δ−531/−609) were made by utilizing PCR primers promRev-1 (5'-TGGGATCCATTGGTTATATA-3'), promRev-2 (5'-TGTGATCATACTATTATGATATG-3'), and promRev-3 (5'-TGT...
FIG. 2. NF-1 binding sites flanking the UL127 TATA box do not mediate transcriptional repression. (A) Binding of various transcription factors (marked by open boxes) to HCMV MIEP sequences between −244 and −781 (based on DNase I footprinting data [reviewed in reference 20]). Locations of the enhancer, usr, NF-1 cluster, modulator, and UL127 TATA box are shown. Numbers refer to nucleotide positions relative to the transcription start site (+1) of the MIEP. CAT reporter constructs [pSnaB-BamCAT, pSnaB-Bam(HI)CAT, pGACC(−673), and pGACC(−531)] with various 5' and 3' deletion endpoints are shown below. (B) HFF, HeLa, U373-MG, and NT-2/D1 cells were transfected with 5 μg of the various promoter-CAT deletion constructs shown in panel A together with 5 μg of the control plasmid pRSV-β-gal. Cell lysates were prepared 30 h after transfection and assayed for β-galactosidase and CAT activity. For the CAT assays, cell extracts containing the same amount of β-galactosidase activity were used. A plot of the normalized percentage of CAT activity calculated for each construct taking as 1 the activity presented by pSnaB-BamCAT is shown. The CAT values shown represent the average ± standard deviation (bars) of three determinations.
GATCAGTCAACATGGCGGT-3'). All PCRs were performed with pMIEP(−1145/+112)CAT and primer promRev-1. Primer promRev-2 was used to create pGL3(Δ−531/+638), and promRev-3 was used to generate pGL3(Δ−531/+609). Primer promRev-1 contains a 5′ BamHI linker, while primers promRev-2 and promRev-3 contain a 5′ BclI linker. The two PCR products generated were cloned between the SpeI and the BamHI sites of pGACC(−531). SmaI/BamHI fragments were excised from the two recombinant plasmids constructed and inserted into SmaI/BglII-digested pGL3-Basic.

The β-galactosidase expression vector pRSV-β-gal and the luciferase expression vector pRL-tk (Promega) were used as internal controls in transfection assays. Transfections were performed by the calcium phosphate precipitation method as described previously (19). Cell lysates were prepared 30 h after transfection. β-Galactosidase and CAT assays were performed as previously described (2). Luciferase activity was determined according to the Promega’s Dual-Luciferase reporter assay system technical manual. The activity of the experimental luciferase reporters was normalized to the activity of the internal control pRL-tk. For CAT assays, cell extracts containing the same amount of β-galactosidase were used. CAT activity was quantitated by using a Molecular Dynamics Phosphor imager system with ImageQuant software.

**BAC mutagenesis and virus construction.** For generation of recombinant viruses UL127-GFP1, UL127-GFP10, and UL127-GFP7, three plasmids were constructed that contained a green fluorescent protein (GFP) reporter gene from pEGFP-C1 (Clontech, Palo Alto, Calif.) downstream of the UL127 sequences which were isolated from reporter plasmids pGACC(−673), pGACC(−531/−638), and pGACC(−531), respectively (Fig. 2 and 6). Note that in these constructs the GFP gene was inserted as a separate ORF upstream of the UL127 ORF but downstream of the UL127 promoter sequences. Briefly, enhanced GFP (EGFP) primers 5′-GGCCCTGCAGATCTGCTAGCGCTACCGGTCGCCA-3′ and 5′-GGCCCTGCAGTTTAAACTCACTTGTACAGCTCGTCCATGCC-3′, containing BglII and Pmel sites, respectively, were used to PCR amplify the EGFP gene. The resultant PCR fragment was digested with PstI and inserted into the XhoI site at position −755 of pMIEP(−1145/+112)CAT. Next a BglII site (introduced from the EGFP primer) and the SmaI site of the wild-type (wt) MIEP sequence (at position −243) were used to replace the wt MIEP sequence
with BamHI/SnaBI fragments containing the deletion mutations of pGACC (~673), pGACC–~531/~638, and pGACC–~531, respectively. A tetracycline resistance gene flanked with FRT sites was excised from plasmid pFL16 (13) and inserted into a unique SnaBI site downstream of the EGFP gene. The resulting constructs were digested with PstI (~1145) and Eagl (~78), generating fragments that contained one of the three different MIEP sequences each, the GFP gene, the tetracycline resistance gene, and 385 bp of downstream sequences complementary to the UL127 ORF. Recombination between the linear fragments and the HCMV bacterial artificial chromosome (BAC) plasmid pHBS (6) was performed in the recombination-prolific Escherichia coli strain JC879 in accordance to a recently described mutagenesis procedure (44). Minipreparations of BAC DNA isolated from 10-mL bacterial cultures were prepared and characterized by digestion with restriction enzymes SpeI, SalI, and SnaBI. HCMV BACs that had received the mutation were then transformed in the recombination-deficient E. coli strain DH10B. Excision of the tetracycline resistance cassette with Flp recombinase was performed essentially as described elsewhere (13) and confirmed by restriction enzyme analysis. In addition, the regions of interest were sequenced in the different BACs. Sequences were as expected for all recombinants except in BAC UL127-GFP10 between nt ~688 and ~676 relative to the transcription start site of the MIEP that contain 10 mismatched nucleotides, disrupting the NF-1 site 5′ proximal to the UL127 TATA box. Midipreparations of BACs were used to transfect MRC-5 cells for generation of infectious recombinant viruses as described previously (6). Infectivity of HFF-1 cells used to generate viral stocks, as well as infectivity of infected cells were determined by standard methods.

**In vitro run-off transcription assay.** Transcription reactions (25 μl) were performed in nuclear extracts prepared from HeLa cells as described previously (18). The templates pMIEP (~66 +/112), pE (~66 +/112), and pEars (~66 +/112) were linearized with EcoRI and used in the transcription reactions at a concentration of 25 μg/ml. Trancript products were analyzed by electrophoresis on a denaturing (8.3 M urea) 6% polyacrylamide gel. Relative levels of transcription were quantitated using a PhosphorImager (Molecular Dynamics). Poly(U) polymerase activity present in the nuclear extract was used as an internal standard to account for variability between samples.

**Reverse transcription (RT)-mediated PCR (RT-PCR).** HFF cells were infected with the different recombinant viruses at a multiplicity of infection (MOI) of 0.5 PFU/cell. On day 3 after infection, infected cells were fixed and permeabilized with 4% paraformaldehyde, 0.2% glutaraldehyde, and 0.2% Triton X-100 in PBS. Cells were incubated with HCMV IE monoclonal antibody 810 (dilution 1/30) for 1 h at 37°C. The coverslips were extensively washed with PBS containing 1% bovine serum albumin and incubated with a secondary antibody, tetramethylrhodamine isothiocyanate-(TRITC)-conjugated goat anti-mouse (dilution 1:800; Sigma Immunochemicals). After a further 60-min incubation at 37°C, the coverslips were washed again with PBS containing 1% bovine serum albumin, rinsed with water, and mounted with Permount. Samples were examined using a Zeiss Axiosplan confocal microscope and a 63× oil immersion objective lens. Data were collected at a resolution of 512 by 512 pixels. Data sets were processed using the NAC 1024 software and then exported for preparation for printing using Adobe Photoshop.

**RESULTS**

**Sequences upstream of the NF-1 cluster mediate transcriptional repression of the UL127 promoter site.** Evidence for direct repression of the MIEP by its own IE2 gene product has been provided to account for the restricted expression of the IE genes in the later stages of infection (14, 30, 31, 34, 38, 39, 42). This repression system is defined by the competitive binding of the IE repressor with the RNA polymerase II recruitment step (31, 42). Our previous studies indicated that a strict requirement for this type of repression is that the position of the repressor binding site should be in the vicinity of the transcription start site (31). In other studies, binding sites 5′ proximal to a TATA box have also been shown to block the recruitment of an RNA polymerase II complex (reviewed in reference 1). The UL127 promoter is located in the center of the NF-1 cluster in which one NF-1 binding site exists 14 bp downstream of the UL127 TATA box in the immediate vicinity of the start site and another lies 10 bp upstream of the TATA box (25, 27). It is possible that these particular TATA proximal NF-1 sites may play a role in maintaining a repressed state. To investigate whether these NF-1 sites are involved in repression, we examined constructs in which promoter-proximal downstream and upstream NF-1 binding sites have been eliminated. Different cell types were cotransfected with an internal control standard and reporter plasmids with the CAT gene under the control of the UL127 promoter with or without the NF-1 binding sites. In the transient transfection assays, the wt UL127 promoter construct pSnaB-BamCAT (containing sequences between nt –243 and –741 of the MIEP) did not show any significant CAT activity in a variety of cell types (Fig. 2) despite the presence of the MIEP enhancer. The construct pSnaB-Bam(HI)CAT lacking the downstream NF-1 binding site also failed to develop significant promoter activity (Fig. 2; compare CAT responses for pSnaB-BamCAT and pSnaB-Bam(HI)).

**Fig. 3.** Identification of the sequences that mediate transcriptional repression of the UL127 promoter. (A) Binding of various transcription factors (marked by open boxes) to HCMV/MIP sequences between –710 and –220 (basepair footprinting data [reviewed in reference 20]). The location of the UL127 TATA box is shown. Numbers refer to nucleotide positions relative to the transcription start site (+1) of the MIEP. Luciferase (LUC) reporter constructs with various 5′ and 3′ deletion endpoints are shown below. (B) HeLa and NT-2/D1 cells were transfected with 5 μg of the various promoter deletion luciferase constructs shown in panel A together with 5 μg of the control plasmid pRL-tk. Cell lysates were prepared 30 h after transfection and assayed for luciferase activity. The activity of each promoter deletion was normalized to the activity of the internal control plasmid. A plot of the normalized percentage of luciferase activity calculated for each construct, taking as 100 the activity presented by pGL3 (~531), is shown. The luciferase values shown represent the average ± standard deviation (bars) of three determinations.

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**Table 1.** Sequences upstream of the NF-1 cluster mediate transcriptional repression of the UL127 promoter site. Evidence for direct repression of the MIEP by its own IE2 gene product has been provided to account for the restricted expression of the IE genes in the later stages of infection (14, 30, 31, 34, 38, 39, 42). This repression system is defined by the competitive binding of the IE repressor with the RNA polymerase II recruitment step (31, 42). Our previous studies indicated that a strict requirement for this type of repression is that the position of the repressor binding site should be in the vicinity of the transcription start site (31). In other studies, binding sites 5′ proximal to a TATA box have also been shown to block the recruitment of an RNA polymerase II complex (reviewed in reference 1). The UL127 promoter is located in the center of the NF-1 cluster in which one NF-1 binding site exists 14 bp downstream of the UL127 TATA box in the immediate vicinity of the start site and another lies 10 bp upstream of the TATA box (25, 27). It is possible that these particular TATA proximal NF-1 sites may play a role in maintaining a repressed state. To investigate whether these NF-1 sites are involved in repression, we examined constructs in which promoter-proximal downstream and upstream NF-1 binding sites have been eliminated. Different cell types were cotransfected with an internal control standard and reporter plasmids with the CAT gene under the control of the UL127 promoter with or without the NF-1 binding sites. In the transient transfection assays, the wt UL127 promoter construct pSnaB-BamCAT (containing sequences between nt –243 and –741 of the MIEP) did not show any significant CAT activity in a variety of cell types (Fig. 2) despite the presence of the MIEP enhancer. The construct pSnaB-Bam(HI)CAT lacking the downstream NF-1 binding site also failed to develop significant promoter activity (Fig. 2; compare CAT responses for pSnaB-BamCAT and pSnaB-Bam(HI)).

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CAT]. In addition, pGACC(−673), a reporter plasmid that contains neither downstream nor upstream NF-1 sites, was found to be equally ineffective in developing detectable levels of reporter gene expression [Fig. 2; compare CAT responses for pSnaB-BamCAT and pGACC(−673)]. While these experiments do not unequivocally exclude the possible involvement of NF-1, it appears that repression is not due to promoter-proximal NF-1 binding sites, indicating that additional cis-acting repression sequences might be involved. In agreement with this suggestion and in marked contrast to deletion of the NF-1 sites described above, deletion of sequences further upstream (to nt −531 relative to the MIEP transcription start site) resulted in high levels of reporter gene expression in a variety of different cell types [Fig. 2; compare CAT responses for pSnaB-BamCAT and pGACC(−531)]. It is noteworthy that the level of reporter gene activity is less than that observed with the MIEP enhancer constructs (A. Angulo and P. Ghazal, unpublished data). The reason for this lower activity is not known, but it may be due to removal of positive elements in the deletion constructs and/or presence of further repressor elements. Alternatively, the UL127 promoter may require viral factors for maximal activity. Nevertheless, the results of these experiments raise the possibility that sequences located 160-bp upstream of the UL127 promoter mediate strong repression of the UL127 promoter site.

**UL127 promoter-proximal upstream sequences block enhancer activation.** We next examined in more detail the UL127 promoter-proximal upstream sequences involved in mediating transcriptional repression. Accordingly, a series of 5′ and 3′ deletion mutants (from nucleotide positions between −531 and −691) were constructed and tested for their effects on reporter (luciferase) activity in transient transfection experiments. The locations of the deletion endpoints are shown in Fig. 3A in nucleotide positions relative to the initiation site (+1) of the MIEP. In transient transcription assays, deletion of sequences between −691 and −634, in pGL3(−634), did not alter reporter activity in these cell lines [Fig. 3B; compare luciferase responses for pGL3(−673) and pGL3(−634)]. When additional sequences between −634 and −694 were deleted, an approximately 10-fold increase in reporter activity was detected [Fig. 3B; compare luciferase responses for pGL3(−673) and pGL3(−604)]. Significantly, when the construct pGL3(−581), which has an additional 23-bp region deleted, was analyzed in transient transfection assays, a further threefold increase in luciferase activity was observed [Fig. 3B; compare pGL3(−581) and pGL3(−604)]. Deletion of sequences between −581 and −556, in pGL3(−556), resulted in levels of reporter gene activity comparable to the ones exhibited by pGL3(−531), in which sequences from −691 to −531 were absent (Fig. 3C). Altogether, these results indicate that a region between nt −556 and −634 of the MIEP mediates transcriptional repression of the UL127 promoter in a cell-type-independent manner.

In good agreement with these results, a 107-bp internal deletion mutant that eliminated sequences between −531 and −638, pGL3(Δ−531/−638), exhibited a 25- to 50-fold enhancement in the transcriptional activity compared with pGL3(−673) (Fig. 3). To verify further the boundary of these cis-acting repressor sequences, a mutant construct containing a shorter internal deletion of 78 bp (from nt −531 to −609) was generated (Fig. 3A). Consistent with the observed activity of pGL3(−604), this construct [pGL3(Δ−531/−609)] presented a lower reporter activity than pGL3(Δ−531/−638), indicating that sequences between −609 and −638 play a role in maintenance of the repression state (Fig. 3B). We note that these constructs collectively accommodate spacing considerations.

For instance, pGL(−604) has deleted 87 bp from the promoter region, and the spacing difference between the enhancer and promoter in pGL(−531) and pGL(−531/−609) is 82 bp. In all cases, in the absence of repressor sequence, the enhancer is capable of efficiently activating the UL127 promoter in a spacing-independent manner. Altogether, we conclude from these transient transfection results that sequences between nt −556 and −634 cooperate in mediating transcriptional repression of the UL127 promoter site. These experiments thus identify within the unique sequence region (usr) a boundary domain as the unit of transcriptional repression.

To investigate whether these sequences could block enhancer activation on a heterologous promoter, the UL127 TATA box region was substituted with a core promoter fragment (from nt −66 to +112) containing the MIEP TATA box. For these experiments, three plasmids, pMIEP(−66/+112)CAT, pE(−66/+112)CAT, and pEusr(−66/+112)CAT, were constructed (Fig. 4A) and analyzed in transient transfection assays in a variety of different cells. Figure 4 shows that the pE(−66/+112)CAT reporter construct containing the enhancer region (from nt −240 to −531) upstream of the core promoter resulted in a marked increase (from 38- to 88-fold) in CAT reporter activity in comparison with the construct without the enhancer [pMIEP(−66/+112)CAT]. Strikingly, the construct pEusr(−66/+112)CAT, containing 67 bp of additional sequence from the usr (nt −531 to −598) between the enhancer and the MIEP TATA box, resulted in approximately 90% inhibition of enhancer-mediated activation of transcription [Fig. 4B; compare CAT responses for pE(−66/+112)CAT and pEusr(−66/+112)CAT]. These results support the conclusion that boundary element sequences are capable of mediating repression of enhancer function on a heterologous promoter.

We next sought to evaluate whether this mode of repression could also be observed in vitro in the context of transcription reactions in reporter plasmids pMIEP(−66/+112)CAT, pE(−66/+112)CAT, and pEusr(−66/+112)CAT were resected with EcoRI and assayed in transcription runoff assays using HeLa cell nuclear extracts. Transcription reactions were normalized to an internal control standard, and the amount of specific initiation of transcription was quantified by PhosphoImager analysis. As shown in Fig. 5A, the template with the MIEP core promoter [pMIEP(−66/+112)CAT] (lane 1) was transcribed at a much (fivefold) reduced level in comparison with the enhancer-containing template, pE(−66/+112)CAT (lane 2), indicating the ability of the CMV enhancer to stimulate transcription in vitro, albeit at a level much lower than observed in vivo. This level of enhancer-mediated activation was significantly lower than for transcription reactions using the pEusr(−66/+112)CAT template (compare lanes 2 and 3 in Fig. 5B). Similar inhibition results were obtained with different concentrations of reporter template (data not shown). These results suggest that cis-acting sequences within the usr repress enhancer activated transcription but do not appear to influence basal promoter activity. This is consistent with the HeLa cell transfection assay, which indicated a reduction of transcriptional activity by as much as 90% (Fig. 5A). These experiments indicate that it is possible to recapitulate, in part, repression by the boundary domain with the use of an in vitro transcription system, supporting the notion that these sequences may interact with repressor proteins to promote the process of transcriptional repression.

**UL127 promoter-proximal upstream sequences mediate repression of transcription in the context of an HCMV infection.** To determine whether this boundary domain is responsible for mediating repression of the UL127 promoter in an HCMV-
infected cell, we constructed a series of HCMV recombinants containing coding sequences for the GFP under the control of various UL127 promoter deletion mutants. To generate these HCMV recombinants, we used the recently described HCMV BAC system (6). In this system, the HCMV genome has been cloned and maintained as a BAC in *E. coli*, whereby viral progeny can be reconstituted after transfection of the HCMV BAC into eukaryotic cells permissive for HCMV. For the purpose of these studies, we constructed three independent HCMV BAC recombinant genomes, UL127-GFP1, UL127-GFP7, and UL127-GFP10. A schematic representation of parental HCMV and the HCMV BAC recombinants generated is shown in Fig. 6A. The recombinant virus, UL127-GFP1, contains the GFP ORF under the control of the UL127 promoter lacking sequences between nt 2673 and 2691, removing the 5'-proximal NF-1 site. Note that in transient transfection assays, deletion of these sequences did not result in any significant promoter activity. The second recombinant HCMV BAC, UL127-GFP10, contains the GFP ORF under the control of the UL127 promoter lacking sequences between nt 2531 and 2691 (Fig. 6A). The structure of the recombinant BACs generated was verified by extensive restriction analysis (*SalI*, *SnaBI*, and *SpeI*) and sequencing. Figure 6B shows the *SalI* restriction patterns. The DNAs from the recombinant HCMV BACs were identical to the DNA from the parental HCMV BAC pHB5 except for the presence of a new *SalI* fragment (7.9 to 8.1 kb) in the place of the natural *SalI* 7.4-kb fragment. This new fragment was generated, as predicted, from the introduction of the GFP ORF downstream of the mutant UL127 promoter sequences (Fig. 6A). These results indicate that the expected recombination events occurred within the UL127 region of the viral genome. MRC-5 cells were subsequently transfected with the HCMV BAC recombinants. Plaques developed in the cultures, and progeny mutant viruses were recovered from infected cells.

We next sought to examine whether deletion of repressor sequences within the UL127 promoter, identified in transient transfection assays, could also lead to an induction of transcription from the UL127 promoter in the context of a viral infection. For these experiments, RT-PCR was performed using RNA from cells infected with the recombinant viruses under study. HFF cells were infected at an equivalent MOI (0.1 PFU/cell) with the different recombinant viruses and parental HCMV RVHB5 reconstituted from the BAC plasmid...
pHB5 as a negative control. RNA from infected cells was harvested at 13 hpi, treated with DNase, and reverse transcribed. Two primer sets were designed to detect the presence of the GFP transcript. Both primer sets contained a common 3′ primer within the GFP gene. In the first primer set, the 5′ primer was chosen to anneal with sequences 29 to 55 bp downstream the predicted UL127 TATA box. This primer pair should detect a 714-bp product if GFP transcripts were present in infected cells. In the second primer set, the 5′ primer was designed to anneal with the predicted UL127 TATA box. This second primer set should either fail to detect a product in infected cells if the predicted UL127 TATA box was used to transcribe the GFP gene or yield a 750-bp product in the case of using an alternative TATA box upstream the predicted UL127 TATA box. As expected, when cells were infected with the parental virus RVHB5, which does not contain the GFP gene, a specific PCR-amplified product was not detected with any of the two primer sets used (Fig. 7A and B, lane 2). Similarly, when cells were infected with UL127-GFP1, the recombinant virus in which sequences from −673 to −691 of the UL127 promoter were absent, amplified PCR products were not obtained (Fig. 7A and B, lanes 4). In marked contrast, in cells infected with UL127-GFP7 and −10, the expected 714-bp PCR product resulting from amplification of GFP sequences using the first primer set was easily detected (Fig. 7A, lanes 6 and 8). However, RT-PCR performed on HFF cells infected with these two viral recombinants using the second primer set that contained the primer designed to anneal within the predicted UL127 TATA box did not yield significant levels of an amplified product (Fig. 7B, lanes 6 and 8). These data show that sequences between −531 and −638 (relative to the MIEP) play an important role in the blockage of the UL127 promoter in the context of a viral infection. In addition, these results indicate that transcription in cells infected with recombinant viruses lacking the boundary repressor sequences predominantly originates approximately 30 bp downstream the UL127 promoter TATA box. Interestingly, transcripts derived from

FIG. 5. Boundary domain sequences within the UL127 promoter mediate repression in an in vitro transcription system. (A) HeLa cells were transfected with 5 μg of either pMIEP(−66/+112)CAT (lane 1), pE(−66/+112)CAT (lane 2), or pEusr(−66/+112)CAT (lane 3) along with 5 μg of the control plasmid pRSV-B-gal. Transfections and CAT assays were performed as described in the legend to Fig. 2 and in Materials and Methods. A plot of the normalized percentage of CAT activity calculated for each construct, taking as 1 the activity presented by pMIEP(−66/+112), is shown. (B) In vitro runoff transcription assays with different DNA templates, pMIEP(−66/+112)CAT (lane 1), pE(−66/+112)CAT (lane 2), and pEusr(−66/+112)CAT (lane 3), linearized with EcoRI. The transcription reactions were performed, processed, and subjected to polyacrylamide gel electrophoresis (18). The 360-nt specific runoff transcript is indicated by an arrow. The specific runoff transcripts were normalized to an internal control, and the amount of transcript was determined by PhosphorImager analysis (42). A plot of the normalized percentage of specific transcription for each DNA template, taking as 100 the activity developed by pE(−66/+112)CAT, is shown.

FIG. 6. Construction of UL127-GFP HCMV BAC recombinant genomes. (A) The top line represents the parental (wt) HCMV genome with the Sfl fragment (nt 169746 to 177147 [11]) containing the MIE region expanded below. Sequences corresponding to the enhancer and usr (open rectangle) and the UL127 ORF (black box) are indicated. Recombinant HCMV BAC genomes, UL127-GFP1, UL127-GFP10, and UL127-GFP7, containing the GFP ORF (open rectangle) under the control of the various UL127 promoter deletion mutants are shown. The diagram is not drawn to scale. (B) Ethidium bromide-stained agarose gels of SalI-digested BAC plasmids UL127-GFP1, UL127-GFP10, UL127-GFP7, and parental HCMV BAC (pHB5) after separation on a 0.5% agarose gel. Positions of size markers are shown at the right; sizes of the natural and new SalI fragments for each virus are shown with arrows.
the UL127 promoter are cycloheximide sensitive, indicating early expression kinetics instead of the anticipated IE expression (data not shown).

To further corroborate whether sequences within the UL127 promoter repress transcription in the context of a viral infection, we examined GFP reporter activity using confocal microscopy and FACS. In the first experiments, HFF cells were infected at an MOI of 0.5 PFU/cell with recombinant viruses UL127-GFP1, UL127-GFP7, and UL127-GFP10 and parental HCMV RVHB5. On day 3 postinfection, cells were fixed, prepared for immunofluorescence analysis using a monoclonal antibody specific for the MIE proteins and a secondary antibody conjugated to TRITC, and visualized by confocal microscopy. As expected, when cells were infected with the parental virus RVHB5, only nuclear staining due to the expression of the MIE proteins could be observed (Fig. 8, panels 1A and 2A). When cells were infected with UL127-GFP1, with sequences from −673 to −691 of the UL127 promoter deleted, GFP fluorescence was not detectable, while cells were clearly positive for expression of the MIE proteins (compare panels 1B and 2B in Fig. 8). In marked contrast, infection of HFF with UL127-GFP7 and UL127-GFP10 resulted in significant amounts of GFP expression. Confocal microscopy images showed that infected HFF cells simultaneously exhibited nuclear staining due to the MIE proteins and expressed GFP in the nucleus and cytoplasm (Fig. 8, panels C1 to -3 and D1 to -3). Therefore, sequences between −531 and −638 (relative to the MIEP) are sufficient in shutting off the UL127 promoter in the context of a viral infection in cells where the divergent MIE promoter is strongly active.

In the next set of experiments, we quantified the levels of reporter activity by directly measuring GFP levels. For these experiments, we infected HFF cells at an MOI of 0.5 PFU/cell with the different recombinant viruses and measured fluorescence on day 3 postinfection by flow cytometry. Figure 9 shows that while all cells were GFP negative after infection with UL127-GFP1 (mean fluorescence intensity with a signal-to-noise ratio of 1.3), HFF cells infected with UL127-GFP7 exhibited approximately twofold-higher levels of GFP in comparison with UL127-GFP10-infected cells (compare mean fluorescence intensities with signal-to-noise ratios of 7.2 and 14.1 for UL127-GFP10 and UL127-GFP7, respectively). These results indicate that maximal derepression of the UL127 promoter in the context of the viral infection requires the removal of sequences from −531 to −673. Thus, these data are consistent with the role of sequences between −556 and −638 of the UL127 promoter as a repressor of gene expression in transient transfection assays. However, sequences between −638 and −673 also appear to contribute to the level of repression of the UL127 promoter but only in the context of an HCMV infection.

DISCUSSION

We have presented evidence for the presence of a boundary domain located within the US of the HCMV MIEP, whereby the putative UL127 promoter is prevented from being activated by the potent CMV enhancer. The borders of the boundary domain are positioned between the UL127 TATA box and the enhancer. Our experiments define a region between −556 and −638 (relative to the MIEP start site) that functions to dramatically repress enhancer activation of transcription at the UL127 promoter site both in transient transfection assays and in the context of virus infection. In addition, sequences between −638 and −673 were found to contribute to the level of repression only in the context of virus infection but not in transient transfection experiments.

Previously, DNase I protection experiments with DNA fragments encompassing US sequences demonstrated (18), on the basis of different chromatographic behavior, the interaction of multiple distinct factors. Later experiments identified one of these binding components as composed of a retinoic acid receptor-retinoid X receptor (RAR-RXR) heterodimer (2). While DNA-bound RAR-RXR complexes are known to recruit transcriptional corepressors in the absence of bound ligand (12, 29), our mapping studies show that this site is not required for mediating repression. Sequences immediately adjacent to the RAR-RXR site, between nt −558 and −602 (relative to the MIEP), interact with a complex of proteins that have similar chromatographic behavior (18) (Fig. 10). These sequences resemble binding sites for the forkhead/winged helix family of transcription factors (8, 16, 32). Many members of this family are known to be potent repressors of transcription. These observations suggest that one of the candidate repressor elements may bind a cluster of winged helix repressors, although our results indicate that the repression domain is not limited to these sites alone (Fig. 10). For instance, the binding site between −604 and −632 participates in mediating repression and is known to interact with a factor that shares characteristics of the CTF family but is distinct from NF-1 (18) (Fig.
In addition, sequences (−638 to −673) between the CTF-like binding site and the NF-1 site proximal to the UL127 TATA box also appear to contribute to repression only in the context of the virus (Fig. 10). This region is not known to interact strongly with any nuclear factors but contains weak binding sites for YY1 (from nucleotide positions −675 to −663 and −655 to −642), a factor also known to mediate transcriptional repression (T. Stamminger, personal communication).

Recently, Lundquist and coworkers presented work on the characterization of a repressor region located upstream of the UL127 promoter (33). They found that sequences between −640 and −694 contribute to strong repression, while sequences between −583 and −640 effected repression to a much lesser extent in the context of the viral infection. The results of our study are in agreement with theirs regarding the presence of a repressor region but differ markedly in the precise sequences shown to be important in mediating the shutoff of UL127 expression. We believe that the conclusions drawn from these two independent studies are complementary. In particular, it is likely that the reason for the weaker repression observed by Lundquist et al. (33) for the distal repression sequences (−583 to −640) is that repression sequences between nt −556 and −583 were not removed. Indeed, resection of sequences between −556 and −638 in the context of an infection leads to a dramatic relief of repression (this study). Also in agreement with the work by Lundquist et al. (33), we found that sequences closer to the UL127 TATA box contribute to repression but only in the context of the virus. However, we note that removal of sequences between nucleotide positions −673 and −691 alone was not sufficient, in our study, to effect activation of the UL127 promoter. Therefore, while our results are different and agree only in part, they are highly complementary.

We note that the inability of the boundary region to repress the MIEP over a relatively large distance is compatible with a short-range repression mechanism (21). In this case, binding of repressors to promoter-proximal elements results in the dominant inhibition of a promoter site by upstream activators. Boundary elements have been identified in numerous drosophila and yeast genes (references 22, 26, and 40 and references therein), but few if any have been formally characterized for mammalian genes. In drosophila, short-range repressors have been elegantly shown to provide flexibility in controlling complex genetic loci. Indeed, Gray and coworkers (23) proposed that short-range repression is central for the evolution of complex promoters that are composed of multiple, autonomous regulatory elements as is prominently exhibited by the CMV MIEPs. Although the nature of the enhancer-repressor-pro-
moter interaction is unclear, the results of this study indicate that the viral boundary domain blocks enhancer activation in a disproportional manner. In other words, there are far more activators bound to the enhancer region than there are potential repressor binding sites in the boundary region. There are several possible explanations for the ability of a limited repressor complex to block many activators, some of which challenge our view of how enhancers work. Perhaps the boundary domain recruits by protein-protein interaction a multifactor repression complex that is able to interfere with many activators. Alternatively, it is possible that multiple activators present in the enhancer coordinate communications with the basal transcription machinery by interacting with a central switching (adapter/nodal) complex. In this scenario, the boundary domain might disrupt the ability of the adapter/node to interact with the promotor. It should be noted that there is a precedent for a central switching unit in coordinating interactions with all other cis-acting regulatory modules, as demonstrated by the Endo16 system of the sea urchin embryo (43). In this example, module A of the Endo16 cis-regulatory system communicates

![Graph showing GFP expression from UL127 GFP HCMV recombinants.](image1)

**FIG. 9.** GFP expression from UL127 GFP HCMV recombinants. HFF cells were infected with UL127-GFP1, UL127-GFP10, and UL127-GFP7 at an MOI of 0.5 PFU/cell. On day 3 postinfection, cells were trypsinized and fixed with 1% formaldehyde. Fluorescence was measured by flow cytometry as described in Materials and Methods. A histogram of FL1 versus the log fluorescence intensity (detected at 530 nm) is shown. A total of 10,000 events were collected for each sample. Mean fluorescence intensities with signal-to-noise ratios of 25.4/19.8, 154.6/18.8, and 222.1/15.8 were obtained for UL127-GFP1, UL127-GFP10, and UL127-GFP7, respectively.

![Diagram showing sites of protein-DNA interaction on MIEP sequences between -538 and -738.](image2)

**FIG. 10.** Sites of protein-DNA interaction on MIEP sequences between -538 and -738. Sequences between -556 and -638, corresponding to the repressor region defined in transient transfection assays, are marked by reverse print. Sequences from -638 to -673, shown to contribute to repression of the UL127 promoter in the context of the infection, are marked by a gray box. Black underlining represents the extent of strong protection observed using the phosphocellulose (p11) gel chromatography fractions P11 0.3, P11 0.6, and P11 1 (18). The cellular factors known to interact with the protected sequences are indicated. The UL127 TATA box is shown. Numbers refer to nucleotide positions relative to the transcription start site (+1) of the MIEP.
the status of the whole regulatory system directly to the basal transcription machinery. All upstream sites work through module A and in the endogenous arrangement do not themselves interact directly with the basal transcription apparatus.

In conclusion, we have shown that repression can play an important role in determining the ability of the enhancer to activate transcription. Until now theusr was characterized as a positive control region for the MIEP (2, 18). We now show that the ability of the HCMV enhancer to bidirectionally activate transcription is stringently regulated in a negative manner by a boundary domain. This work thus provides a molecular explanation for limiting the action of the enhancer to the MIEP. Restricting the boundary of action of the CMV enhancer is likely to have important biological implications for ensuring sequential and coordinate regulation of transcription. It may not be advantageous or may even be deleterious to the virus to have genes, other than select IE genes, influenced by the proximal and long-range effects of an innately strong enhancer. The identification of the repressor proteins and target sequences, the precise mechanism for effecting inhibition, and the functional consequences of altered physiological states on UL127 promoter activation are topics that remain to be explored.

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